# The Case for Phytoplankton Blooms Under Antarctic Sea Ice

Christopher  $Horvat^{1,1,1}$ , Sarah Seabrook<sup>2,2,2</sup>, Antonia Cristi<sup>2,2,2</sup>, Lisa Matthes<sup>3,3,3</sup>, and Kelsey M Bisson<sup>4,4,4</sup>

November 30, 2022

#### Abstract

Areas covered in compact sea ice are often assumed to prohibit upper ocean photosynthesis. Yet under-ice phytoplankton blooms (UIBs) have increasingly been observed in the Arctic, driven by anthropogenic changes to the optical properties of Arctic sea ice. Here we show the Southern Ocean can also support widespread UIBs. Using under ice-enabled BGC-Argo float data, we detail numerous high phytoplankton biomass events below compact sea ice preceding seasonal ice retreat, and classify 12 distinct UIB events. Using joint light, sea ice, and ocean conditions obtained from the ICESat-2 laser altimeter and 11 climate model contributions to CMIP6, we find that more than 4 million square kilometers of the compact-ice-covered Southern Ocean could support these events in late spring and early summer.

<sup>&</sup>lt;sup>1</sup>Brown University

<sup>&</sup>lt;sup>2</sup>National Institute of Water and Atmospheric Research

<sup>&</sup>lt;sup>3</sup>University of Manitoba

<sup>&</sup>lt;sup>4</sup>Oregon State University

# Phytoplankton Blooms Under Antarctic Sea Ice

Christopher Horvat,<sup>1\*</sup>, Kelsey Bisson<sup>2</sup>, Sarah Seabrook,<sup>3,4</sup>, Antonia Cristi, <sup>3,5</sup> Lisa Matthes<sup>6</sup>

<sup>1</sup>Institute at Brown for Environment and Society, Brown University, Providence, RI, USA

<sup>2</sup>Department of Botany and Plant Pathology, Oregon State University, Corvallis, OR, USA

<sup>3</sup>National Institute of Water and Atmospheric Research, Wellington, NZ

<sup>4</sup>School of the Environment, University of Auckland, Auckland, NZ

<sup>5</sup>Department of Marine Sciences, University of Otago, Dunedin, NZ

<sup>6</sup>Takuvik Joint International Laboratory, Université Laval, Quebec City, QC, CA

\*To whom correspondence should be addressed; E-mail: horvat@brown.edu.

3

under-ice phytoplankton blooms (UIBs) have increasingly been observed in the Arctic, driven by anthropogenic changes to the optical properties of Arctic sea ice. Here we show the Southern

Areas covered in compact sea ice are often assumed to prohibit upper ocean photosynthesis. Yet

Ocean can also support widespread UIBs. Using under ice-enabled BGC-Argo float data, we

8 detail numerous high phytoplankton biomass events below compact sea ice preceding seasonal

ice retreat, and classify 12 distinct UIB events. Using joint light, sea ice, and ocean conditions

obtained from the ICESat-2 laser altimeter and 11 climate model contributions to CMIP6, we

find that more than 4 million square kilometers of the compact-ice-covered Southern Ocean

could support these events in late spring and early summer.

## 1 Introduction

Observations of under-ice phytoplankton blooms (UIBs) in the Arctic Ocean [1] have highlighted the need to understand ecological communities living under compact (local concentration greater than 80%) sea ice, now and under future climate change scenarios [2, and references
within]. Regions supporting UIBs in the Arctic have likely expanded as sea ice has thinned and
become more seasonal. Yet to date, no studies have described nor quantified the potential for
widespread UIBs under Antarctic sea ice, where annual and seasonal variability has changed
less than in the Arctic over the satellite period [3] and where sea ice is typically thinner, more
seasonal, and more fragmented.

Antarctic sea ice typically has a higher albedo than Arctic sea ice [4, 5]. Thus a limited 22 amount of photosynthetically available radiation (PAR, 400-700 nm) can reach the upper ocean 23 directly through sea ice, especially compared to the Arctic, where light transmission through 24 melt-pond-covered sea ice is thought to be a primary cause of UIBs [6]. Still, spring-summer 25 solar irradiance is high: recently a bloom of nanoflagellates was observed under highly reflective landfast sea ice [7]. Floating sea ice in the Southern Ocean is also fractured, thin, and mobile. 27 Small areas of open water, like leads or small openings within the floe mosaic, can allow substantial amounts of light to reach the upper ocean. Sunlight entering the ocean through leads in the Arctic has been shown to initiate phytoplankton blooms, even in areas where sea ice is thick and snow-covered [8]. We therefore investigate this possibility across the sea-ice-covered 31 Southern Ocean. 32

Phytoplankton communities in the Southern Ocean respond rapidly to changes in light conditions, with phytoplankton blooms often observed as soon as the sea ice edge retreats in spring, flooding the mixed layer with light and leaving freshwater rich in iron, main limiters of primary production [9, 10, 11]. In the Arctic, a crucial factor in the development of UIBs is a stable surface mixed layer, which can be induced by melt water and/or increased solar heating of the surface layer [12, 13]. Observations using tagged seals in the Ross Sea show the initiation of a shallow (20 m) surface mixed layer driven by ice melt, preceding the seasonal retreat of sea ice [14]. Yet while shallow mixed layer depths may be necessary to keep phytoplankton in the well-lit surface layer, observations from under-ice Argo floats [15, 16] demonstrate that primary production can be initiated before seasonal sea ice retreat, and even before the restratification of surface waters. This challenges the notion that too-deep surface mixed layers in ice-covered regions of the Southern Ocean limit productivity. Together these factors present the possibility that non-coastal regions of the Southern Ocean, much like the Arctic, are productive before sea ice retreats in summer.

Here we assess the potential for widespread phytoplankton growth under compact, floating,
sea ice in the Southern Ocean. Our primary lines of evidence for observing UIBs are particulatebackscatter-derived phytoplankton carbon (PC) and fluorometric Chlorophyll-a (Chl-a) derived
from under-ice biogeochemical Argo float data (BGC-Argo) (see methods), and we report both
Chl<sub>max</sub>, the maximum Chl-a recorded in each profile, and PC values at the depth of Chl<sub>max</sub>
[PC<sub>max</sub>). While Chl-a is a pigment common to all phytoplankton and has historically been
favored as a metric for phytoplankton, including in under ice studies [17, 18], attributing all
changes in Chl-a to biomass may be biased because of mechanistic (i.e., photoacclimation,
nutrient conditions, growth stage) and methodological concerns [19, 20, 21]. Thus we use
Chl-a in conjunction with nitrate and dissolved oxygen, as data to validate and contextualize
observations of enhanced PC.

After quality control, we examine 38 BGC-Argo floats that operated under sea ice in the Southern Ocean over the period from 2014-2020. 29 could have recorded UIBs as they returned profiles under compact sea ice in austral spring-summer. Nearly all such floats (24/29) record enhanced levels of phytoplankton carbon under compact ice, and 11 record UIBs. These 11

account for nearly half (196/403) of profiles examined for UIBs.

Using supporting Argo, PAR, and sea ice data, we show that UIB measurements follow representative dynamics for light-limited blooms, and use this to define a series of simple criteria
for under-ice photosynthetic activity which is applied to data from ICESat-2 laser altimetry and
11 climate model assessments of Southern Ocean sea ice, light and ocean conditions. Supporting BGC-Argo data, we find the conditions required for such light-limited phytoplankton
blooms are predicted across nearly 50% of regions with compact ice in spring and summer.

These results suggest that in compact, but not completely ice-covered regions of the South-

orn Ocean, enough light reaches into the upper water column to permit primary production, as found in the Arctic [8]. We identify potential sampling regions for examining under-ice primary production and community composition in the Ross Sea, and discuss the implications for sampling strategies and cruise timing.

# Observations of phytoplankton blooms under compact sea ice

Biogeochemical Argo floats record many instances of high phytoplankton biomass under compact sea ice in spring-summer. In Fig. 1(a) we scatter maximum Chl-a measurements for BGC-Argo profiles (Chl<sub>max</sub>, units mg/m<sup>3</sup> see methods) for the months of September to December when local sea ice concentration (SIC) exceeds 80%, overlaid on September-November climatological SIC. For consistency with previous BGC-Argo work [22], and to support Chl<sub>max</sub> observations, we compare Chl<sub>max</sub> measurements with particulate backscattering data ( $b_{bp}$ , at 700nm, units m<sup>-1</sup>) taken at the depth of maximum Chl-a, which we use to compute phytoplankton carbon PC<sub>max</sub> (units mg/m<sup>3</sup>, see methods). We found comparable seasonal cycles under compact ice and a high association (Spearman's R=0.7) between the two (see Supporting Information, Fig. S2). Example profiles of Chl-a and  $b_{bp}$  are provided as Supporting Figure S4

with  $Chl_{max}$  varying from 0.1 to 3.5 mg/m<sup>3</sup>, showing typical covariance of  $b_{bp}$  with Chl-a.

We considered 38 BGC-Argo floats that operated under sea ice in the Southern Ocean, in total recording 1117 profiles. 34 of these floats recorded 873 profiles under compact sea ice, 32 floats recorded 522 profiles during the period from September-December, leaving 29 that recorded 403 profiles under compact sea ice in this period. We define two thresholds to characterize under-ice phytoplankton biomass, based on Chl-a and PC measurements. First we define profiles where both  $PC_{max}$  and  $Chl_{max}$  exceed the interquartile range (of all 1117 profiles) of 13.3 mg/m<sup>3</sup> and 0.13 mg/m<sup>3</sup>, respectively, as having "elevated" photosynthetic activity. 24 out of 29 floats under compact ice from September-December recorded such profiles. The six floats that did not observe elevated  $PC_{max}$  and  $Chl_{max}$  accounted for just 28 total profiles under compact sea ice.

We further define "UIBs" when September-December profiles under compact sea ice record 97 both  $PC_{max}$  and  $Chl_{max}$  three inter-quartile ranges above the all-profile median, or 22.8 mg/m<sup>3</sup> and 0.36 mg/m<sup>3</sup>, respectively. Such Chl-a values are similar to those used to characterize UIBs in the seasonally ice-covered Arctic [23, 24, 25], and the  $PC_{max}$  threshold chosen here exceeds 100 the majority of global phytoplankton carbon observations [26]. In total, 11 of 30 floats recorded 101 at least one UIB, and we record 23 UIB profiles across 12 distinct locations and time periods. 102 Note here we define blooms only in a relative sense, as compared to background values mea-103 sured in the Southern Ocean (i.e, not relative to the global phytoplankton carbon measurements) [19, 27], however these are not globally small values. UIB profiles record an average PC<sub>max</sub> of 105  $36.5 \text{ mg/m}^3$  and average  $\text{Chl}_{max}$  of  $1.2 \text{ mg/m}^3$ . For comparison, phytoplankton carbon calculated from BGC-Argo  $b_{bp}$  in the same manner ranges from 40-90 mg/m<sup>3</sup> during the spring North 107 Atlantic bloom [28], and the highest  $PC_{max}$  observed here in September-December measure-108 ments is 55.0 mg/m<sup>3</sup>. Chl-a, a photophysiological expression of adjustments to light throughout 109 the water column, has greater uncertainty than  $PC_{max}$  for both mechanistic and observational

reasons [21], and so we use it only qualitatively to describe phytoplankton, following [17].

Of all under-ice profiles, many of the highest  $PC_{max}$  values were recorded under compact sea ice, though higher  $PC_{max}$  values were observed on average areas with a looser sea ice cover. Considering the 522 under-ice profiles taken between the months of September-December, we show box plots of  $PC_{max}$  in Figure 1(b), grouped by sea ice concentration (SIC, in 20% bin intervals) to illustrate this fact. A total of 43 profiles under sea ice recorded  $PC_{max}$  above 22.8 mg/m<sup>3</sup>: 5 for SIC from 20%-40%, 5 for SIC from 40% to 60%, 10 for SIC from 60% to 80%, and 23 for SIC from 80% to 100% (compact ice). Median  $PC_{max}$  values are below 12.5 mg/m<sup>3</sup> in each SIC category, and lowest for SIC from 80-100% (9.81 mg/m<sup>3</sup>).

The seasonal cycle in median  $PC_{max}$  increases with the seasonal cycle of downwelling so-120 lar irradiance, though most recorded UIBs occur in November, before the seasonal maximum 121 irradiance. Box plots of  $PC_{max}$  in each month are given in Figure 1(c) for the 903 total profiles 122 under compact sea ice. Median  $PC_{max}$  under compact ice ranged from 9.4 mg/m<sup>3</sup> (n=156) in 123 August, to a high of 17.1 mg/m<sup>3</sup> in December (n=24). The number of recorded UIBs was 0 124 of 553 profiles from July-September, 3 out of 140 profiles in October, 15 out of 95 profiles in 125 November, and 5 out of 24 profiles in December. UIB measurements were recorded under an 126 average SIC of 94%. Because BGC-Argo float records can span multiple years, the "bloom" 127 measurements were recorded in 12 distinct times and locations. In 7 instances, at least 2 succes-128 sive profiles (Argo dives are spaced 10 days apart) were classified as UIBs, 6 including profiles taken in November, with the other including October profiles alone. 130

To examine the drivers of high biomass events under compact sea ice, and permit an analysis of their potential extent using climate model and observational data, we also consider the
composite behavior of supplementary measurements for these profiles. In Fig. 2(a), we plot estimated surface, 25-meter average, and mixed-layer average PAR values (see methods) for each
of the 12 distinct recorded UIBs, referenced in time to the first dive that recorded  $PC_{max}$  above

22.8 mg/m<sup>3</sup>. While there is high variability in estimated surface PAR values for individual floats (grey lines), averaged across all floats, there is a noted increase in PAR towards the UIB observation. By the time of the UIB measurement, both mixed-layer average PAR and 25-meter average PAR cross a threshold of 10  $\mu$ mol photons /m<sup>2</sup>/s, which we use in Sec3 to evaluate the likelihood of observing blooms in model data. The median irradiance at this depth for all UIB measurements is 2.0  $\mu$ mol photons /m<sup>2</sup>/s, similar to observed compensation irradiance in Arctic waters [29, ] and within the range of reported values in the North Atlantic [30, ].

The UIBs found here have an observational signature consistent with light-limited under-ice 143 blooms. In Fig. 2(b-d) we plot dissolved oxygen (b) and nitrate (c) at the depth of  $Chl_{max}$ , and 144 (d) mixed layer depth, all as grey lines, with the average across different UIB measurements as 145 a solid black line and the standard deviation denoted as dashed lines. Leading up to UIB measurements we note decreasing nitrate concentration, increasing dissolved oxygen, and rapidly 147 declining mixed layer depths, covariant with increases in PAR prior to the first UIB profile. 148 Light-limited blooms are often associated with shoaling mixed layers which keep phytoplankton in the euphotic layer [31, ]. The autotrophy rate is set by light and nutrient status, and 150 decreasing nitrate concentrations in the surface compared to deeper in the water column evince 151 photosynthetic activity (noting that nitrate concentrations from deep mixing in the Southern 152 Ocean would give surface values 35 umol/kg, prior to biological uptake). Autotrophy will ini-153 tially exceed heterotrophy in the water column, also leading to increases in dissolved oxygen at the surface. Note that oxygen concentrations are also a function of atmospheric exchange and increased solubility of oxygen with colder waters [32,], which we do not consider here. These results may be compatible with the "disturbance-recovery" hypothesis [33, 15, ], as phytoplank-157 ton are initially in deep MLD, which might act to dilute them from predators where they may 158 accumulate prior to receiving enhanced light with shoaling MLDs. Because of the poor tem-159 poral adjacency of float observations, however, we cannot assess phytoplankton accumulation rates needed to test that hypothesis.

## 162 3 The prevalence of blooms under Antarctic sea ice

The presented BGC-Argo float data showed numerous elevated phytoplankton carbon events under compact sea ice in the Southern Ocean, and we found 12 distinct instances which we classify as under-ice blooms, recorded under sea ice with an average SIC of 94%. Many of these UIBs were recorded in November, when Antarctic sea ice is near its seasonal maximum extent. We next quantify if conditions that support UIBs are widespread across the sea-ice covered Southern Ocean before sea ice retreat.

In Figure 3(a), we show ICESat-2-(IS2)-derived average ocean surface PAR values in the 169 Southern Ocean in November, in which we make the simplifying assumption that no PAR 170 reaches the upper ocean directly through sea ice (see methods). A solid line outlines the com-171 pact sea ice zone (CIZ, SIC > 80%) defined using the NSIDC-CDR SIC product [34]. We also 172 plot the 15% SIC contour, marking the edge of total sea ice extent (SIE). Regions lying inside 173 the SIE contour but outside the CIZ are defined as marginal ice zones (MIZs), which due to 174 the lower albedo of open water, receive higher PAR in the surface water layer compared to the CIZ. Figure 3(b) shows pre-industrial November PAR values for the CESM2 climate model 176 (see methods), with CIZ and MIZ defined from the CESM2 model climatology. Both IS-2 and 177 CESM2 show large areas within the CIZ where ocean surface PAR estimates exceed a "bloom" threshod of 23  $\mu$ mol photons/m<sup>2</sup>/s, sufficient for average insolation within the top 25 meters to exceed 10  $\mu$ mol photons/m<sup>2</sup>/s (see above, methods), representative of the mixed-layer PAR 180 conditions found in BGC-Argo UIBs (Fig. 2a). For the IS-2 estimate of ocean surface PAR, 181 6.9 million km<sup>2</sup> of the November CIZ exceeds that PAR threshold, versus 5.9 million km<sup>2</sup> for CESM2. Because we do not have coincident ocean and sea ice melt observations at the scale of IS2 observations, IS2 estimates only indicate the presence of light in the upper ocean and may

overestimate the area that permits an UIB.

