A tale of three islands: downstream natural iron fertilization in the Southern Ocean

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Key points:

• Iron fertilization of blooms downstream of Southern Ocean islands studied with Lagrangian

6 modelling

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• Ocean areas fertilized by simulated iron transport overlap with observed spatial extent of

9 blooms

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• Inter-annual variability of iron input explains blooms at Crozet but not Kerguelen or South

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Abstract. Iron limitation of primary productivity prevails across much 14 of the Southern Ocean but there are exceptions; in particular, the phy-15 toplankton blooms associated with the Kerguelen Plateau, Crozet Islands 16 and South Georgia. These blooms occur annually, fertilized by iron and 17 nutrient-rich shelf waters that are transported downstream from the is-18 lands. Here we use a high-resolution $(1/12^{\circ})$ ocean general circulation 19 model and Lagrangian particle tracking to investigate whether inter-annual 20 variability in the potential lateral advection of iron, could explain the inter-21 annual variability in the spatial extent of the blooms. Comparison with 22 ocean color data, 1998 to 2007, suggests that iron fertilization via advec-23 tion can explain the extent of each island's annual bloom, but only the 24 inter-annual variability of the Crozet bloom. The area that could potentially be fertilized by iron from Kerguelen was much larger than the bloom, suggesting that there is another primary limiting factor, potentially sili-27 cate, that controls the inter-annual variability of bloom spatial extent. For 28 South Georgia, there are differences in the year-to-year timing of advection 29 and consequently fertilization, but no clear explanation of the inter-annual 30 variability observed in the bloom's spatial extent has been identified. The 31 model results suggest that the Kerguelen and Crozet blooms are terminated 32 by nutrient exhaustion, probably iron and or silicate, whereas the deepening 33 of the mixed layer in winter terminates the South Georgia bloom. There-34 fore, iron fertilization via lateral advection alone can explain the annual 35 variability of the Crozet bloom, but not fully that of the Kerguelen and 36

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- ³⁷ South Georgia blooms.
- 38

1. Introduction

It is now generally accepted that iron, in conjunction with light, is a major limiting 30 factor of primary production in the Southern Ocean, indirectly controlling the biologi-40 cal pump and drawdown of carbon dioxide from the atmosphere [Takahashi et al., 2009; 41 Blain et al., 2007; Boyd et al., 2007; de Baar et al., 1995; Martin, 1990; Martin et al., 42 1990]. However, there are exceptions to the high nutrient, low chlorophyll conditions that 43 prevail across most of the Southern Ocean. Large phytoplankton blooms are observed downstream of continental shelf and land mass [Blain et al., 2007; Pollard et al., 2007; 45 Korb et al., 2008, where iron is suggested to be supplied to surface waters predominately from ocean sediments [Bakker et al., 2007; Tyrrell et al., 2005; Thomalla et al., 2011]. In 47 order to understand these important high productivity regions, we need to characterize 48 the timescales and mechanisms that transport iron to where primary production occurs 49 [Boyd et al., 2012; d'Ovidio et al., 2015; Wadley et al., 2014]. Here we focus on three 50 Southern Ocean islands groups, the Kerguelen Plateau, Crozet Islands and South Georgia 51 and Shag Rocks, outlined by black boxes in Figure 1a, specifically looking at the role of 52 advection in determining the spatial extent of the downstream blooms. 53

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Iron supply in the Southern Ocean comes from a variety of different sources including: aeolian input; brine rejection and drainage from sea ice; sediments; entrainment from the deep ocean via winter mixing, Ekman pumping, and upwelling at ocean fronts; and it is also constantly resupplied via rapid recycling of organic material [*Boyd and Ellwood*, 2010; *Gille et al.*, 2014; *Graham et al.*, 2015; *Korb et al.*, 2008; *Schallenberg et al.*, 2015;

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Tagliabue et al., 2014]. A recent study by Graham et al. [2015] suggests that coastlines, 60 continental and island, are key sources of iron to the Southern Ocean, and also provides a 61 comprehensive description of the behavior of iron in sediment pore waters and the mech-62 anisms behind its flux into over-lying bottom water. Another source, recently found to 63 be relevant to the Kerguelen Plateau, is riverine input associated with snowmelt. This 64 source is important during spring, as there is increased rainfall and runoff, whereas freez-65 ing conditions during the winter inhibit this iron supply [van der Merwe et al., 2015]. In 66 this paper we focus on the potential for iron fertilization from island sources, primarily 67 from sediments and run-off. 68