We next consider how frequently an individual grid cell would permit an UIB (see methods),
a metric we term the UIB%. A spatial map of UIB% in November months is given in Figure 3(c)
for CESM2. Areas within the climatological November CIZ (solid line), which has an area of
8.3 million km², permit an UIB 46.4% of the time. Because of year-to-year variability of the
CIZ contour, areas outside of the climatological CIZ also have non-zero UIB%. In those areas,
average UIB% is 19.3%.

We accumulate climatological statistics of UIB-permitting regions in Figure 3(d), compar-192 ing the climatological extent of compact sea ice (red) to the extent of UIB-permitting regions 193 (blue). Large areas support UIBs, peaking at 5.9 million km<sup>2</sup> of compact ice-covered regions 194 in November. The fraction of the CIZ that permits an UIB, the UIB fraction (UIBF), is exam-195 ined in Figure 3(d) (black line, right axis), which peaks in November at an UIBF of 77%. By 196 point of comparison, we reproduce Figure 3(d) as Figure 3(e) for the Arctic Ocean. Up to 4.3 197 million km<sup>2</sup> of the pre-industrial Arctic CIZ is permissive to UIBs, repeating the finding in [2], 198 that large regions of the pre-industrial Arctic also supported UIBs. The seasonal maximum of 199 Arctic UIB area occurs in June, at the peak of the solar cycle, with a peak UIBF of 52% in 200 July. Generally, in the CESM2 picontrol experiments, we find that UIB-permitting regions in 201 the Antarctic are (1) larger, (2) constitute a larger percentage of the CIZ, and (3) peak earlier in 202 the annual solar cycle (November in the Antarctic versus June in the Arctic) than in the Arctic. 203

#### Southern ocean UIB statistics across CMIP6 models

In Figure 4(a,b), we plot the climatological seasonal cycle of Southern Ocean UIB area (a) and UIBF (b) for the 11 CMIP6 models (listed in Supporting Table S2). Across these models, we find a similar seasonal cycle. None of the CMIP6 models have large UIB areas before October, but 10 of 11 have a maximum UIB area in November. Only the MRI-ESM2 model shows a

maximum UIB area in December. Each has a climatological UIB area exceeding 2.66 million km<sup>2</sup>, with a median of 4.75 million km<sup>2</sup>. In Figure 4(c), we show box plots of annual maximum UIB area in the Antarctic for each of the models (filled), compared to annual maximum UIB area in the Arctic (unfilled) for the same years. Out of 11 models, 8 have median Antarctic UIB areas that exceed Arctic UIB areas.

We repeat Figure 4(a,c) in Figure 4(b,d) for the UIBF, with Figure 4(d) showing UIBF values 214 during the month where UIB area is at its maximum (November or December in the Antarctic, 215 June or July in the Arctic). Seasonal cycles of UIBF are similar between models, with most 216 models peaking in December as the CIZ reduces in extent and ocean surface PAR increases. In 217 10 of 11 models, a higher fraction of the Antarctic CIZ permits an UIB than of the Arctic CIZ. 218 Average values of UIBF range from 27-86% (average 57%) in the Antarctic, compared to 26-219 66% in the Arctic (average 37%). Each of the three models in which Antarctic UIB areas were 220 less than Arctic UIB areas have higher UIBF in the Antarctic. Thus we suggest that the reason 221 for differences in the overall magnitude of Antarctic UIB areas is due to differences in model 222 representations of Antarctic and Arctic sea ice, not disagreements about whether sufficient PAR 223 is available under the compact sea ice there. 224

# 4 The potential for observing UIBs in situ

Using BGC-Argo float data, we have demonstrated that high phytoplankton biomass events exist under compact sea ice in the Southern Hemisphere, preceding the seasonal loss of sea ice by several months as well as the seasonal maximum downwelling solar irradiance. Examining a series of climate model estimates of upper ocean light and sea ice conditions, we found that under-ice phytoplankton growth is permitted across wide swaths of the compact ice-covered Southern Ocean. We also found that areas permitting UIBs make up a larger percentage of compact sea ice zones in the Southern Ocean than the Arctic, with an earlier peak in the seasonal

cycle. To validate these results further, observational campaigns will be needed. We specifically focus on the potential of the Ross Sea region (see methods) to support such an event, as it is seasonally ice free, is among the highest-productivity regions of the Southern Ocean, and is known for supporting large ice-algal communities [35, 36].

In Figure 5, we plot UIB% for each of the 11 models during the model period with highest 237 Ross Sea UIB area, which is November in 7 models and December in 4 models. All models 238 have high UIB% in the coastal region near Cape Adare in the Western Ross Sea, which has 239 compact sea ice into January. We identify a region at 72°S, 178.5°E with a blue square in 240 Figure 5. A box plot of UIB% in this location for these 11 models is given in Figure 5 (bottom 241 right), showing a median UIB% there of 62% with a minimum of 40%. Across the CMIP6 242 models, a mean area of 0.55 million km<sup>2</sup> of the Ross Sea is expected to permit UIBs, although 243 the borders of UIB-permitting areas vary by model, and range from 0.29 million km<sup>2</sup> (MRI-244 ESM2, 49% of the Ross CIZ) to 0.95 million km<sup>2</sup> (NorESM2-LM, 65% of the Ross CIZ), with 245 inter-annual variability. Independent of modeled sea-ice area coverage, a large fraction of the 246 Ross CIZ permits UIBs in each year in all models. Figure 4 is repeated as Supporting Figure S1 247 for the Ross Sea region, showing that during the month of highest Ross Sea UIB area, at least 248 49% of the Ross CIZ permits UIBs in each model, on average 60%. 249

## 50 Discussion

Here we explored the potential for under-ice phytoplankton blooms beneath compact sea ice in the Southern Ocean using model simulations, altimetric measurements of sea ice coverage, and BCG-Argo data. We show that on 12 distinct occasions from October-December, BCG-Argo floats recorded UIBs with average maximum Chl-a measurements of 1.8 mg/m³, and derived phytoplankton carbon at the depth of the Chl-a max of 38.0 mg/m³, in areas with an average sea ice concentration of 93%. In addition to these direct high carbon measurements, findings of

elevated ( $PC_{max} > 10 \text{ mg/m}^3$ ,  $Chl_{max} > 0.12 \text{ mg/m}^3$ ) phytoplankton biomass in a large fraction of analyzed float data demonstrates the likelihood for primary production and blooms beneath 258 Antarctic sea ice predating seasonal sea ice retreat. These elevated levels of  $PC_{max}$  under compact ice suggest that even for areas with low open water fraction, incident solar radiation is high 260 enough to promote photosynthetic activity. This is similar to findings in the Arctic where small 261 lead features were sufficient to support under-ice blooming [8], and suggests that small regions 262 of open water are sufficient to relax light limitations on blooming in the summertime South-263 ern Ocean. We supplemented the  $PC_{max}$  observations with observations of nitrate and oxygen 264 taken at the depth as the Chl-a maximum and mixed-layer depths. Increasing oxygen towards 265 the time of peak  $PC_{max}$ , together with decreasing nitrate, supplies evidence for photosynthetic 266 activity. The high association between Chl-a and phytoplankton carbon (Spearman's R = 0.7, 267 see Supporting Info Fig. S2), reinforces that these bio-optical events are associated with higher 268 phytoplankton (i.e., biomass) concentration in the water column. 269

We further used ICESat-2 and an ensemble of climate model estimates of sea ice, light, and oceanographic conditions across the compact-ice-covered Southern Ocean to show that indeed, conditions are favorable for under-ice blooms over wide regions, with a median estimate UIB area of 4.75 million km<sup>2</sup> across the model ensemble. In using ICESat-2 data, we assumed no light reached through sea ice, and all light available for photosynthesis came through open water regions near compact ice. Thus these findings indicate that even in regions with local sea ice concentrations above 80%, and with no light passing directly through the ice, enough open water exists that light does not limit growth in the upper Southern Ocean [16]. As modeled and observed in the Arctic, widespread under-ice productivity, preceding the retreat of seasonal sea ice, may indicate a different ecological system under sea ice than previously understood.

270

271

272

273

274

275

278

279

280

The climate models considered here have inter-related sea ice and light schemes (see Supporting Table S2), and provide estimates of the light conditions in the Southern Ocean. They

may not be accurate if systematic biases in modeled Southern Ocean climate or sea ice proper-282 ties exist. Still, compared against the estimate of upper ocean PAR derived from ICESat-2 data, 283 models produce similar PAR estimates and areas of high surface light levels. We adopted a 284 simple diagnostic criteria for when sufficient light is available to support a bloom, using a fixed 285 PAR threshold in model data based on our observations of UIBs in BGC-Argo data, in line with 286 Arctic modeling studies and observations of acclimation in key Antarctic phytoplankton species 287 [37]. While some BGC-Argo floats do report PAR values, none of the ice-enabled floats used 288 here do. Further observations and modeling of radiative transfer of PAR specifically focused on 289 variable Antarctic sea ice (as in, for example, [38, 39]) would help constrain and evaluate PAR 290 levels needed to trigger blooms in concert with BGC-Argo data. While the Argo data confirms 291 that under-ice regions can be productive, because of their uneven spatial and temporal cover-292 age (Supporting Info Fig. S3), it is not yet possible to directly compare geographic estimates of 293 model-predicted conditions to float data, although we note that locations with the highest profile 294 density also record more frequent UIB events. Thus here we use the Argo data in conjunction 295 with models for understanding the possibility of under-ice blooms, but cannot directly validate 296 model predictions of pan-Antarctic UIB extents. 297

The work we presented here raises an important question: if conditions beneath compact sea ice are favorable for supporting UIBs, and Antarctic sea ice coverage and downwelling irradiance has remained largely stable over the past several decades, why are there no reported observations of under-ice blooms in the Southern Ocean by underway cruises or moorings? We suggest two potential answers.

First, the detection of UIBs requires a dedicated effort to collect in situ chlorophyll data under compact sea ice. An analogy can be drawn to the Arctic Ocean, where spring-summer ice-breaker research expeditions are more common. UIBs are now thought to have been widespread dating back to at least the 1950s (with an overall area coverage that has doubled since 1970 [2]).

303

304

305

But these phenomena, which can have some of the highest levels of integrated biomass of any ecological system [1], were rarely observed before the report of a massive under-ice bloom in the Chukchi Sea in 2011. As we show here, BGC-Argo floats permit a broader sampling of biological parameters across the Southern Hemisphere using consistent methodologies and calibrations. Mining of existing and previous under-ice Chlorophyll data, for example from the BCO-DMO archive, will be a focus of future work to understand whether such events have been observed in the past.

Second, it is possible that UIBs do not occur regularly. While nearly all BGC-Argo floats 314 operating from September-December show elevated phytoplankton carbon measurements pre-315 dating sea ice retreat, only 23 profiles, taken 11 of 30 floats operating in the right time and sea 316 ice conditions, exceed our defined threshold for a "bloom". The threshold established here is 317 defined in both phytoplankton carbon and Chl-a measurements - it exceeds typical values in 318 the global oceans [26] generally and greatly exceeds typical values reported in open water and 319 under-ice conditions in the Southern Ocean [19, 27] specifically. Of these UIB profiles, we 320 recorded 6 events with two or more subsequent measurements meeting the "bloom" threshold. 321 These multi-measurement events occurred in November-December 2016 (Argo id 5904767 and 322 5904180, which was previously discussed in [17]), October-November 2017 (Argo ids 5905100 323 and 5904180), and October-December 2018 (Argo ids 5905375 and 5905636). Further research 324 into these six specific events will be necessary to rule out that the high levels of phytoplankton 325 carbon recorded by these floats were not, for example, advected from a bloom occurring in open water. Our assessment of favorable underwater light conditions for UIBs over large areas of the compact ice zone is also based on a simple set of diagnostic criteria, not detailed biogeochem-328 ical modeling, and uses bulk estimates for light transmission and stratification. We do not take 329 into account iron or other nutrient limitations, nor grazing pressure by higher trophic levels. 330 Instead we follow the perspective of [11] that primary production is primarily light-limited in summer. Thus the UIB-permitting area of the Southern Ocean estimated by CMIP6 models in this study is likely an upper bound on the areas that might bloom in a given year.

This work suggests there is potentially unexplored ecological variability beneath South-334 ern Ocean sea ice, with several million square kilometers of the ice-covered Southern Ocean 335 potentially permitting blooms before the seasonal retreat of the sea ice edge. We paid spe-336 cial attention to the frequently visited Ross Sea region, and suggest detailed measurements 337 of physical and biogeochemical variables to study under-ice phytoplankton bloom phenology, 338 magnitude and community composition and to compare those to known bloom dynamics in the 339 Arctic Ocean [40]. Sampling during the sea ice-covered season will be challenging, especially 340 as remote sensing technologies presently cannot measure chlorophyll under sea ice. Continued 341 targeted deployment of Remotely Operated Vehicles (ROV) and other autonomous profiling 342 floats [15, 16] to measure under-ice light availability and bio-optical parameters can be comple-343 mentary to ship-based sampling, supported by ICESat-2 measurements used to remotely sense 344 particulate backscatter in ice-free conditions [41] extended into sea-ice-covered regions.

### 6 Methods

346

**Ice-enabled BGC-Argo float data** Autonomous profiling Argo floats equipped with biological sensors are a foundation of Southern Ocean biogeochemical observations because they 348 provide observations with consistent sampling methodologies in places (and at sampling fre-349 quencies) inaccessible via ships, and with depth resolution inaccessible via satellite, while ex-350 periencing minimal biofouling and lateral drift [42]. Because Argo floats drift with the currents 351 during their transit, a portion of floats deployed in open water drift into ice-covered regions. To 352 protect the floats from ice damage, an ice-avoidant algorithm (based on a temperature threshold) 353 was implemented to initiate a float's descent when it encounters near freezing surface waters 354 [43]. 355

We calculate phytoplankton carbon from particulate backscattering data  $b_{bp}$  (700 nm) from 356 biogeochemical Argo floats (BGC-Argo) [44]. Particulate backscattering covaries with phy-357 toplankton biomass as phytoplankton scatter light proportional to their concentration and size 358 [45], although  $b_{bp}$  observations do not necessarily imply the presence of only phytoplankton 359 because  $b_{bp}$  can be elevated due to the presence of non-algal particles, especially deeper in the 360 water column where there is enhanced particle sinking. Particulate backscatter has been shown 361 to be a better proxy for phytoplankton carbon compared to fluorometric Chl-a ([26] with less 362 measurement uncertainty for  $b_{bp}$  (on the order of 15%), [46] than for Argo Chl-a. We use 363 the empirically derived phytoplankton carbon relationship in [26], after employing a standard 364 conversion of  $b_{bp}$  (700 nm) to  $b_{bp}$  (470 nm) [47]. 365

BGC-Argo pre-processing and quality control As in [46, 48], all float profiles of Chl-a and  $b_{bp}$  (700 nm, m<sup>-1</sup>) were despiked with a 3-point moving median and we visually confirmed that peak Chl-a and phytoplankton carbon (from  $b_{bp}$ ) values were not from noise in the profile. Examples of four Chl-a and  $b_{bp}$  profiles are given in Supporting Figure S4. We select phy-

toplankton carbon ( $PC_{max}$ ) observations at the depth of the Chl-a maximum to confirm high backscatter measurements correspond to phytoplankton, and that high Chl-a measurements are not the result of photoacclimation.

We exclude profiles if  $Chl_{max}$  is recorded at a depth below 200 meters, or if  $b_{max}$  exceeds 373  $0.01 \text{ m}^{-1}$ , which is in excess of natural values of  $b_{bp}$  found in phytoplankton, possibly indi-374 cating the influence of bubbles or large particles (zooplankton) attracted to the instrumentation 375 [49]. We also include oxygen and nitrate data (units  $\mu$ mol/kg) for comparison with the optical 376 variables (see [20]). The main results of this study, namely the number of observed UIB pro-377 files and their provenance, are insensitive to their presentation in either Chl-a measurements or 378 phytoplankton carbon, and we remake Fig. 1 using  $Chl_{max}$  as Supporting Figure S2 to illustrate 379 the similar seasonal cycle and number of UIBs recorded using Chl-a data alone. 380

We use corrected [20] and quality-controlled data distributed through SOCCOM with a 381 quality flag of either '0' or '1,' indicating it was not checked or received a 'good' quality rating. 382 While most of data within a profile has a flag of '0,', Chl-a has high numbers of 'bad' flags in 383 near-surface observations compared to other variates (of all rated Chl-a observations, 67% were 384 rated 'bad'). We masked and removed any 'bad' data prior to analysis in all cases. Parameters 385 for the floats are sampled at 2m vertical resolution. We did not impose geographical constraints 386 on the data other than that float data come from under ice regions in the Southern Ocean. Obser-387 vations of other variates are reported at the depth of the maximum Chl-a concentration, because 388 while the exact magnitudes of Chl-a may be uncertain, the location of maximum Chl-a in the 389 water column is useful to explore these co-located biologically relevant variables [50, 51]. 390

Values of  $PC_{max}$  and  $Chl_{max}$  reported here may underestimate of the true maximum phytoplankton carbon in the water column, as it is not possible to assess backscatter and Chl-a closer to the surface under sea ice because of the ice-avoidant nature of the Argo floats. Typically, ocean phytoplankton blooms are surface-intensified [1, 52]. For example, the mean depth of

391

392

393

Chl<sub>max</sub> for the UIB profiles was 45 meters. Additionally, as we report  $PC_{max}$  as the value of phytoplankton carbon at the depth of  $Chl_{max}$ , it may not be the same as that of peak PC.

Location information for a float under sea ice is imprecise, as the latitude and longitude 397 coordinates are calculated via a linear interpolation of the pre- and post-sea ice coordinates of a 398 specific float. In some cases, the float will not surface in open water and post-sea ice coordinates 399 are unavailable. Some floats lack under-ice geographic coordinates if they do not surface under 400 open water conditions following a period under ice. Thus it is not possible to interpolate all 401 float location while it transits an ice-covered region, and we remove such floats/dives. We 402 excluded locations where local estimated SIC is less than 15%, profiles where  $b_{bp}$  at the depth 403 of maximum Chl-a exceeded 0.01 m<sup>-1</sup>, and profiles where a mixed layer depth could not be 404 estimated. A list of all floats is provided in the Supporting Information Table S2. 405

CMIP6 model data Remote sensing technologies presently do not directly measure light or chlorophyll beneath sea ice, and most sampling strategies for Southern Ocean photosynthetic communities associated with sea ice focus on in-ice algae communities in coastal regions [53, 54, 55, 56]. We must instead turn to model estimates to describe the joint climatological light, sea ice, and ocean conditions underneath sea ice. We used an ensemble of current-generation coupled climate models contributing to the 6th Coupled Model Intercomparison Project (CMIP6).

While observations show Antarctic sea ice has been stable or increased in extent over the
satellite period (1978-present), CMIP6 models consistently simulate a declining annual-average
Antarctic sea-ice cover over this period [57]. Thus we did not consider it feasible to examine
present-day model estimates of Antarctic sea ice state, which might incorporate biased depictions of sea ice albedo and extent. Instead we postulate that light conditions under Antarctic sea
ice have remained stable over the industrial period, and use data from pre-industrial control run

simulations (CMIP6 runs titled picontrol) in this analysis. Of the full CMIP6 model dataset, 11 simulations (see Supporting Table S1) submitted the required model output we used here.

The ensemble of 11 models produced variable estimates of climate and sea ice state, de-421 spite high interrelation between their sea ice and radiative transfer model components. Differ-422 ent versions of the Community Sea Ice Model (CICE) are used as sea ice model components 423 in 9 of 11 models. There are three substantively different light models, the improved [58] 424 (B+L)  $\delta$ -Eddington multiple-scattering scheme found in CICE versions 5 and above (CESM2 425 and NorESM2 simulations), an earlier version of the B+L scheme found in CICE version 4 426 (CAS), or implementations of simpler Beer-Lambert exponential attenuation of light in ice 427 (CERFACS, MRI). 428

For each CMIP6 model, we defined a climatology of light and sea ice properties using the final 100 years of their respective pre-industrial spinup experiments. In Fig. 3 we specifically examined the Community Earth System Model version 2 (CESM2, [59]) model run, as it uses the more recent version of CICE and the more advanced B+L  $\delta$ -Eddington light scheme. CESM2 produces an overall mean state of Antarctic sea ice that is broadly realistic compared to other CMIP6 models [57, 60], and similar output from CESM2 was analyzed to evaluate the potential for Arctic UIBs in [2].

ICESat-2 light data To supplement model estimates of light under sea ice, we approximated the light field under sea ice using the ICESat-2 laser altimeter. We utilized the L3A along-track sea ice type product (ATL07, [61]) derived from Level 2A ATL03 photon heights [62]. Sea ice types are determined using an empirical decision tree, which identifies whether a given segment is sea ice or water. We developed an estimate of SIC as the ratio of total ice segment length to total segment length. This quantity, the linear concentration  $c^*$ , is related to the SIC, which is defined over a 2-dimensional region. Given the random orientation of crack and open water

features relative to frequent satellite tracks, many repeat 1-D measurements can approximate a 2-D field when sampled sufficiently. In [63], we found global sea ice area metrics derived from passive microwave (PM) satellites were well-approximated by this method in regions where IS2 records at least 1000 individual segments per month. We adopted this same threshold in this study to define  $c^*$ . An advantage of using ICESat-2 segments instead of PM is that ICESat-2 is capable of resolving small cracks and leads that are difficult to observe in PM estimates of local SIC, particularly in summer [64, 65, 66].

From a gridded dataset of  $c^*$ , we estimated the total shortwave irradiance,  $I_0$  ( $\sim$  300–3000 nm), reaching the upper ocean,  $I_0$  (averaged monthly),

$$I_0 = SW(1 - c^*)(1 - \alpha_{oc}) \tag{1}$$

where  $\alpha_{oc} = 0.06$  is the open water albedo and SW is the downwelling solar irradiance at the surface. This shortwave irradiance is then converted to a PAR (400-700 nm) estimate as in the CMIP6 model data (see methods below). This simple model assumes no light passes through the sea ice surface, and the only light available in ice-covered regions comes through the open water part of the area. For this reason we expect ICESat-2 derived downwelling irradiances may be conservative. For SW, we use the reanalyzed estimate of diurnal-average downwelling shortwave irradiance from [67]. We use IS2 data from January 2019-December 2020 to form the present-day climatology of  $I_0$  that is presented in Figure 3.

Argo float sea ice concentration This study includes under-ice profiles initially obtained from 41 BGC-Argo floats suspected to be under ice based on a quality flag noting the float's position was interpolated. To obtain sea ice concentrations (SIC) in the area of float deployment, we matched geographic coordinates for each float to the daily 25-km resolution NSIDC Climate data record SIC product [34].

Argo float mixed layer depths To compute mixed layer depths, we use a density gradient method designed for Southern Ocean mixed-layer depths observed in Argo float data [68]. This method is preferred to standard threshold methods as it prevents near-surface temperature inversions associated with sea ice from impacting depth estimates. In each profile, water column density is computed from temperature and salinity observations, and the mixed layer depth is the first depth where the density gradient exceeds 0.05 kilograms /m<sup>4</sup>. Profiles for which a mixed layer depth cannot be established are masked out as described above.

Argo PAR estimates As no under-ice BGC-Argo floats record onboard PAR estimates, we obtain an estimate of local PAR using the same formalism as with IS-2 PAR estimates 1, using the NSIDC SIC instead of IS-2-derived SIC, assuming no shortwave irradiance penetrates sea ice. Using the Argo-derived mixed layer depth H, we then define mixed-layer average PAR  $I_{ML}$  as,

$$I_{ML} = \frac{I_0}{\kappa H} \left( 1 - e^{-\kappa H} \right). \tag{2}$$

Note that when referring to irradiance we refer to a diurnal average.

- Criteria for permitting an UIB We define an area as "permitting" an under-ice bloom if it meets three criteria:
- 480 **Compact sea ice** Local sea ice concentration exceeds 80%.
- An illuminated upper ocean. Average PAR in the top 25 meters of the ocean exceeds  $10 \mu \text{mol}$  photons/m<sup>2</sup>/s.
- A stable or stratifying surface mixed layer. Sea ice is not refreezing and the upper ocean is non-convecting.

The UIB% is therefore defined as the percentage of model years where a grid cell meets all three criteria together. Thus the UIB% can be low if a region both if it is not frequently covered 486 by compact sea ice, or if the light conditions and ocean stratification are not permissive of a 487 bloom. 488

We focused on those ocean regions under "compact" sea ice to differentiate from phyto-489 plankton growth known to occur as the ice edge retreats in marginal ice zones [69, 70]. Marginal 490 ice zones are typically defined as areas where sea ice concentration is less than 80% [71, e.g.,], 491 thus we used this cutoff to define regions that are "compact" ice. 492

To establish a threshold for upper-ocean PAR, we estimated average PAR,  $\overline{I}$ , at a depth D 493 as, 494

$$\overline{I} = \frac{I_0}{\kappa D} \left[ 1 - \exp(-\kappa D) \right]. \tag{3}$$

Here we assumed that PAR is attenuated exponentially in water with a coefficient  $\kappa$ . We assumed positive photosynthesis (gains outweigh losses) occurs when the average PAR over a 496 25-m deep water column exceeds a threshold value of 10  $\mu$ mol photons / m<sup>2</sup>/s that is exceeded 497 by UIBs recorded in Fig. 2. This value is approximately twice the threshold of integrated 498 daily irradiance of 4.8  $\mu$ mol photons / m<sup>2</sup>/s considered to initiate a phytoplankton bloom in 499 [72, 73, 13], and higher than the levels found to initiate growth in the Southern Ocean [15, 16]. 500 Using  $\kappa = 0.081/m$  [74] for PAR extinction in clear under-ice waters and  $D=25~\mathrm{m}$ 501 establishes a surface PAR threshold value for blooms of  $I_0^* \approx 23 \ \mu \text{mol photons /m}^2\text{/s}$ . CMIP6 502 models typically store and output full-spectrum solar forcing to the upper ocean, but not PAR. 503 We therefore had to convert full spectrum solar irradiance to PAR using a factor of 1.9975  $\mu$ mol 504 photons/J [75, 74]. 505

We also included a threshold for the termination of upper-ocean convection. Under-ice 506 blooms are unlikely to occur when active convection extends below the euphotic zone, such as when leads are actively refreezing with the ocean at its freezing point [12]. The requirement

507

that the upper ocean is non-convecting is similar to the "turbulent shutdown" theory used to explain mid-latitude phytoplankton blooms [76]. GCMs used here are too coarse to resolve 510 the complex boundary layer dynamics that result from surface melting of sea ice [77, 78, 79], 511 and thus they are not suited for determining the convective state of the upper ocean in the 512 presence of sea ice leads. Instead, we considered the ocean to be non-convecting if sea ice was 513 melting at its base, which would lead to stratification of the upper ocean, consistent with Argo 514 observations of high negative covariance between shoaling MLD and phytoplankton biomass 515 under ice [18]. In practice, simply non-zero basal melting does not restrict the location of UIBs 516 as small monthly-averaged basal melt rates occur whenever sea ice is present. We therefore 517 set a positive threshold for the sea ice basal melt rate  $\dot{h}$ , which we expressed as an equivalent 518 heat flux  $Q=\rho_i L_f \dot{h}$ , with  $\rho_i$  = 920 kg/m $^3$  the sea ice density and  $L_f=3.34\times 10^{-5}$  J/kg the 519 latent heat of fusion. As a result Q is required to exceed 5 W/m<sup>2</sup>, for an approximate basal melt 520 rate of  $\dot{h}=5$  cm/month. While turbulent vertical mixing related to sea ice motion can have 521 a significant impact on local circulation, it does not typically extend beyond several meters in 522 the ocean [80, 81], and therefore likely does not impact convection at the depths of  $Chl_{max}$ 523 considered here. 524

The Ross Sea region To define the "Ross Sea region", we roughly followed the convention established by the NIWA Ross Sea Trophic Model [82], taking the ocean region south of 69°S and between 160°W and 170°E longitude. Because of grid variations, the area of this region can vary between CMIP6 models, but its surface area is approximately 1.5 million km<sup>2</sup>.

## **References**

[1] Arrigo, R. Phytoplankton K. al. Massive **Blooms** Unet 530 der Arctic Sea Ice. Science 336. 1408 (2012).**URL** 531

- http://www.sciencemag.org/cgi/doi/10.1126/science.1215065.
  9605103.
- [2] Ardyna, M. *et al.* Under-Ice Phytoplankton Blooms: Shedding Light on the "Invisible" Part of Arctic Primary Production. *Frontiers in Marine Science* **7**, 1–25 (2020). URL https://www.frontiersin.org/articles/10.3389/fmars.2020.608032/full.
- [3] Parkinson, C. L. A 40-y record reveals gradual Antarctic sea ice increases followed by decreases at rates far exceeding the rates seen in the Arctic. *Proceedings of the National Academy of Sciences of the United States of America* **116**, 14414–14423 (2019).
- [4] Brandt, R. E., Warren, S. G., Worby, A. P. & Grenfell, T. C. Surface Albedo
  of the Antarctic Sea Ice Zone. *Journal of Climate* 18, 3606–3622 (2005). URL
  https://journals.ametsoc.org/jcli/article/18/17/3606/30648/Surface-Albe
- 543 [5] Arndt, S. et al. Influence of snow depth and surface flooding on light trans544 mission through Antarctic pack ice. Journal of Geophysical Research: Oceans
  545 122, 2108–2119 (2017). URL http://www.nature.com/articles/175238c0
  546 http://doi.wiley.com/10.1002/2016JC012325.
- [6] Arrigo, K. R. et al. Phytoplankton blooms beneath the sea ice in the Chukchi sea.

  Deep-Sea Research Part II: Topical Studies in Oceanography 105, 1–16 (2014). URL

  http://linkinghub.elsevier.com/retrieve/pii/S0967064514000836.
- 550 [7] Saggiomo, M., Escalera, L., Saggiomo, V., Bolinesi, F. & Mangoni, O. Phytoplankton Blooms Below the Antarctic Landfast Ice During the Melt Season Between
  Late Spring and Early Summer. *Journal of Phycology* 57, 541–550 (2021). URL

  https://onlinelibrary.wiley.com/doi/10.1111/jpy.13112.

- [8] Assmy, P. et al. Leads in Arctic pack ice enable early phytoplankton blooms below snow-covered sea ice. Scientific Reports 7, 40850 (2017). URL http://www.nature.com/articles/srep40850.
- [9] Martin, J. H., Fitzwater, S. E. & Gordon, R. M. Iron deficiency limits phytoplank ton growth in Antarctic waters. *Global Biogeochemical Cycles* 4, 5–12 (1990). URL
   http://doi.wiley.com/10.1029/GB004i001p00005.
- [10] Comiso, J. C., McClain, C. R., Sullivan, C. W., Ryan, J. P. & Leonard, C. L. Coastal
   zone color scanner pigment concentrations in the Southern Ocean and relationships to
   geophysical surface features. *Journal of Geophysical Research: Oceans* 98, 2419–2451
   (1993). URL http://doi.wiley.com/10.1029/92JC02505.
- [11] van Oijen, T. Light rather than iron controls photosynthate production allocation Southern Ocean phytoplankton populations during and aus-565 Journal of Plankton Research 26, 885–900 (2004). **URL** tral autumn. 566 https://academic.oup.com/plankt/article-lookup/doi/10.1093/plankt/fbh08 567
- 568 [12] Lowry, K. E. *et al.* Under-Ice Phytoplankton Blooms Inhibited by Spring Convective
  569 Mixing in Refreezing Leads. *Journal of Geophysical Research: Oceans* **123**, 90–109
  570 (2018). URL http://doi.wiley.com/10.1002/2016JC012575.
- [13] Oziel, L. al. Environmental factors influencing the et seasonal 571 spring algal blooms in and beneath sea ice in western Bafnamics of 572 fin Bay. Science of the Anthropocene 7, 34 (2019). **URL** Elementa: 573 https://online.ucpress.edu/elementa/article/doi/10.1525/elementa.372/11 574

- For [14] Porter, D. F. *et al.* Evolution of the Seasonal Surface Mixed Layer of the Ross Sea, Antarctica, Observed With Autonomous Profiling Floats. *Journal of Geophysical Research:*Oceans 124, 4934–4953 (2019).
- 578 [15] Arteaga, L. A., Boss, E., Behrenfeld, M. J., Westberry, T. K. & Sarmiento, J. L. Seasonal 579 modulation of phytoplankton biomass in the Southern Ocean. *Nature Communications* 11 580 (2020). URL http://dx.doi.org/10.1038/s41467-020-19157-2.
- [16] Hague, M. & Vichi, M. Southern Ocean Biogeochemical Argo detect under-ice phytoplankton growth before sea ice retreat. *Biogeosciences* **18**, 25–38 (2021). URL https://bg.copernicus.org/articles/18/25/2021/.
- [17] Briggs, E. M., Martz, T. R., Talley, L. D., Mazloff, M. R. & Johnson,

  K. S. Physical and Biological Drivers of Biogeochemical Tracers Within

  the Seasonal Sea Ice Zone of the Southern Ocean From Profiling Floats.

  Journal of Geophysical Research: Oceans 123, 746–758 (2018). URL

  https://onlinelibrary.wiley.com/doi/10.1002/2017JC012846.
- 589 [18] Bisson, K. M. & Cael, B. B. How are under ice phytoplankton related to sea 590 ice in the Southern Ocean? *Geophysical Research Letters* 1–14 (2021). URL 591 https://onlinelibrary.wiley.com/doi/10.1029/2021GL095051.
- 592 [19] Haëntjens, N., Boss, E. & Talley, L. D. Revisiting Ocean Color algorithms for 593 chlorophyll-a and particulate organic carbon in the Southern Ocean using biogeochem-594 ical floats. *Journal of Geophysical Research: Oceans* **122**, 6583–6593 (2017). URL 595 https://onlinelibrary.wiley.com/doi/10.1002/2017JC012844.

- 596 [20] Johnson, K. S. *et al.* Biogeochemical sensor performance in the SOCCOM profiling
  597 float array. *Journal of Geophysical Research: Oceans* **122**, 6416–6436 (2017). URL
  598 https://onlinelibrary.wiley.com/doi/10.1002/2017JC012838.
- mates from in situ chlorophyll fluorometers: A global analysis of WET Labs ECO sensors. *Limnology and Oceanography: Methods* **15**, 572–585 (2017). URL https://onlinelibrary.wiley.com/doi/10.1002/lom3.10185.
- [22] Mayot, N. et al. Assessing Phytoplankton Activities in the Seasonal Ice Zone of the
   Greenland Sea Over an Annual Cycle. Journal of Geophysical Research: Oceans 123,
   8004–8025 (2018). URL http://doi.wiley.com/10.1029/2018JC014271.
- 606 [23] Apollonio, S. Hydrobiological measurements on IGY drifting station Bravo. *Trans. Am.*607 *Geophys. Union* **40**, 316–3 (1959).
- [24] Laney, S. R. *et al.* Assessing algal biomass and bio-optical distributions in perennially ice-covered polar ocean ecosystems. *Polar Science* **8**, 73–85 (2014). URL

  https://linkinghub.elsevier.com/retrieve/pii/S1873965213000510.
- G11 [25] Boles, E. et al. Under-Ice Phytoplankton Blooms in the Central Arctic

  Ocean: Insights From the First Biogeochemical IAOOS Platform Drift in 2017.

  Journal of Geophysical Research: Oceans 125, 6069–6079 (2020). URL

  https://onlinelibrary.wiley.com/doi/10.1029/2019JC015608.
- carbon [26] Graff, R. Analytical phytoplankton J. al. measure-615 Research ments spanning diverse ecosystems. Deep Sea Part 616 (2015).**URL** *I*: Oceanographic Research **Papers** 102, 16 - 25617 https://linkinghub.elsevier.com/retrieve/pii/S0967063715000801. 618

- 619 [27] Moore, J. K. & Abbott, M. R. Phytoplankton chlorophyll distributions and primary pro-620 duction in the Southern Ocean. *Journal of Geophysical Research: Oceans* **105**, 28709– 621 28722 (2000). URL http://doi.wiley.com/10.1029/1999JC000043.
- [28] Cetinić, I. *et al.* Particulate organic carbon and inherent optical properties during 2008 North Atlantic bloom experiment. *Journal of Geophysical Research: Oceans* **117** (2012).
- [29] Tremblay, J. É., Michel, C., Hobson, K. A., Gosselin, M. & Price, N. M. Bloom dynamics
   in early opening waters of the Arctic Ocean. *Limnology and Oceanography* 51, 900–912
   (2006).
- [30] Siegel, D. A., Doney, S. C. & Yoder, J. A. The North Atlantic Spring Phytoplankton Bloom and Sverdrup's Critical Depth Hypothesis. *Science* **296**, 730–733 (2002). URL http://www.sciencemag.org/cgi/doi/10.1126/science.1069174.
- [31] Sverdrup, H. U. On Conditions for the Vernal Blooming of Phytoplankton. ICES Journal of Marine Science 18, 287-295 (1953). URL

  https://academic.oup.com/icesjms/article-lookup/doi/10.1093/icesjms/18.
- G33 [32] Glud, R. N., Rysgaard, S., Turner, G., McGinnis, D. F. & Leakey, R. J. G. Biological- and physical-induced oxygen dynamics in melting sea ice of the Fram Strait. *Limnology and Oceanography* **59**, 1097–1111 (2014). URL http://doi.wiley.com/10.4319/lo.2014.59.4.1097.
- [33] Behrenfeld, M. J. Abandoning sverdrup's critical depth hypothe-637 on phytoplankton blooms. Ecology 91, 977–989 (2010).URL 638 http://www.esajournals.org/doi/abs/10.1890/09-1207.1 639 http://doi.wiley.com/10.1890/09-1207.1. 640

- [34] Meier, W. N., Fetterer, F., Windnagel., A. K. & Stewart, J. S. NOAA/NSIDC Climate Data
   Record of Passive Microwave Sea Ice Concentration, Version 4. (2021).
- [35] Lizotte, M. P. The Contributions of Sea Ice Algae to Antarctic Marine Primary Production. American Zoologist 41, 57–73 (2001). URL https://academic.oup.com/icb/article-lookup/doi/10.1093/icb/41.1.57.
- 646 [36] Arrigo, K. R. Physical control of chlorophyll a , POC, and TPN distributions in the pack 647 ice of the Ross Sea, Antarctica. *Journal of Geophysical Research* **108**, 3316 (2003). URL 648 https://onlinelibrary.wiley.com/doi/10.1029/2001JC001138.
- 649 [37] Arrigo, K. R. *et al.* Photophysiology in two major southern ocean phytoplankton taxa:
  650 Photosynthesis and growth of phaeocystis antarctica and fragilariopsis cylindrus under
  651 different irradiance levels. *Integrative and Comparative Biology* **50**, 950–966 (2010).
- Horvat, C., Flocco, D., Rees Jones, D. W., Roach, L. & Golden, K. M.

  The Effect of Melt Pond Geometry on the Distribution of Solar Energy Under First-Year Sea Ice. *Geophysical Research Letters* **47** (2020). URL

  https://onlinelibrary.wiley.com/doi/abs/10.1029/2019GL085956.
- sights into radiative transfer within sea ice derived from autonomous optical propagation measurements. *The Cryosphere* **15**, 183–198 (2021). URL https://tc.copernicus.org/articles/15/183/2021/.
- 660 [40] Chase, A. P. *et al.* Evaluation of diagnostic pigments to estimate phytoplankton size classes. *Limnology and Oceanography: Methods* (2020).

- 662 [41] Lu, X. et al. Antarctic spring ice-edge blooms observed from space by
  663 ICESat-2. Remote Sensing of Environment 245, 111827 (2020). URL
  664 https://doi.org/10.1016/j.rse.2020.111827.
- Poteau, A., Boss, E. & Claustre, H. Particulate concentration and seasonal dynamics in the mesopelagic ocean based on the backscattering coefficient measured with Biogeochemical-Argo floats. *Geophysical Research Letters* **44**, 6933–6939 (2017). URL http://doi.wiley.com/10.1002/2017GL073949.
- 669 [43] Klatt, O., Boebel, O. & Fahrbach, E. A profiling float's sense of ice. *Jour-*670 nal of Atmospheric and Oceanic Technology 24, 1301–1308 (2007). URL
  671 http://journals.ametsoc.org/doi/abs/10.1175/JTECH2026.1.
- 672 [44] Claustre, H. *et al.* Bio-Optical Profiling Floats as New Observational Tools for
  673 Biogeochemical and Ecosystem Studies: Potential Synergies with Ocean Color Re674 mote Sensing. In *Proceedings of OceanObs'09: Sustained Ocean Observations*675 *and Information for Society*, 1, 177–183 (European Space Agency, 2010). URL
  676 http://www.oceanobs09.net/proceedings/cwp/cwp17.
- [45] Hergert, W. & Wriedt, T. The Mie theory: basics and applications (Springer, 2012).
- ing satellite estimates of particulate backscatter in the global open ocean using autonomous profiling floats. *Optics Express* **27**, 30191 (2019). URL

  https://www.osapublishing.org/abstract.cfm?URI=oe-27-21-30191.
- Deriving inherent opti-[47] Lee, Z., Carder, K. L. & Arnone, R. A. 682 properties from water color: multiband quasi-analytical algorithm a 683

- for optically deep waters. *Applied Optics* **41**, 5755 (2002). URL https://www.osapublishing.org/abstract.cfm?URI=ao-41-27-5755.
- Particulate Backscattering in the Global Ocean: A Comparison of Independent Assessments. *Geophysical Research Letters* **48** (2021). URL https://onlinelibrary.wiley.com/doi/10.1029/2020GL090909.
- [49] Haëntjens, N. et al. Detecting Mesopelagic Organisms Using Biogeochemical-Argo
   Floats. Geophysical Research Letters 47 (2020).
- [50] Ardyna, M. *et al.* Parameterization of vertical chlorophyll a in the Arctic Ocean: Impact of the subsurface chlorophyll maximum on regional, seasonal, and annual primary production estimates. *Biogeosciences* **10**, 4383–4404 (2013).
- [51] Brown, Z. W. et al. Characterizing the subsurface chlorophyll a maximum in the Chukchi
   Sea and Canada Basin. Deep-Sea Research Part II: Topical Studies in Oceanography 118,
   88–104 (2015). URL http://dx.doi.org/10.1016/j.dsr2.2015.02.010.
- [52] SMITH, W. O. & NELSON. D. M. Phytoplankton Bloom Pro-698 a Receding Ice Edge in the Ross Sea: **Spatial** Coherby 699 ence with the Density Field. Science 227, 163–166 **URL** 700 http://www.sciencemag.org/content/227/4683/163.shorthttp://www.sciencem 701
- [53] Smetacek, V. et al. Early spring phytoplankton blooms in ice platelet 702 layers of the southern Weddell Sea, Antarctica. Deep Sea Research 703 *Oceanographic* Research Papers 39, **URL** 153–168 (1992).704 https://linkinghub.elsevier.com/retrieve/pii/019801499290102Y. 705

- [54] Arrigo, K. R. & Thomas, D. N. Large scale importance of sea ice biology in the Southern
   Ocean. *Antarctic Science* 16, 471–486 (2004).
- [55] McMinn, A., Martin, A. & Ryan, K. Phytoplankton and sea ice al-708 gal biomass and physiology during the transition between winter and 709 (McMurdo Sound, Antarctica). Polar Biology 33, 1547-1556 spring 710 **URL** http://www.nature.com/articles/175238c0 (2010).http://doi.wiley.com/10.1002/2016JC012325 712

http://link.springer.com/10.1007/s00300-010-0844-6.

713

- To prince [56] Cummings, V. J. et al. In situ response of Antarctic under-ice prince mary producers to experimentally altered ph. Scientific Reports 9, 6069 (2019). URL http://dx.doi.org/10.1038/s41598-019-42329-0 http://www.nature.com/articles/s41598-019-42329-0.
- 718 [57] Roach, L. A. *et al.* Antarctic Sea Ice Area in CMIP6. *Geophysical Research Letters* 1–24 (2020).
- [58] Briegleb, B. P. & Light, B. A Delta-Eddington multiple scattering parameterization for
   solar radiation in the sea ice component of the Community Climate System Model. Tech.
   Rep. February, National Center for Atmospheric Research (2007).
- [59] Danabasoglu, G. et al. The Community Earth System Model Version 2 (CESM2). Journal
   of Advances in Modeling Earth Systems 12, 1–35 (2020).
- 725 [60] Singh, H. K. A., Landrum, L., Holland, M. M., Bailey, D. A. & DuVivier, A. K. An

  Overview of Antarctic Sea Ice in the Community Earth System Model version 2, Part

  I: Analysis of the Seasonal Cycle in the Context of Sea Ice Thermodynamics and Cou-

- pled Atmosphere-Ocean-Ice Processes. *Journal of Advances in Modeling Earth Systems* (2020).
- [61] Kwok, R. *et al.* ATLAS/ICESat-2 L3A Sea Ice Height, Version 1. Boulder, Colorado
   USA. Tech. Rep. May, NSIDC, Boulder, Colorado USA (2019).
- 732 [62] Neumann, T. A. *et al.* The Ice, Cloud, and Land Elevation Satellite 2 mis-733 sion: A global geolocated photon product derived from the Advanced Topographic 734 Laser Altimeter System. *Remote Sensing of Environment* **233**, 111325 (2019). URL 735 https://linkinghub.elsevier.com/retrieve/pii/S003442571930344X.
- 736 [63] Horvat, C., Blanchard-Wrigglesworth, E. & Petty, A. Observing waves
  737 in sea ice with ICESat-2. *Geophysical Research Letters* (2020). URL
  738 https://onlinelibrary.wiley.com/doi/abs/10.1029/2020GL087629.
- 739 [64] Kwok, R. Sea ice concentration estimates from satellite passive microwave radiometry 740 and openings from SAR ice motion. *Geophysical Research Letters* **29**, 25–1–25–4 (2002). 741 URL http://doi.wiley.com/10.1029/2002GL014787.
- 742 [65] Notz, D., Haumann, F. A., Haak, H., Jungclaus, J. H. & Marotzke, J.

  743 Arctic sea-ice evolution as modeled by Max Planck Institute for Meteorol744 ogy's Earth system model. *Journal of Advances in Modeling Earth Sys-*745 *tems* 5, 173–194 (2013). URL http://dx.doi.org/10.1002/jame.20016

  746 http://doi.wiley.com/10.1002/jame.20016.
- 747 [66] Kern, S., Lavergne, T., Notz, D., Pedersen, L. T. & Tonboe, R. Satel148 lite passive microwave sea-ice concentration data set inter-comparison for Arc149 tic summer conditions. *The Cryosphere* 14, 2469–2493 (2020). URL
  150 https://tc.copernicus.org/articles/14/2469/2020/.

- Tsujino, H. et al. JRA-55 based surface dataset for driving ocean—seaice models (JRA55-do). Ocean Modelling 130, 79–139 (2018). URL

  https://linkinghub.elsevier.com/retrieve/pii/S146350031830235X.
- Dong, S., Sprintall, J., Gille, S. T. & Talley, L. Southern ocean mixed-layer depth from Argo float profiles. *Journal of Geophysical Research: Oceans* **113**, 1–12 (2008).
- 756 [69] Smith, W. O. & Nelson, D. M. Importance of Ice Edge Phytoplankton Pro757 duction in the Southern Ocean. *BioScience* 36, 251–257 (1986). URL
  758 https://academic.oup.com/bioscience/article-lookup/doi/10.2307/1310215.
- <sup>759</sup> [70] Perrette, M., Yool, A., Quartly, G. D. & Popova, E. E. Near-ubiquity of ice-edge blooms in the Arctic. *Biogeosciences* **8**, 515–524 (2011).
- 761 [71] Strong, C. & Rigor, I. G. Arctic marginal ice zone trending wider in summer and
  762 narrower in winter. *Geophysical Research Letters* **40**, 4864–4868 (2013). URL
  763 http://doi.wiley.com/10.1002/grl.50928.
- [72] Letelier, R. M., Karl, D. M., Abbott, M. R. & Bidigare, R. R. Light driven seasonal patterns of chlorophyll and nitrate in the lower euphotic zone of the North Pacific Subtropical
   Gyre. *Limnology and Oceanography* 49, 508–519 (2004).
- Poss, E. & Behrenfeld, M. In situ evaluation of the initiation of the North Atlantic phytoplankton bloom. *Geophysical Research Letters* **37**, 1–5 (2010).
- Matthes, L. C. *et al.* Average cosine coefficient and spectral distribution of the light field under sea ice: Implications for primary production. *Elementa* **7** (2019).
- 771 [75] Yu, X., Wu, Z., Jiang, W. & Guo, X. Predicting daily photosyntheti-772 cally active radiation from global solar radiation in the Contiguous United

- States. Energy Conversion and Management 89, 71-82 (2015). URL

  https://linkinghub.elsevier.com/retrieve/pii/S0196890414008395.
- 775 [76] Taylor, J. R. & Ferrari, R. Shutdown of turbulent convection as a new criterion for the 776 onset of spring phytoplankton blooms. *Limnology and Oceanography* **56**, 2293–2307 777 (2011).
- [77] Holland, improved single-column model representation M. M. 778 associated with summertime leads: Results from a SHEBA ocean mixing 779 Journal of Geophysical Research 108, 3107 (2003). case study. **URL** 780 http://doi.wiley.com/10.1029/2002JC001557. 781
- Horvat, C., Tziperman, E. & Campin, J.-M. Interaction of sea ice floe size, ocean eddies, and sea ice melting. *Geophysical Research Letters* **43**, 8083–8090 (2016). URL http://doi.wiley.com/10.1002/2016GL069742.
- 785 [79] Pellichero, V., Sallée, J.-B., Schmidtko, S., Roquet, F. & Charrassin, J.-B.

  The ocean mixed layer under Southern Ocean sea-ice: Seasonal cycle and forc
  ing. *Journal of Geophysical Research: Oceans* 122, 1608–1633 (2017). URL

  http://doi.wiley.com/10.1002/2016JC011970.
- [80] Smith, M. & Thomson, J. Ocean Surface Turbulence in Newly Formed Marginal Ice
   Zones. Journal of Geophysical Research: Oceans 124, 1382–1398 (2019). URL
   https://onlinelibrary.wiley.com/doi/10.1029/2018JC014405.
- Parameterizations of Ice-Ocean Drag Through an Annual Cycle Across the Beaufort Sea.

  Journal of Geophysical Research: Oceans 126 (2021).

- Pinkerton, M. H., Bradford-Grieve, J. M. & Hanchet, S. M. A balanced model of the food
   web of the Ross Sea, Antarctica. *CCAMLR Science* 17, 1–31 (2010).
- [83] Horvat, C., Seabrook, S., Cristi, A., Matthes, L. & Bisson, K. Code for: The Case for
   Phytoplankton Blooms Under Antarctic Sea Ice (2021).

## **799 7 Corresponding Author**

All correspondence and requests for material should be addressed to Christopher Horvat, christopher\_horvat

## 8 Acknowledgements

CH was supported by NASA grant 80NSSC20K0959 and by Schmidt Futures – a philanthropic initiative that seeks to improve societal outcomes through the development of emerging sci-803 ence and technologies. CH thanks the National Institute of Water and Atmospheric Research 804 in Wellington, NZ for their hospitality during parts of this work. KB was supported by NASA 805 grant 80NSSC20K0970 and thanks Tanya Maurer at MBARI for help with Argo under ice 806 identification. We acknowledge the World Climate Research Programme, which, through its 807 Working Group on Coupled Modelling, coordinated and promoted CMIP6. We thank the cli-808 mate modeling groups for producing and making available their model output, the Earth System 809 Grid Federation (ESGF) for archiving the data and providing access, and the multiple funding 810 agencies who support CMIP6 and ESGF. These Argo data were collected and made freely available by the International Argo Program and the national programs that contribute to it. 812 (https://argo.ucsd.edu, https://www.ocean-ops.org). The Argo Program is part of the Global 813 Ocean Observing System.

#### 9 Author Contributions

- 816 CH and SS concieved of the study. CH performed the data analysis and prepared the manuscript.
- KB provided BGC-Argo data processing and analysis. All authors assisted with study design
- 818 and manuscript writing.

## 819 10 Competing Interests

The authors declare no competing interests.

## 11 Data Availability

- sea ice type product is found online at https://nsidc.org/data/ATL07/versions/3. JRA55-do and
- 824 CMIP6 data used in this study are available at the Earth System Federated Grid at https://esgf-node.lln
- and https://esgf-node.llnl.gov/projects/cmip6/, respectively. Argo data were

ICESat-2 data are available through the National Snow and Ice Data Center (NSIDC). The

- 826 collected and made freely available by the Southern Ocean Carbon and Climate Observations
- and Modeling (SOCCOM) Project funded by the National Science Foundation, Division of Po-
- lar Programs (NSF PLR -1425989 and OPP-1936222), supplemented by NASA, and by the In-
- ternational Argo Program and the NOAA programs that contribute to it. http://www.argo.ucsd.edu,
- https://soccompu.princeton.edu/www/index.html. The Argo Program is part
- of the Global Ocean Observing System. Processed Argo data for making figures is included in
- the code repository [83].

## 33 12 Code Availability

- 834 Code for processing data and producing Antarctic under-ice light fields and UIB-permitting
- criteria is publicly available on github at https://github.com/chhorvat/Antarctic-Light/, with re-

836 leases archived in the Zenodo repository [83]. Data required to produce figures in this manuscript

is included within this repository.

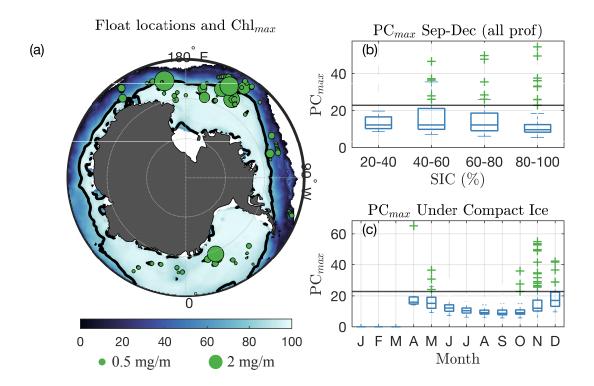


Figure 1: Chlorophyll-a and Phytoplankton Carbon recorded under compact sea ice by BGC-Argo floats. (a) Climatological sea ice coverage in September-November, 2014-2020. Black line shows 80% concentration contour. Green circles are locations of under-ice Argo float profiles under compact sea ice from September-December, with sizes scaled with value of  $Chl_{max}$ . Green dots outside of map shows sizes corresponding to 0.5 and 2.0 mg/m³. (b) Box plot of  $Chl_{max}$  for all BGC-Argo measurements under sea ice, indexed by sea ice concentration. Whiskers extend boxes  $\pm 3$  standard deviations from the mean in each month and vertical blue line is ensemble median. Crosses show values identified as UIBs. Black line is  $PC_{max}$  bloom threshold. (c) Same as (b), but for  $PC_{max}$  recorded under compact sea ice (concentration > 80%) only, indexed by month.

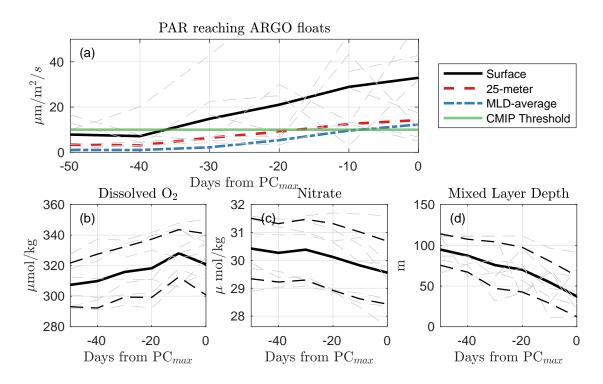


Figure 2: **Under-ice conditions preceding UIBs.** (a) averaged estimated PAR for each of the 13 measured UIBs for the first profile identified as a UIB and the 5 preceding under-ice profiles. Black line is average surface PAR. Red dash line is average PAR over top 25 meters. Blue dot-dash line is PAR averaged over Argo-measured mixed layer depth. Green line indicates the PAR threshold used to define UIBs in CMIP6 data. (b-d) Same as (a, black line) for (b) dissolved oxygen at depth of Chlorophyll-a maximum, (c) nitrate at depth of  $Chl_{max}$ , and (d) mixed layer depth. Dashed lines indicate standard deviation of profile measurements.

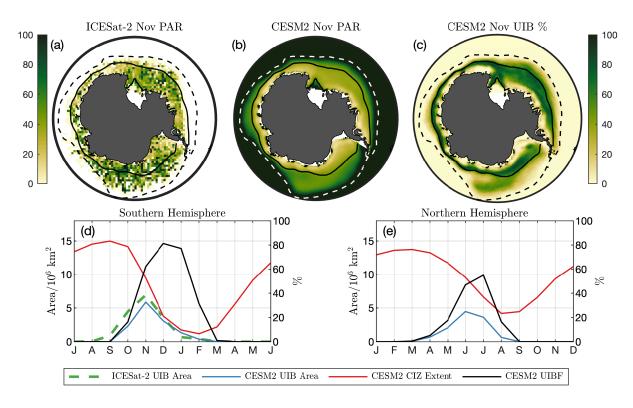


Figure 3: **Light field and UIB potential under Southern Ocean sea ice** (a) 2018-2020 November surface PAR ( $\mu$ mol photons/m²/s) estimate from ICESat-2. (b) CESM2 climatological PAR from pre-industrial simulation. Solid lines in (a-b) are CESM2 climatological CIZ (concentration above 80%). Dashed lines are climatological SIE (concentration above 15%). (c) CESM2 November UIB%. (d,left axis) Seasonal cycle of CESM2 (red) CIZ extent and (blue) UIB extent. Dashed green line is UIB area from ICESat-2. (d,right axis) CESM2 UIBF. (e) As in (d), but for the Northern Hemisphere. Axes in (d) and (e) are offset by 6 months.

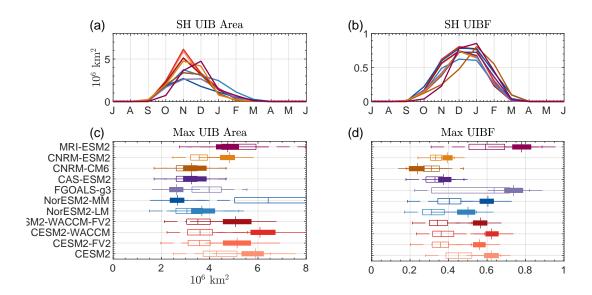


Figure 4: **Statistics of bloom-permitting area for CMIP6 models**. (a) Seasonal cycle of UIB-permitting area in the Southern Hemisphere. (b) Seasonal cycle of UIBF. (c) Box plots of maximum annual UIB area in (filled) the Southern Hemisphere or (unfilled) the Northern Hemisphere. (d) Box plots of UIBF during month of maximum UIB area. Colors of lines in (a,b) correspond to boxes in (c,d)

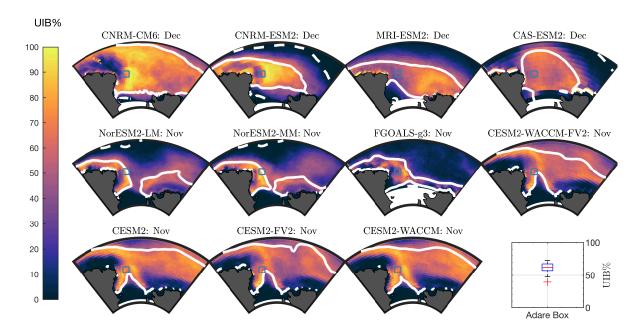


Figure 5: **UIB likelihoods in the Ross Sea.** Ross Sea UIB% for each model in the month of maximum UIB area. Solid lines are climatological CIZ. Dashed lines are climatological SIE. Blue square highlights location of interest at 72°S,178°E. Box plot (bottom right) is of UIB% at square location.

# Phytoplankton Blooms Under Antarctic Sea Ice

Christopher Horvat,<sup>1\*</sup>, Kelsey Bisson<sup>2</sup>, Sarah Seabrook,<sup>3,4</sup>, Antonia Cristi, <sup>3,5</sup> Lisa Matthes,<sup>6</sup>

<sup>1</sup>Institute at Brown for Environment and Society, Brown University, Providence, RI, USA

<sup>2</sup>Department of Botany and Plant Pathology, Oregon State University, Corvallis, OR, USA

<sup>3</sup>National Institute of Water and Atmospheric Research, Wellington, NZ

<sup>4</sup>School of the Environment, University of Auckland, Auckland, NZ

<sup>5</sup>Department of Marine Sciences, University of Otago, Dunedin, NZ

<sup>6</sup>Takuvik Joint International Laboratory, Université Laval, Quebec City, QC, CA

\*To whom correspondence should be addressed; E-mail: horvat@brown.edu.

3

summer.

Areas covered in compact sea ice are often assumed to prohibit upper ocean photosynthesis. Yet under-ice phytoplankton blooms (UIBs) have increasingly been observed in the Arctic, driven by changes to the optical properties of Arctic sea ice as it has thinned and become more seasonal. Here we show the Southern Ocean can also support widespread UIBs. Using under ice-enabled BGC-Argo float data, we detail numerous high chlorophyll and particulate backscatter events below compact sea ice preceding seasonal ice retreat, classifying 16 distinct UIB events. Using joint light, sea ice, and ocean conditions obtained from the ICESat-2 laser altimeter and 11 climate model contributions to CMIP6, we find that more than 4 million square kilometers of the compact-ice-covered Southern Ocean could support these events in late spring and early

#### 4 1 Introduction

Observations of under-ice phytoplankton blooms (UIBs) in the Arctic Ocean [1] have high-lighted the need to understand ecological communities living under compact (local concentration greater than 80%) sea ice, now and under future climate change scenarios [2, and references within]. Regions supporting UIBs in the Arctic have likely expanded as sea ice has thinned and become more seasonal. Yet to date, no studies have quantified the potential for widespread UIBs under Antarctic sea ice, where annual and seasonal variability has changed less than in the Arctic over the satellite period [3] and where sea ice is typically thinner, more seasonal, and more fragmented.

Antarctic sea ice typically has a higher albedo than Arctic sea ice [4, 5]. Thus a limited 23 amount of photosynthetically available radiation (PAR, 400-700 nm) can reach the upper ocean 24 directly through sea ice, especially compared to the Arctic, where light transmission through 25 melt-pond-covered sea ice is thought to be a primary cause of UIBs [6]. Still, spring-summer 26 solar irradiance is high: recently a bloom of nanoflagellates was observed under highly reflective landfast sea ice [7]. Floating sea ice in the Southern Ocean is also fractured, thin, and mobile. 28 Small areas of open water, like leads or small openings within the floe mosaic, can allow substantial amounts of light to reach the upper ocean. Sunlight entering the ocean through leads in the Arctic has been shown to initiate phytoplankton blooms, even in areas where sea ice is thick and snow-covered [8]. We therefore investigate this possibility across the sea-ice-covered 32 Southern Ocean.

Phytoplankton communities in the Southern Ocean respond rapidly to changes in light conditions, with phytoplankton blooms often observed as soon as the sea ice edge retreats in spring, flooding the mixed layer with light and leaving freshwater rich in iron, main limiters of primary production [9, 10, 11]. In the Arctic, a crucial factor in the development of UIBs is a stable surface mixed layer, which can be induced by melt water and/or increased solar heating of the surface layer [12, 13]. Observations using tagged seals in the Ross Sea show the initiation of a shallow (20 m) surface mixed layer driven by ice melt, preceding the seasonal retreat of sea ice [14]. Yet while shallow mixed layer depths may be necessary to keep phytoplankton in the well-lit surface layer, observations from under-ice Argo floats [15, 16] demonstrate that primary production can be initiated before seasonal sea ice retreat, and even before the restratification of surface waters. This challenges the notion that too-deep surface mixed layers in ice-covered regions of the Southern Ocean limit productivity. Together these factors present the possibility that non-coastal regions of the Southern Ocean, like the Arctic, may be productive before sea ice retreats in summer.

Here we assess the potential for widespread phytoplankton growth under compact (and floating) sea ice in the Southern Ocean. We examine 39 BGC-Argo floats that operated under sea ice in the Southern Ocean, 30 of which returned profiles under compact sea ice in austral springsummer. Of these 30 floats, 27 recorded sub-surface Chl-a maxima that exceeded 0.1 mg/m<sup>3</sup>. A 51 further 12 recorded Chl-a values that exceeded 0.5 mg/m<sup>3</sup>, which we define as a UIB, a threshold used to define UIBs in the seasonally ice-covered Arctic [17, 18, 19]. In total, we observed 28 UIB profiles in 16 distinct locations and time periods. To evaluate the general ability of the ice-covered Southern Ocean to permit UIBs, we combine ICESat-2 laser altimetry and 11 climate model assessments of Southern Ocean sea ice, light and ocean conditions, finding the conditions required for light-limited phytoplankton blooms are predicted across nearly 50% of regions with compact ice in spring and summer. These results suggest that in compact, but not completely ice-covered regions of the Southern Ocean, enough light reaches into the upper water column to permit primary production, as found in the Arctic [8]. We identify potential sampling regions for examining under-ice primary production and community composition in the Ross Sea, and discuss the implications for sampling strategies and cruise timing.

# Observations of phytoplankton blooms under compact sea ice

Figure 1(a) scatters the maximum Chl-a measurement for BGC-Argo profiles ( $Chl_{max}$ , see methods) in the months of September to December when local sea ice concentration (SIC) exceeds 80%, overlaid on September-November climatological SIC. Due to the east of interpretation, we define high-productivity measurements using  $Chl_{max}$ , however Chl-a is an imperfect measure of phytoplankton biomass because it varies due to photoacclimation, nutrient conditions and growth stage, and several studies have identified uncertainty with Argo float fluorometers [20, 21]. For consistency with previous BGC-Argo work [22], and to support  $Chl_{max}$  observations, we compare  $Chl_{max}$  measurements with particulate backscattering data ( $b_{bp}$ , at 700nm, units m<sup>-1</sup>) taken at the depth of maximum Chl-a. We found comparable seasonal cycles under compact ice and a high association (Spearman's R=0.7) between the two (see Supporting Information, Fig. S2). Example profiles of Chl-a and  $b_{bp}$  are provided as Supporting Figure S4 with  $Chl_{max}$  varying from 0.1 to 3.5 mg/m³, showing the typical covariance of  $b_{bp}$  with Chl-a. The depth of Chl-a maxima are predominantly within the top 60 meters of the water column (Supporting Figure S3), implying the association between high Chl-a values and high particle concentrations is not due to photoacclimation.

In total, we considered 39 BGC-Argo floats that recorded 1153 profiles under sea ice in the Southern Ocean, 903 of which were under compact sea ice. 33 floats recorded 549 profiles during the period from September-December, of which 30 recorded 426 profiles under compact sea ice. Of these 30, 27 recorded Chl<sub>max</sub> greater than 0.1 mg/m³, indicating some photosynthetic activity, and 12 record Chl<sub>max</sub> greater than 0.5 mg/m³ at least once, a threshold we take to represent initiation of a under-ice "bloom". We note that we define blooms here only in a relative sense to background concentrations in the Southern Ocean (i.e, not relative to the North Atlantic

bloom) [20, 23]. The three floats recording profiles under compact ice in spring-summer that did not observe a value of  $Chl_{max}$  above 0.1 mg/m<sup>3</sup> accounted for just 14 total profiles, with just one under compact sea ice.

Many of the highest  $Chl_{max}$  values were recorded under compact sea ice. Considering 549 under-ice profiles taken between the months of September-December, we show box plots of  $Chl_{max}$  in Figure 1(b), grouped by sea ice concentration (SIC, in 20% bin intervals). A total of 52 profiles under sea ice recorded  $Chl_{max}$  above 0.5 mg/m<sup>3</sup>: 11 for SIC from 20%-40%, 6 for SIC from 40% to 60%, 13 for SIC from 60% to 80%, and 28 for SIC from 80% to 100% (compact ice). Median  $Chl_{max}$  values are below 0.2 mg/m<sup>3</sup> in each SIC category, with the median for SIC from 20%-40% (0.15 mg/m<sup>3</sup>, n=31) similar to that for SIC from 80-100% (0.12 mg/m<sup>3</sup>, n=426).

Box plots of Chl<sub>max</sub> in each month are given in Figure 1(c) for the 903 total profiles under compact sea ice. The seasonal cycle in median  $Chl_{max}$ , matches the seasonal cycle of downwelling solar irradiance. Median  $Chl_{max}$  under compact ice ranged from 0.03 mg/m<sup>3</sup> (n=163) 100 in August, to a high of 0.38 mg/m<sup>3</sup> in December (n=24). The number of recorded Chl<sub>max</sub> values 101 exceeding 0.5 mg/m<sup>3</sup> was 0 through September, 3 out of 149 profiles in October, 17 out of 99 102 profiles in November, and 8 out of 24 profiles in December. These 28 "bloom" Chl-a measure-103 ments averaged 1.34 mg/m<sup>3</sup>, were recorded by 12 different floats, and were under an average SIC of 94%. As BGC-Argo float records can span multiple years, these "bloom" measurements were recorded in 16 distinct times and locations. We found 7 instances where at least 2 suc-106 cessive profiles (Argo dives are spaced 10 days apart) recorded both  $Chl_{max} > 0.5 \text{ mg/m}^3$  and SIC>80%. Of the 7 "events" spanning 10 or more days, 6 included profiles taken in November - the other spanned only the month of December.

## 10 3 The prevalence of blooms under Antarctic sea ice

The presented BGC-Argo float data showed numerous elevated Chl-a events under compact sea ice in the Southern Ocean, and we found 16 distinct instances of Chl-a concentrations exceeding 0.5 mg/m³ that were recorded under sea ice with an average SIC of 94%. Many of these high-Chlorophyll events were recorded in November, when Antarctic sea ice is near its seasonal maximum extent. We next quantify if conditions that support UIBs are widespread across the sea-ice covered Southern Ocean before sea ice retreat.

In Figure 2(a), we show ICESat-2-(IS2)-derived average ocean surface PAR values in the 117 Southern Ocean in November, in which we make the simplifying assumption that no PAR reaches the upper ocean directly through sea ice (see methods). A solid line outlines the com-119 pact sea ice zone (CIZ, SIC > 80%) defined using the NSIDC-CDR SIC product [24]. We also 120 plot the 15% SIC contour, marking the edge of total sea ice extent (SIE). Regions lying inside 121 the SIE contour but outside the CIZ are defined as marginal ice zones (MIZs), which due to 122 the lower albedo of open water, receive higher PAR in the surface water layer compared to the 123 CIZ. Figure 2(b) shows pre-industrial November PAR values for the CESM2 climate model 124 (see methods), with CIZ and MIZ defined from the CESM2 model climatology. Both IS-2 and 125 CESM2 show large areas within the CIZ where ocean surface PAR estimates exceed a "bloom" 126 threshod of 23  $\mu$ mol photons/m<sup>2</sup>/s (see methods). For the IS-2 estimate of ocean surface PAR, 127 6.9 million km<sup>2</sup> of the November CIZ exceeds that PAR threshold, versus 5.9 million km<sup>2</sup> for 128 CESM2. Because we do not have coincident ocean and sea ice melt observations at the scale of 129 IS2 observations, IS2 estimates only indicate the presence of light in the upper ocean and may 130 overestimate the area that permits an UIB. 131

We next consider how frequently an individual grid cell would permit an UIB (see methods), a metric we term the UIB%. A spatial map of UIB% in November months is given in Figure 2(c)

for CESM2. Areas within the climatological November CIZ (solid line), which has an area of 8.3 million km<sup>2</sup>, permit an UIB 46.4% of the time. Because of year-to-year variability of the CIZ contour, areas outside of the climatological CIZ also have non-zero UIB%. In those areas, average UIB% is 19.3%. We accumulate climatological statistics of UIB-permitting regions in Figure 2(d), comparing the climatological extent of compact sea ice (red) to the extent of UIBpermitting regions (blue). Large areas support UIBs, peaking at 5.9 million km<sup>2</sup> of compact ice-covered regions in November. The fraction of the CIZ that permits an UIB, the UIB fraction 140 (UIBF), is examined in Figure 2(d) (black line, right axis), which peaks in November at an UIBF 141 of 77%. By point of comparison, we reproduce Figure 2(d) as Figure 2(e) for the Arctic Ocean. Up to 4.3 million km<sup>2</sup> of the pre-industrial Arctic CIZ is permissive to UIBs, repeating the finding in [2], that large regions of the pre-industrial Arctic also supported UIBs. The seasonal maximum of Arctic UIB area occurs in June, at the peak of the solar cycle, with a peak UIBF 145 of 52% in July. Generally, in the CESM2 picontrol experiments, we find that UIB-permitting regions in the Antarctic are (1) larger, (2) constitute a larger percentage of the CIZ, and (3) peak earlier in the annual solar cycle (November in the Antarctic versus June in the Arctic) than in the Arctic.

#### 50 Southern ocean UIB statistics across CMIP6 models

In Figure 3(a,b), we plot the climatological seasonal cycle of Southern Ocean UIB area (a) and UIBF (b) for the 11 CMIP6 models (listed in Supporting Table S2). Across these models, we find a similar seasonal cycle. None of the CMIP6 models have large UIB areas before October, but 10 of 11 have a maximum UIB area in November. Only the MRI-ESM2 model shows a maximum UIB area in December. Each has a climatological UIB area exceeding 2.66 million km², with a median of 4.75 million km². In Figure 3(c), we show box plots of annual maximum UIB area in the Antarctic for each of the models (filled), compared to annual maximum UIB

area in the Arctic (unfilled) for the same years. Out of 11 models, 8 have median Antarctic UIB areas that exceed Arctic UIB areas.

We repeat Figure 3(a,c) in Figure 3(b,d) for the UIBF, with Figure 3(d) showing UIBF values 160 during the month where UIB area is at its maximum (November or December in the Antarctic, 161 June or July in the Arctic). Seasonal cycles of UIBF are similar between models, with most 162 models peaking in December as the CIZ reduces in extent and ocean surface PAR increases. In 163 10 of 11 models, a higher fraction of the Antarctic CIZ permits an UIB than of the Arctic CIZ. 164 Average values of UIBF range from 27-86% (average 57%) in the Antarctic, compared to 26-165 66% in the Arctic (average 37%). Each of the three models in which Antarctic UIB areas were 166 less than Arctic UIB areas have higher UIBF in the Antarctic. Thus we suggest that the reason 167 for differences in the overall magnitude of Antarctic UIB areas is due to differences in model 168 representations of Antarctic and Arctic sea ice, not disagreements about whether sufficient PAR 169 is available under the compact sea ice there. 170

#### 4 The potential for observing UIBs in situ

171

Using BGC-Argo float data, we have demonstrated that high phytoplankton biomass events exist under compact sea ice in the Southern Hemisphere, preceding the seasonal loss of sea ice by several months. Examining a series of climate model estimates of upper ocean light and sea ice conditions, we found that under-ice phytoplankton growth is permitted across wide swaths of the compact ice-covered Southern Ocean. We also found that areas permitting UIBs make up a larger percentage of compact sea ice zones in the Southern Ocean than the Arctic, with an earlier peak in the seasonal cycle. To validate these results further, observational campaigns will be needed. We specifically focus on the potential of the Ross Sea region (see methods) to support such an event, as it is seasonally ice free, and is among the highest-productivity regions of the Southern Ocean, and is known for supporting large ice-algal communities [25, 26].

In Figure 4, we plot UIB% for each of the 11 models during the model period with highest 182 Ross Sea UIB area, which is November in 7 models and December in 4 models. All models have high UIB% in the coastal region near Cape Adare in the Western Ross Sea, which has 184 compact sea ice into January. We identify a region at 72°S, 178.5°E with a blue square in 185 Figure 4. A box plot of UIB% in this location for these 11 models is given in Figure 4 (bottom 186 right), showing a median UIB% there of 62% with a minimum of 40%. Across the CMIP6 187 models, a mean area of 0.55 million km<sup>2</sup> of the Ross Sea is expected to permit UIBs, although 188 the borders of UIB-permitting areas vary by model, and range from 0.29 million km<sup>2</sup> (MRI-189 ESM2, 49% of the Ross CIZ) to 0.95 million km<sup>2</sup> (NorESM2-LM, 65% of the Ross CIZ), with 190 inter-annual variability. Independent of modeled sea-ice area coverage, a large fraction of the 191 Ross CIZ permits UIBs in each year in all models. Figure 3 is repeated as Supporting Figure S1 192 for the Ross Sea region, showing that during the month of highest Ross Sea UIB area, at least 193 49% of the Ross CIZ permits UIBs in each model, on average 60%.

#### 5 Discussion

Here we explored the potential for under-ice phytoplankton blooms beneath compact sea ice in the Southern Ocean using model simulations, altimetric measurements of sea ice coverage, 197 and BCG-Argo data. We show that on at least 16 distinct occasions from October-December, 198 BCG-Argo floats recorded Chl-a concentrations above 0.5 mg/m<sup>3</sup> in areas with an average sea ice concentration of 94%. In addition to these direct high Chl-a measurements, findings of elevated (Chl<sub>max</sub> > 0.1 mg/m<sup>3</sup>) Chl-a concentrations in a large fraction of analyzed float data 201 demonstrates the likelihood for primary production and blooms beneath Antarctic sea ice pre-202 dating seasonal sea ice retreat. These elevated levels of  $Chl_{max}$  under compact ice suggest that 203 even for areas with low open water fraction, incident solar radiation is high enough to pro-204 mote photosynthetic activity. This is similar to findings in the Arctic where small lead features were sufficient to support under-ice blooming [8], and suggests that small regions of open water are sufficient to relax light limitations on blooming in the summertime Southern Ocean. We supplemented the Chl-a observations with observations of  $b_{max}$ , values of particulate backscatter  $b_{bp}$  taken at the same depth as  $Chl_{max}$ . We find high association between Chl-a and  $b_{max}$ (Spearman's R = 0.7, see Supporting Info Fig. S2), reinforcing that these high Chl-a events are associated with higher particle (i.e., biomass) concentration in the water column.

We further used ICESat-2 and an ensemble of climate model estimates of sea ice, light, and 212 oceanographic conditions across the compact-ice-covered Southern Ocean to show that indeed, 213 conditions are favorable for under-ice blooms over wide regions, with a median estimate UIB 214 area of 4.75 million km<sup>2</sup> across the model ensemble. In using ICESat-2 data, we assumed that 215 no light reached through sea ice, and all light available for photosynthesis came through open 216 water regions near compact ice. Thus these findings indicate that even in regions with local sea 217 ice concentrations above 80%, and with no light passing directly through the ice, enough open 218 water exists that light does not limit growth in the upper Southern Ocean [16]. As modeled and 219 observed in the Arctic, such widespread under-ice productivity, preceding the retreat of seasonal 220 sea ice, may reveal a markedly different ecological system than was previously understood 221 to exist there. Since sea ice conditions in the Antarctic have not changed dramatically over 222 the global warming period, under-ice phytoplankton may play an important role in driving or 223 limiting blooms observed to occur when sea ice retreats. 224

The climate models considered here have inter-related sea ice and light schemes (see Supporting Table S2), and provide estimates of the light conditions in the Southern Ocean. They may not be accurate if systematic biases in modeled Southern Ocean climate or sea ice properties exist. Still, compared against the estimate of upper ocean PAR derived from ICESat-2 data, models produce similar PAR estimates and areas of high surface light levels. We adopted a simple diagnostic criteria for when sufficient light is available to support a bloom, using a fixed

225

227

228

229

PAR threshold in model data, in line with Arctic model studies and observations of acclimation in key Antarctic phytoplankton species [27]. While some BGC-Argo floats do report PAR 232 values, their ice-avoidant algorithms mean near-surface PAR was not observed - and further ob-233 servations and modeling of radiative transfer of PAR specifically focused on variable Antarctic 234 sea ice (as in, for example, [28, 29]) would help constrain and evaluate PAR levels needed to 235 trigger blooms in concert with BGC-Argo data. While the Argo data confirms that under-ice 236 regions can be productive, because of their uneven spatial and temporal coverage (Supporting 237 Info Fig. S3), it is not yet possible to directly compare geographic estimates of model-predicted 238 conditions to float data. Thus here we use the Argo data in conjunction with models for under-239 standing the possibility of under-ice blooms, but cannot directly validate model predictions of 240 pan-Antarctic UIB extents. 241

The work we presented here raises an important question: if conditions beneath compact sea ice are favorable for supporting UIBs, and Antarctic sea ice coverage and downwelling irradiance has remained largely stable over the past several decades, why are there no reported observations of under-ice blooms in the Southern Ocean by underway cruises or moorings? We suggest two potential answers.

First, the detection of UIBs requires a dedicated effort to collect in situ chlorophyll data under compact sea ice. An analogy can be drawn to the Arctic Ocean, where spring-summer icebreaker research expeditions are more common. UIBs are now thought to have been widespread
dating back to at least the 1950s (with an overall area coverage that has doubled since 1970 [2]).

But these phenomena, which can have some of the highest levels of integrated biomass of any
ecological system [1], were rarely observed before the report of a massive under-ice bloom
in the Chukchi Sea in 2011. As we show here, BGC-Argo floats permit a broader sampling
of biological parameters across the Southern Hemisphere using consistent methodologies and
calibrations. Mining of existing and previous under-ice Chlorophyll data, for example from the

BCO-DMO archive, will be a focus of future work to understand whether such events have been observed in the past.

Second, it is possible that UIBs do not occur regularly. While nearly all BGC-Argo floats 258 operating from September-December show elevated Chl-a measurements predating sea ice re-259 treat, only 28 profiles, from 12 of 30 floats operating in the right time and sea ice condi-260 tions, exceed our defined Chl-a threshold for a "bloom", which here is defined relative to 261 typical open water and under-ice conditions in the Southern Ocean [20, 23]. Of these, we 262 recorded 7 events with two or more subsequent measurements meeting the "bloom" threshold. 263 These multi-measurement events occurred in December 2014 (Argo id 5904180), November-264 December 2016 (Argo id 5904767 and 5904180, which was previously discussed in [30]), 265 October-November 2017 (Argo ids 5905100 and 5904180), November 2019 (Argo id 5905379), 266 and November-December 2018 (Argo id 5905636). Further research into these specific events 267 will be necessary to rule out that the high levels of Chl-a recorded by these floats were not, 268 for example, advected from a bloom occurring in open water. Our assessment of favorable un-269 derwater light conditions for UIBs over large areas of the compact ice zone is also based on a 270 simple set of diagnostic criteria, not detailed biogeochemical modeling, and uses bulk estimates 271 for light transmission and stratification. We do not take into account iron or other nutrient limi-272 tations, nor grazing pressure by higher trophic levels. Instead we follow the perspective of [11] 273 that primary production is primarily light-limited in summer. Thus the UIB-permitting area of 274 the Southern Ocean in this study is likely an upper bound on the areas that might bloom in a given year.

This work suggests there is potentially unexplored ecological variability beneath Southern Ocean sea ice, with several million square kilometers of the ice-covered Southern Ocean potentially permitting blooms before the seasonal retreat of the sea ice edge. We paid special attention to the frequently visited Ross Sea region, and suggest detailed measurements

of physical and biogeochemical variables to study under-ice phytoplankton bloom phenology, 281 magnitude and community composition and to compare those to known bloom dynamics in the 282 Arctic Ocean [31]. Sampling during the sea ice-covered season will be challenging, especially 283 as remote sensing technologies presently cannot measure chlorophyll under sea ice. Continued 284 targeted deployment of Remotely Operated Vehicles (ROV) and other autonomous profiling 285 floats [15, 16] to measure under-ice light availability and bio-optical parameters can be comple-286 mentary to ship-based sampling, supported by ICESat-2 measurements used to remotely sense 287 particulate backscatter in ice-free conditions [32] extended into sea-ice-covered regions. 288

#### 6 Methods

306

307

308

309

310

**Ice-enabled BGC-Argo float data** Autonomous profiling Argo floats equipped with biological sensors are a foundation of Southern Ocean biogeochemical observations because they 291 provide observations with consistent sampling methodologies in places (and at sampling fre-292 quencies) inaccessible via ships, and with depth resolution inaccessible via satellite, while ex-293 periencing minimal biofouling and lateral drift [33]. Because Argo floats drift with the currents 294 during their transit, a portion of floats deployed in open water drift into ice-covered regions. To protect the floats from ice damage, an ice-avoidant algorithm (based on a temperature threshold) 296 was implemented to initiate a float's descent when it encounters near freezing surface waters 297 [34]. We used Chl-a flourescence data and particulate backscattering data  $b_{bp}$  (700 nm) col-298 lected with biogeochemical Argo floats (BGC-Argo) [35], which have been shown to be consis-299 tent with both satellite-derived Chl-a and in situ Chl-a values measured via High Performance 300 Liquid Chromatography in the Southern Ocean [20, 21] following bias and non-photochemical 301 quenching corrections [36]. In this study we use quality controlled and calibrated [21] delayed-302 mode data distributed through SOCCOM. Parameters for the floats are sampled at 2m vertical 303 resolution. We did not impose geographical constraints on the data, and required only that float 304 data come from compact (SIC > 80%) regions in the Southern Ocean. 305

Chl-a observations are supported by particle backscatter ( $b_{bp}$ ) data because while Chl-a is a pigment common to all phytoplankton, its observed value varies can vary based on both phytoplankton physiology and measurement error. Particulate backscattering covaries with phytoplankton biomass as phytoplankton scatter light proportional to their concentration and size [37], although  $b_{bp}$  observations do not necessarily imply the presence of phytoplankton because  $b_{bp}$  can be elevated due to the presence of non-algal particles. Backscatter has been shown to be a better proxy for phytoplankton carbon compared to Chl-a ([38] with Argo floats having less

measurement uncertainty for  $b_{bp}$  (on the order of 15%), [39] than for Argo Chl-a.

As in [39, 40], all float profiles of Chl-a and  $b_{bp}$  (700 nm, m<sup>-1</sup>) were despiked with a 3-point 314 moving median and we visually confirmed that maximum Chl-a values were not from noise in 315 the profile. We report the maximum adjusted Chl-a,  $Chl_{max}$ , within a profile due to unavailable 316 profile data near the surface (float descended prior to reaching the surface due to the ice avoidant 317 algorithm, see also [41]). Examples of four Chl-a and  $b_{bp}$  profiles are given in Supporting Figure 318 S4. We also examine values of  $b_{max}$ , the value of  $b_{bp}$  at the depth of  $Chl_{max}$ , to confirm that the 319 high Chl-a is due to phytoplankton, with Fig. 1(a) repeated as Fig. S2(b) for  $b_{max}$ . Profiles of 320 Chl-a are not considered if  $Chl_{max}$  is recorded at a depth above 200 meters, or if  $b_{max}$  exceeds 321  $0.01 \text{ m}^{-1}$ , which exceeds natural values of  $b_{bp}$  found in phytoplankton and possibly reflects the 322 influence of bubbles or large particles (zooplankton) that were attracted to the instrumentation 323 [42]. 324

Maximum values of Chl-a reported here may represent an underestimate of the true maxi-325 mum Chl-a in the water column, as it is not possible to assess Chl-a concentrations closer to the 326 surface under sea ice because of the ice-avoidant nature of the Argo floats, and typically ocean 327 phytoplankton blooms are surface-intensified [1, 43]. For example, the mean depth of  $Chl_{max}$ 328 for the 28 profiles with  $Chl_{max}$  above 0.5 mg/m<sup>3</sup> under compact ice was 45 meters. Location 329 information for a float under sea ice is imprecise, as the latitude and longitude coordinates are 330 calculated via a linear interpolation of the pre- and post-sea ice coordinates of a specific float. 331 In some cases, the float will not surface in open water and post-sea ice coordinates are unavailable. Some floats lack under-ice geographic coordinates if they do not surface under open water 333 conditions following a period under ice. Thus it is not possible to interpolate all float location 334 while it transits an ice-covered region, and we remove such floats/dives. 335

This study includes under-ice profiles initially obtained from 41 BGC-Argo floats. To obtain sea ice concentrations (SIC) in the area of float deployment, we matched geographic coordinates

336

for each float to the daily 25-km resolution NSIDC Climate data record SIC product [24]. We excluded locations with a local estimated SIC below 15%, and outlier profiles where  $b_{bp}$  at the depth of maximum Chl-a exceeded 0.01 m<sup>-1</sup>. In total, that leaves 1153 profiles under sea ice from 39 floats. Of all profiles, 35 floats recorded 903 profiles with local SIC above 80% in all months. Of all profiles, 33 floats recorded 549 profiles during the months of September-December. Combining these criteria, 30 floats recorded 426 profiles with local SIC above 80% between September and December. A list of all floats is provided in the Supporting Information Table S2.

CMIP6 model data Remote sensing technologies presently do not directly measure light or chlorophyll beneath sea ice, and most sampling strategies for Southern Ocean photosynthetic communities associated with sea ice focus on in-ice algae communities in coastal regions [44, 45, 46, 47]. We must instead turn to model estimates to describe the joint climatological light, sea ice, and ocean conditions underneath sea ice. We used an ensemble of current-generation coupled climate models contributing to the 6th Coupled Model Intercomparison Project (CMIP6).

While observations show Antarctic sea ice has been stable or increased in extent over the 353 satellite period (1978-present), CMIP6 models consistently simulate a declining annual-average 354 Antarctic sea-ice cover over this period [48]. Thus we did not consider it feasible to examine 355 present-day model estimates of Antarctic sea ice state, which might incorporate biased depic-356 tions of sea ice albedo and extent. Instead we postulate that light conditions under Antarctic sea 357 ice have remained stable over the industrial period, and use data from pre-industrial control run 358 simulations (CMIP6 runs titled picontrol) in this analysis. Of the full CMIP6 model dataset, 11 359 simulations (see Supporting Table S1) submitted the required model output we used here. 360

The ensemble of 11 models produced variable estimates of climate and sea ice state, de-

361

spite high interrelation between their sea ice and radiative transfer model components. Different versions of the Community Sea Ice Model (CICE) are used as sea ice model components in 9 of 11 models. There are three substantively different light models, the improved [49] (B+L)  $\delta$ -Eddington multiple-scattering scheme found in CICE versions 5 and above (CESM2 and NorESM2 simulations), an earlier version of the B+L scheme found in CICE version 4 (CAS), or implementations of simpler Beer-Lambert exponential attenuation of light in ice (CERFACS,MRI).

For each CMIP6 model, we defined a climatology of light and sea ice properties using the final 100 years of their respective pre-industrial spinup experiments. In Fig. 2 we specifically examined the Community Earth System Model version 2 (CESM2, [50]) model run, as it uses the more recent version of CICE and the more advanced B+L  $\delta$ -Eddington light scheme. CESM2 produces an overall mean state of Antarctic sea ice that is broadly realistic compared to other CMIP6 models [48, 51], and similar output from CESM2 was analyzed to evaluate the potential for Arctic UIBs in [2].

To supplement model estimates of light under sea ice, we approximated ICESat-2 light data 376 the light field under sea ice using the ICESat-2 laser altimeter. We utilized the L3A along-track 377 sea ice type product (ATL07, [52]) derived from Level 2A ATL03 photon heights [53]. Sea ice 378 types are determined using an empirical decision tree, which identifies whether a given segment 379 is sea ice or water. We developed an estimate of SIC as the ratio of total ice segment length to 380 total segment length. This quantity, the linear concentration  $c^*$ , is related to the SIC, defined 381 over a 2-dimensional region. Given the random orientation of crack and open water features 382 relative to frequent satellite tracks, many repeat 1-D measurements can approximate a 2-D field 383 when sampled sufficiently. In [54], we found global sea ice area metrics derived from passive 384 microwave (PM) satellites were well-approximated by this method in regions where IS2 records at least 1000 individual segments per month. We adopted this same threshold in this study to define  $c^*$ . An advantage of using ICESat-2 segments instead of PM is that ICESat-2 is capable of resolving small cracks and leads that are difficult to observe in PM estimates of local SIC, particularly in summer [55, 56, 57].

From a gridded dataset of  $c^*$ , we estimated the total shortwave irradiance,  $I_0$  ( $\sim$  300–3000 nm), reaching the upper ocean,  $I_0$ ,

$$I_0 = SW(1 - c^*)(1 - \alpha_{oc}) \tag{1}$$

where  $\alpha_{oc} = 0.06$  is the open water albedo and SW is the downwelling solar irradiance at 392 the surface. This shortwave irradiance is then converted to a PAR (400-700 nm) estimate as 393 in the CMIP6 model data (see methods below). This simple model assumes no light passes 394 through the sea ice surface, and the only light available in ice-covered regions comes through 395 the open water part of the area. For this reason we expect ICESat-2 derived downwelling 396 irradiances may be conservative. For SW, we use the reanalyzed estimate of downwelling 397 shortwave irradiance from [58]. We use IS2 data from January 2019-December 2020 to form 398 the present-day climatology of  $I_0$  that is presented in Figure 2. 399

- Criteria for permitting an UIB We define an area as "permitting" an under-ice bloom if it
  meets three criteria:
- 402 **Compact sea ice** Local sea ice concentration exceeds 80%.
- An illuminated upper ocean. Average PAR in the top 25 meters of the ocean exceeds  $10 \mu mol$  photons/m<sup>2</sup>/s.
- A stable or stratifying surface mixed layer. Sea ice is not refreezing and the upper ocean is non-convecting.

The UIB% is therefore defined as the percentage of model years where a grid cell meets all three criteria together. Thus the UIB% can be low if a region both if it is not frequently covered by compact sea ice, or if the light conditions and ocean stratification are not permissive of a bloom.

We focused on those ocean regions under "compact" sea ice to differentiate from phytoplankton growth known to occur as the ice edge retreats in marginal ice zones [59, 60]. Marginal ice zones are typically defined as areas where sea ice concentration is less than 80% [61, e.g.,], thus we used this cutoff to define regions that are "compact" ice.

To establish a threshold for upper-ocean PAR, we estimated average PAR,  $\overline{I}$ , at a depth D as,

$$\overline{I} = \frac{I_0}{\kappa D} \left[ 1 - \exp(-\kappa D) \right]. \tag{2}$$

Here we assumed that PAR is attenuated exponentially in water with a coefficient  $\kappa$ . We assumed positive photosynthesis (gains outweigh losses) occurs when the average PAR over a 418 25-m deep water column exceeds 10  $\mu$ mol photons / m<sup>2</sup>/s. This is approximately twice the threshold of integrated daily irradiance of 4.8  $\mu$ mol photons / m<sup>2</sup>/s considered to initiate a 420 phytoplankton bloom in [62, 63, 13], and higher than the levels found to initiate growth in the 421 Southern Ocean [15, 16]. Using  $\kappa = 0.081/m$  [64] for PAR extinction in clear waters and 422 D=25 m established a PAR threshold value for blooms of  $I_0^*\approx 23~\mu \text{mol}$  photons /m<sup>2</sup>/s. 423 CMIP6 models typically store and output full-spectrum solar forcing to the upper ocean, but 424 not PAR. We therefore had to convert full spectrum solar irradiance to PAR using a factor of 425 1.9975  $\mu$ mol photons/J [65, 64]. 426 We also included a threshold for the termination of upper-ocean convection. Under-ice 427 blooms are unlikely to occur when active convection extends below the euphotic zone, such as 428 when leads are actively refreezing with the ocean at its freezing point [12]. The requirement 429

that the upper ocean is non-convecting is similar to the "turbulent shutdown" theory used to

explain mid-latitude phytoplankton blooms [66]. GCMs used here are too coarse to resolve 431 the complex boundary layer dynamics that result from surface melting of sea ice [67, 68, 69], 432 and thus they are not suited for determining the convective state of the upper ocean in the 433 presence of sea ice leads. Instead, we considered the ocean to be non-convecting if sea ice was 434 melting at its base, which would lead to stratification of the upper ocean, consistent with Argo 435 observations of high negative covariance between shoaling MLD and phytoplankton biomass 436 under ice [41]. In practice, simply non-zero basal melting does not restrict the location of UIBs 437 as small monthly-averaged basal melt rates occur whenever sea ice is present. We therefore set 438 a positive threshold for the sea ice basal melt rate  $\dot{h}$ , which we expressed as an equivalent heat 439 flux  $Q = \rho_i L_f \dot{h}$ , with  $\rho_i = 920 \text{ kg/m}^3$  the sea ice density and  $L_f = 3.34 \times 10^{-5} \text{ J/kg}$  the latent 440 heat of fusion. As a result Q is required to exceed 5 W/m<sup>2</sup>, for an approximate basal melt rate of  $\dot{h} = 5$  cm/month.

The Ross Sea region To define the "Ross Sea region", we roughly followed the convention established by the NIWA Ross Sea Trophic Model [70], taking the ocean region south of 69°S and between 160°W and 170°E longitude. Because of grid variations, the area of this region can vary between CMIP6 models, but its surface area is approximately 1.5 million km<sup>2</sup>.

#### References

- [1] Arrigo, K. R. et al. Massive Phytoplankton Blooms Un-448 Sea 1408 der Arctic Ice. Science 336, (2012).**URL** 449 http://www.sciencemag.org/cgi/doi/10.1126/science.1215065. 450 9605103. 451
- [2] Ardyna, M. *et al.* Under-Ice Phytoplankton Blooms: Shedding Light on the "Invisible" Part of Arctic Primary Production. *Frontiers in Marine Science* **7**, 1–25 (2020). URL

- https://www.frontiersin.org/articles/10.3389/fmars.2020.608032/full.
- Parkinson, C. L. A 40-y record reveals gradual Antarctic sea ice increases followed by decreases at rates far exceeding the rates seen in the Arctic. *Proceedings of the National Academy of Sciences of the United States of America* **116**, 14414–14423 (2019).
- 458 [4] Brandt, R. E., Warren, S. G., Worby, A. P. & Grenfell, T. C. Surface Albedo
  459 of the Antarctic Sea Ice Zone. *Journal of Climate* 18, 3606-3622 (2005). URL
  460 https://journals.ametsoc.org/jcli/article/18/17/3606/30648/Surface-Albe
- mission through Antarctic pack ice. Journal of Geophysical Research: Oceans

  122, 2108–2119 (2017). URL http://www.nature.com/articles/175238c0

  http://doi.wiley.com/10.1002/2016JC012325.
- [6] Arrigo, K. R. et al. Phytoplankton blooms beneath the sea ice in the Chukchi sea.

  Deep-Sea Research Part II: Topical Studies in Oceanography 105, 1–16 (2014). URL

  http://linkinghub.elsevier.com/retrieve/pii/S0967064514000836.
- for Item 15 Saggiomo, M., Escalera, L., Saggiomo, V., Bolinesi, F. & Mangoni, O. Phytoplankton Blooms Below the Antarctic Landfast Ice During the Melt Season Between Late Spring and Early Summer. *Journal of Phycology* 57, 541–550 (2021). URL https://onlinelibrary.wiley.com/doi/10.1111/jpy.13112.
- [8] Assmy, P. et al. Leads in Arctic pack ice enable early phytoplankton blooms below snow-covered sea ice. Scientific Reports 7, 40850 (2017). URL http://www.nature.com/articles/srep40850.

- 475 [9] Martin, J. H., Fitzwater, S. E. & Gordon, R. M. Iron deficiency limits phytoplank-476 ton growth in Antarctic waters. *Global Biogeochemical Cycles* **4**, 5–12 (1990). URL 477 http://doi.wiley.com/10.1029/GB004i001p00005.
- [10] Comiso, J. C., McClain, C. R., Sullivan, C. W., Ryan, J. P. & Leonard, C. L. Coastal
   zone color scanner pigment concentrations in the Southern Ocean and relationships to
   geophysical surface features. *Journal of Geophysical Research: Oceans* 98, 2419–2451
   (1993). URL http://doi.wiley.com/10.1029/92JC02505.
- Oijen, T. Light rather than iron controls photosynthate production [11] van 482 and allocation Southern Ocean phytoplankton populations during aus-483 tral autumn. Journal of Plankton Research 26, 885–900 (2004). **URL** 484 https://academic.oup.com/plankt/article-lookup/doi/10.1093/plankt/fbh08 485
- Inhibited by Spring Convective Leads. Leads. Journal of Geophysical Research: Oceans 123, 90–109
   URL http://doi.wiley.com/10.1002/2016JC012575.
- [13] Oziel, influencing the L. al.Environmental factors seasonal dy-489 namics of spring algal blooms in and beneath sea ice in western Baf-490 fin Bay. Elementa: Science of the Anthropocene 7, 34 (2019). **URL** 491 https://online.ucpress.edu/elementa/article/doi/10.1525/elementa.372/11 492
- [14] Porter, D. F. et al. Evolution of the Seasonal Surface Mixed Layer of the Ross Sea, Antarctica, Observed With Autonomous Profiling Floats. Journal of Geophysical Research:
   Oceans 124, 4934–4953 (2019).

- 496 [15] Arteaga, L. A., Boss, E., Behrenfeld, M. J., Westberry, T. K. & Sarmiento, J. L. Seasonal 497 modulation of phytoplankton biomass in the Southern Ocean. *Nature Communications* **11** 498 (2020). URL http://dx.doi.org/10.1038/s41467-020-19157-2.
- toplankton growth before sea ice retreat. *Biogeosciences* **18**, 25–38 (2021). URL https://bg.copernicus.org/articles/18/25/2021/.
- 502 [17] Apollonio, S. Hydrobiological measurements on IGY drifting station Bravo. *Trans. Am.*503 *Geophys. Union* **40**, 316–3 (1959).
- [18] Laney, S. R. *et al.* Assessing algal biomass and bio-optical distributions in perennially ice-covered polar ocean ecosystems. *Polar Science* **8**, 73–85 (2014). URL

  https://linkinghub.elsevier.com/retrieve/pii/S1873965213000510.
- 507 [19] Boles, E. et al. Under-Ice Phytoplankton Blooms in the Central Arctic

  508 Ocean: Insights From the First Biogeochemical IAOOS Platform Drift in 2017.

  509 Journal of Geophysical Research: Oceans 125, 6069–6079 (2020). URL

  510 https://onlinelibrary.wiley.com/doi/10.1029/2019JC015608.
- [20] Haëntjens, N., Boss, E. & Talley, L. D. Revisiting Ocean Color algorithms for chlorophyll-a and particulate organic carbon in the Southern Ocean using biogeochemical floats. *Journal of Geophysical Research: Oceans* 122, 6583–6593 (2017). URL https://onlinelibrary.wiley.com/doi/10.1002/2017JC012844.
- 515 [21] Johnson, K. S. *et al.* Biogeochemical sensor performance in the SOCCOM profiling
  516 float array. *Journal of Geophysical Research: Oceans* **122**, 6416–6436 (2017). URL
  517 https://onlinelibrary.wiley.com/doi/10.1002/2017JC012838.

- 518 [22] Mayot, N. et al. Assessing Phytoplankton Activities in the Seasonal Ice Zone of the

  Greenland Sea Over an Annual Cycle. Journal of Geophysical Research: Oceans 123,

  8004–8025 (2018). URL http://doi.wiley.com/10.1029/2018JC014271.
- [23] Moore, J. K. & Abbott, M. R. Phytoplankton chlorophyll distributions and primary production in the Southern Ocean. *Journal of Geophysical Research: Oceans* **105**, 28709–28722 (2000). URL http://doi.wiley.com/10.1029/1999JC000043.
- [24] Meier, W. N., Fetterer, F., Windnagel., A. K. & Stewart, J. S. NOAA/NSIDC Climate Data
   Record of Passive Microwave Sea Ice Concentration, Version 4. (2021).
- 526 [25] Lizotte, M. P. The Contributions of Sea Ice Algae to Antarctic Ma-527 rine Primary Production. American Zoologist 41, 57–73 (2001). URL 528 https://academic.oup.com/icb/article-lookup/doi/10.1093/icb/41.1.57.
- [26] Arrigo, K. R. Physical control of chlorophyll a, POC, and TPN distributions in the pack ice of the Ross Sea, Antarctica. *Journal of Geophysical Research* **108**, 3316 (2003). URL https://onlinelibrary.wiley.com/doi/10.1029/2001JC001138.
- <sup>532</sup> [27] Arrigo, K. R. *et al.* Photophysiology in two major southern ocean phytoplankton taxa:

  Photosynthesis and growth of phaeocystis antarctica and fragilariopsis cylindrus under

  different irradiance levels. *Integrative and Comparative Biology* **50**, 950–966 (2010).
- The Effect of Melt Pond Geometry on the Distribution of Solar Energy Under First-Year Sea Ice. *Geophysical Research Letters* **47** (2020). URL https://onlinelibrary.wiley.com/doi/abs/10.1029/2019GL085956.
- 539 [29] Katlein, C., Valcic, L., Lambert-Girard, S. & Hoppmann, M. New in-540 sights into radiative transfer within sea ice derived from autonomous opti-

- cal propagation measurements. *The Cryosphere* **15**, 183–198 (2021). URL https://tc.copernicus.org/articles/15/183/2021/.
- [30] Briggs, E. M., Martz, T. R., Talley, L. D., Mazloff, M. R. & Johnson,

  K. S. Physical and Biological Drivers of Biogeochemical Tracers Within

  the Seasonal Sea Ice Zone of the Southern Ocean From Profiling Floats.

  Journal of Geophysical Research: Oceans 123, 746–758 (2018). URL

  https://onlinelibrary.wiley.com/doi/10.1002/2017JC012846.
- <sup>548</sup> [31] Chase, A. P. *et al.* Evaluation of diagnostic pigments to estimate phytoplankton size classes. *Limnology and Oceanography: Methods* (2020).
- ICESat-2. Remote Sensing of Environment 245, 111827 (2020). URL https://doi.org/10.1016/j.rse.2020.111827.
- poteau, A., Boss, E. & Claustre, H. Particulate concentration and seasonal dynamics in the mesopelagic ocean based on the backscattering coefficient measured with Biogeochemical-Argo floats. *Geophysical Research Letters* **44**, 6933–6939 (2017). URL http://doi.wiley.com/10.1002/2017GL073949.
- 557 [34] Klatt, O., Boebel, O. & Fahrbach, E. A profiling float's sense of ice. *Jour-*558 nal of Atmospheric and Oceanic Technology 24, 1301–1308 (2007). URL
  559 http://journals.ametsoc.org/doi/abs/10.1175/JTECH2026.1.
- 560 [35] Claustre, H. et al. Bio-Optical Profiling Floats as New Observational Tools for

  Biogeochemical and Ecosystem Studies: Potential Synergies with Ocean Color Re
  mote Sensing. In Proceedings of OceanObs'09: Sustained Ocean Observations

- and Information for Society, 1, 177–183 (European Space Agency, 2010). URL http://www.oceanobs09.net/proceedings/cwp/cwp17.
- Recommendations for obtaining unbiased chlorophyll estimates from in situ chlorophyll fluorometers: A global analysis of WET Labs ECO sensors. *Limnology and Oceanography: Methods* **15**, 572–585 (2017). URL https://onlinelibrary.wiley.com/doi/10.1002/lom3.10185.
- [37] Hergert, W. & Wriedt, T. The Mie theory: basics and applications (Springer, 2012).
- al.[38] Graff, J. R. etAnalytical phytoplankton carbon measure-570 Sea Research ments spanning diverse ecosystems. Deep Part 571 102, 16 - 25**URL** I: Oceanographic Research **Papers** (2015).572 https://linkinghub.elsevier.com/retrieve/pii/S0967063715000801. 573
- ing satellite estimates of particulate backscatter in the global open ocean using autonomous profiling floats. *Optics Express* **27**, 30191 (2019). URL
- Fig. [40] Bisson, K. M., Boss, E., Werdell, P. J., Ibrahim, A. & Behrenfeld, M. J.

  Particulate Backscattering in the Global Ocean: A Comparison of Independent Assessments. *Geophysical Research Letters* **48** (2021). URL

  https://onlinelibrary.wiley.com/doi/10.1029/2020GL090909.
- ice in the Southern Ocean? Geophysical Research Letters 1–14 (2021). URL https://onlinelibrary.wiley.com/doi/10.1029/2021GL095051.

- [42] Haëntjens, N. et al. Detecting Mesopelagic Organisms Using Biogeochemical-Argo
   Floats. Geophysical Research Letters 47 (2020).
- [43] SMITH, W. O. & NELSON, D. M. Phytoplankton Bloom Pro-587 duced by a Receding Ice Edge in the Ross Sea: **Spatial** Coher-588 ence with the Density Field. *Science* **227**, 163–166 (1985).**URL** 589 http://www.sciencemag.org/content/227/4683/163.shorthttp://www.sciencem 590
- [44] Smetacek, V. et al. Early spring phytoplankton blooms in ice platelet 591 Weddell Sea, the southern Antarctica. Deep Sea Research layers of 592 Part A. Oceanographic Research Papers 39, 153–168 (1992). URL 593 https://linkinghub.elsevier.com/retrieve/pii/019801499290102Y. 594
- [45] Arrigo, K. R. & Thomas, D. N. Large scale importance of sea ice biology in the Southern
   Ocean. *Antarctic Science* 16, 471–486 (2004).
- [46] McMinn, A., Martin, A. & Ryan, K. Phytoplankton and sea ice al-597 transition between gal biomass and physiology during the winter and 598 Polar Biology spring (McMurdo Sound, Antarctica). 33, 1547–1556 599 (2010).URL http://www.nature.com/articles/175238c0 http://doi.wiley.com/10.1002/2016JC012325 601 http://link.springer.com/10.1007/s00300-010-0844-6. 602
- [47] Cummings, V. J. *et al.* In situ response of Antarctic under-ice primary producers to experimentally altered pH. *Scientific Reports* **9**, 6069 (2019). URL http://dx.doi.org/10.1038/s41598-019-42329-0 http://www.nature.com/articles/s41598-019-42329-0.

- [48] Roach, L. A. *et al.* Antarctic Sea Ice Area in CMIP6. *Geophysical Research Letters* 1–24 (2020).
- [49] Briegleb, B. P. & Light, B. A Delta-Eddington multiple scattering parameterization for
   solar radiation in the sea ice component of the Community Climate System Model. Tech.
   Rep. February, National Center for Atmospheric Research (2007).
- [50] Danabasoglu, G. et al. The Community Earth System Model Version 2 (CESM2). Journal
   of Advances in Modeling Earth Systems 12, 1–35 (2020).
- [51] Singh, H. K. A., Landrum, L., Holland, M. M., Bailey, D. A. & DuVivier, A. K. An
   Overview of Antarctic Sea Ice in the Community Earth System Model version 2, Part
   I: Analysis of the Seasonal Cycle in the Context of Sea Ice Thermodynamics and Coupled Atmosphere-Ocean-Ice Processes. *Journal of Advances in Modeling Earth Systems* (2020).
- [52] Kwok, R. *et al.* ATLAS/ICESat-2 L3A Sea Ice Height, Version 1. Boulder, Colorado
   USA. Tech. Rep. May, NSIDC, Boulder, Colorado USA (2019).
- [53] Neumann, T. A. *et al.* The Ice, Cloud, and Land Elevation Satellite 2 mission: A global geolocated photon product derived from the Advanced Topographic

  Laser Altimeter System. *Remote Sensing of Environment* 233, 111325 (2019). URL

  https://linkinghub.elsevier.com/retrieve/pii/S003442571930344X.
- 625 [54] Horvat, C., Blanchard-Wrigglesworth, E. & Petty, A. Observing waves
  626 in sea ice with ICESat-2. *Geophysical Research Letters* (2020). URL
  627 https://onlinelibrary.wiley.com/doi/abs/10.1029/2020GL087629.

- [55] Kwok, R. Sea ice concentration estimates from satellite passive microwave radiometry and openings from SAR ice motion. *Geophysical Research Letters* **29**, 25–1–25–4 (2002).

  URL http://doi.wiley.com/10.1029/2002GL014787.
- [56] Notz, D., Haumann, F. A., Haak, H., Jungclaus, J. H. & Marotzke, J. Arctic sea-ice evolution as modeled by Max Planck Institute for Meteorology's Earth system model. *Journal of Advances in Modeling Earth Systems* 5, 173–194 (2013). URL http://dx.doi.org/10.1002/jame.20016 http://doi.wiley.com/10.1002/jame.20016.
- lite passive microwave sea-ice concentration data set inter-comparison for Arctic summer conditions. *The Cryosphere* **14**, 2469–2493 (2020). URL https://tc.copernicus.org/articles/14/2469/2020/.
- [58] Tsujino, H. et al. JRA-55 based surface dataset for driving ocean—sea ice models (JRA55-do). Ocean Modelling 130, 79–139 (2018). URL
   https://linkinghub.elsevier.com/retrieve/pii/S146350031830235X.
- [59] Smith, W. O. & Nelson, D. M. Importance of Ice Edge Phytoplankton Production in the Southern Ocean. *BioScience* 36, 251–257 (1986). URL

  https://academic.oup.com/bioscience/article-lookup/doi/10.2307/1310215.
- <sup>646</sup> [60] Perrette, M., Yool, A., Quartly, G. D. & Popova, E. E. Near-ubiquity of ice-edge blooms in the Arctic. *Biogeosciences* **8**, 515–524 (2011).
- 648 [61] Strong, C. & Rigor, I. G. Arctic marginal ice zone trending wider in summer and
  649 narrower in winter. *Geophysical Research Letters* **40**, 4864–4868 (2013). URL
  650 http://doi.wiley.com/10.1002/grl.50928.

- [62] Letelier, R. M., Karl, D. M., Abbott, M. R. & Bidigare, R. R. Light driven seasonal patterns of chlorophyll and nitrate in the lower euphotic zone of the North Pacific Subtropical
   Gyre. Limnology and Oceanography 49, 508–519 (2004).
- [63] Boss, E. & Behrenfeld, M. In situ evaluation of the initiation of the North Atlantic phyto plankton bloom. *Geophysical Research Letters* 37, 1–5 (2010).
- 656 [64] Matthes, L. C. *et al.* Average cosine coefficient and spectral distribution of the light field 657 under sea ice: Implications for primary production. *Elementa* **7** (2019).
- 658 [65] Yu, X., Wu, Z., Jiang, W. & Guo, X. Predicting daily photosynthetically active radiation from global solar radiation in the Contiguous United

  660 States. Energy Conversion and Management 89, 71–82 (2015). URL

  661 https://linkinghub.elsevier.com/retrieve/pii/S0196890414008395.
- 662 [66] Taylor, J. R. & Ferrari, R. Shutdown of turbulent convection as a new criterion for the 663 onset of spring phytoplankton blooms. *Limnology and Oceanography* **56**, 2293–2307 664 (2011).
- [67] Holland, single-column model representation of Μ. M. improved 665 associated with summertime leads: ocean mixing Results from a **SHEBA** 666 Journal of Geophysical Research 108, case study. 3107 (2003).URL 667 http://doi.wiley.com/10.1029/2002JC001557. 668
- 669 [68] Horvat, C., Tziperman, E. & Campin, J.-M. Interaction of sea ice floe size, ocean ed-670 dies, and sea ice melting. *Geophysical Research Letters* 43, 8083–8090 (2016). URL 671 http://doi.wiley.com/10.1002/2016GL069742.
- 672 [69] Pellichero, V., Sallée, J.-B., Schmidtko, S., Roquet, F. & Charrassin, J.-B.
  673 The ocean mixed layer under Southern Ocean sea-ice: Seasonal cycle and forc-

- ing. Journal of Geophysical Research: Oceans **122**, 1608–1633 (2017). URL http://doi.wiley.com/10.1002/2016JC011970.
- [70] Pinkerton, M. H., Bradford-Grieve, J. M. & Hanchet, S. M. A balanced model of the food
   web of the Ross Sea, Antarctica. *CCAMLR Science* 17, 1–31 (2010).
- 678 [71] Horvat, C., Seabrook, S., Cristi, A., Matthes, L. & Bisson, K. Code for: The Case for

  Phytoplankton Blooms Under Antarctic Sea Ice (2021).

## **7** Corresponding Author

All correspondence and requests for material should be addressed to Christopher Horvat, christopher\_horvat

#### 8 Acknowledgements

CH was supported by NASA grant 80NSSC20K0959 and in part by Schmidt Futures – a philan-683 thropic initiative that seeks to improve societal outcomes through the development of emerging 684 science and technologies. CH thanks the National Institute of Water and Atmospheric Re-685 search in Wellington, NZ for their hospitality during parts of this work. KB was supported 686 by NASA grant 80NSSC20K0970. We acknowledge the World Climate Research Programme, 687 which, through its Working Group on Coupled Modelling, coordinated and promoted CMIP6. 688 We thank the climate modeling groups for producing and making available their model output, 689 the Earth System Grid Federation (ESGF) for archiving the data and providing access, and the 690 multiple funding agencies who support CMIP6 and ESGF. 691

#### 9 Author Contributions

- 693 CH and SS concieved of the study. CH performed the data analysis and prepared the manuscript.
- KB provided BGC-Argo data processing and analysis. All authors assisted with study design

695 and manuscript writing.

## 696 10 Competing Interests

697 The authors declare no competing interests.

## 698 11 Data Availability

ICESat-2 data are available through the National Snow and Ice Data Center (NSIDC). The 699 sea ice type product is found online at https://nsidc.org/data/ATL07/versions/3. JRA55-do and 700 CMIP6 data used in this study are available at the Earth System Federated Grid at https://esqf-node.lln 701 and https://esgf-node.llnl.gov/projects/cmip6/, respectively. Argo data were 702 collected and made freely available by the Southern Ocean Carbon and Climate Observations 703 and Modeling (SOCCOM) Project funded by the National Science Foundation, Division of Po-704 lar Programs (NSF PLR -1425989 and OPP-1936222), supplemented by NASA, and by the In-705 ternational Argo Program and the NOAA programs that contribute to it. http://www.argo.ucsd.edu, 706 https://soccompu.princeton.edu/www/index.html. The Argo Program is part 707 of the Global Ocean Observing System. Processed Argo data for making figures is included in 708 the code repository [71].

## 12 Code Availability

Code for processing data and producing Antarctic under-ice light fields and UIB-permitting criteria is publicly available on github at https://github.com/chhorvat/Antarctic-Light/, with releases archived in the Zenodo repository [71]. Data required to produce figures in this manuscript is included within this repository.

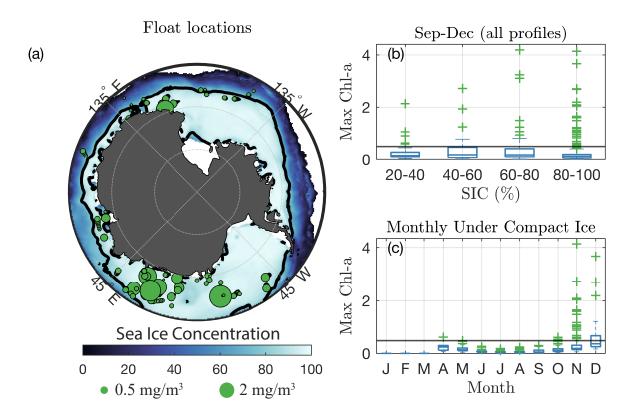


Figure 1: Chlorophyll-a recorded under compact sea ice by BGC-Argo floats. (a) Climatological sea ice coverage in September-November, 2014-2020. Black line shows 80% concentration contour. Green circles are locations of under-ice Argo float profiles under compact sea ice from September-December, with sizes scaled with value of  $Chl_{max}$ . Green dots outside of map shows sizes corresponding to 0.5 and 2.0 mg/m³. (b) Box plot of  $Chl_{max}$  for all BGC-Argo measurements under sea ice, indexed by sea ice concentration. Whiskers extend boxes  $\pm 3$  standard deviations from the mean in each month and vertical blue line is ensemble median. Crosses are outlier values ( $\sigma > 3$ ). Black line is 0.5 mg Chl-a/m³. (c) Same as (b), but for  $Chl_{max}$  recorded under compact sea ice (concentration > 80%) only, indexed by month.

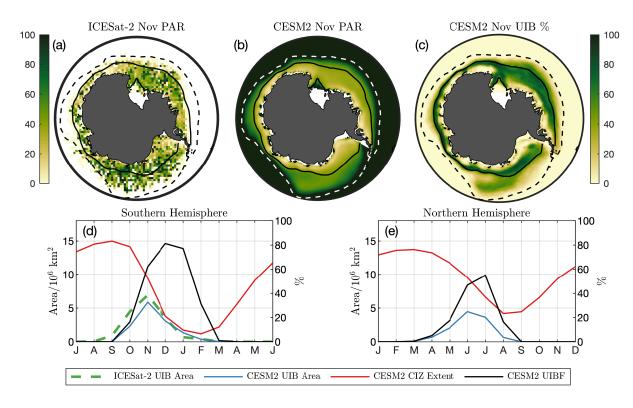


Figure 2: **Light field and UIB potential under Southern Ocean sea ice** (a) 2018-2020 November surface PAR ( $\mu$ mol photons/m²/s) estimate from ICESat-2. (b) CESM2 climatological PAR from pre-industrial simulation. Solid lines in (a-b) are CESM2 climatological CIZ (concentration above 80%). Dashed lines are climatological SIE (concentration above 15%). (c) CESM2 November UIB%. (d,left axis) Seasonal cycle of CESM2 (red) CIZ extent and (blue) UIB extent. Dashed green line is UIB area from ICESat-2. (d,right axis) CESM2 UIBF. (e) As in (d), but for the Northern Hemisphere. Axes in (d) and (e) are offset by 6 months.

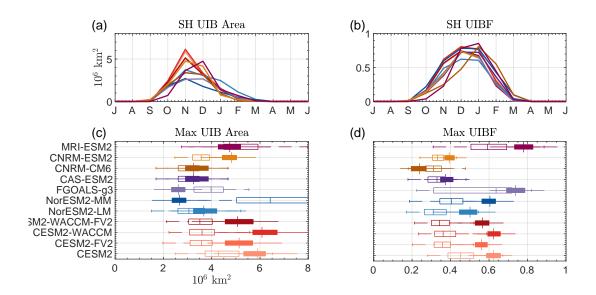


Figure 3: **Statistics of bloom-permitting area for CMIP6 models**. (a) Seasonal cycle of UIB-permitting area in the Southern Hemisphere. (b) Seasonal cycle of UIBF. (c) Box plots of maximum annual UIB area in (filled) the Southern Hemisphere or (unfilled) the Northern Hemisphere. (d) Box plots of UIBF during month of maximum UIB area. Colors of lines in (a,b) correspond to boxes in (c,d)

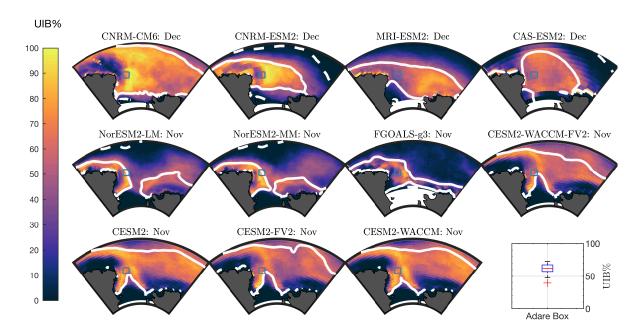


Figure 4: **UIB likelihoods in the Ross Sea.** Ross Sea UIB% for each model in the month of maximum UIB area. Solid lines are climatological CIZ. Dashed lines are climatological SIE. Blue square highlights location of interest at 72°S,178°E. Box plot (bottom right) is of UIB% at square location.