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As iron is released from island sediments, internal waves and turbulence mix the iron 70 up into surface waters which then fuels phytoplankton production [Bowie et al., 2015; 71 Boyd, 2007; Korb et al., 2008; Park et al., 2008]. Iron that is not immediately utilized by 72 biota or scavenged from the water column can be transported downstream of its source 73 via lateral advection within the local circulation and also by stirring within mesoscale fea-74 tures [Abraham et al., 2000; d'Ovidio et al., 2015]. As it advects, iron can undergo many 75 processes and transformations as part of the complex iron cycle, which can alter both the 76 transport and bioavailability of iron. For instance, iron can be diluted by physical mixing, 77 it can be kept in circulation by iron–binding ligands, or there can be luxury uptake of the 78 iron by biota and hence "internal advection" [Mongin et al., 2008]. In various forms, iron 79 can be lost from the surface by sinking or it can be retained in the surface water and then 80 remineralized downstream of the original source and supply a new area with iron [Boyd81

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⁸² et al., 2000; Boyd, 2007].

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In order to test the hypothesis that inter-annual variability observed in the spatial ex-84 tent of downstream island blooms could be explained by horizontal advection, the details 85 of the iron cycle are not considered here. In this paper the term "iron advection" refers 86 to any iron from island sources in a form that can be laterally transported, via either ad-87 vection or stirring, and is also bioavailable at the bloom site, hence what is demonstrated 88 in this paper is the potential for iron fertilization. To diagnose the advection around 89 each island, Lagrangian particles were released within velocity fields from the NEMO 90 (Nucleus for European Modelling of the Ocean) $1/12^{\circ}$ ocean general circulation model, a 91 resolution high enough to resolve eddies and small scale circulation patterns around the 92 islands. In the analysis the Lagrangian trajectories, representing water mass potentially 93 fertilized with iron, are compared against the observed bloom areas in the satellite data. Additionally, the possible causes for bloom termination will be considered for each island, 95 utilising the model diagnostics and also World Ocean Atlas nutrient data.

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2. Methodology

In order to assess the impact of iron that could potentially be advected downstream of Southern Ocean islands, satellite derived data (chlorophyll–a concentrations and sea surface currents) were compared with Lagrangian particle trajectories within velocity output from the NEMO $1/12^{\circ}$ model. Here we give a brief description of each of the three study sites, the tools used, and explain the experimental design.

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2.1. Study sites

The Kerguelen Plateau and Heard Island (southeast of the Kerguelen Island), depicted 104 by the box on the right of Figure 1d (bathymetry plot), is a major bathymetry feature 105 within the Indian Ocean sector of the Southern Ocean, extending from $46^{\circ}S$ to $64^{\circ}S$ at the 106 3000 m isobath. It forms a major barrier to the eastward flowing Antarctic Circumpolar 107 Current (ACC), with most of the flow being deflected to the north of the plateau (\sim 108 100 Sv), and the substantial remainder to the south (30-40 Sv), steered primarily by the 109 topography. The circulation over the plateau between the two islands is rather stagnant, 110 <5 cm $^{-s}$ on average. A major circulation feature within the region is the Polar Front 111 (PF), which cuts between the two islands, flowing close to the southeast Kerguelen Island 112 [Park et al., 2008, 2014]. The Kerguelen bloom occurs on decadal average during Novem-113 ber to January, as demonstrated in Figure 3a., and is predominately made up of diatom 114 species above the plateau [Blain et al., 2001]. However, note that the Kerguelen bloom 115 can persist for much longer periods, due to a concurrent resupply of essential nutrients via 116 remineralisation and entrainment from the deep ocean during vertical mixing [Boyd, 2007]. 117

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¹¹⁹ Crozet Islands (46°S, 52°E), depicted by the central black box in Figure 1d, is separated ¹²⁰ from the Del Cano Rise plateau to the west by the Subantarctic Front (SAF), which is the ¹²¹ dominant circulation feature in the area. The SAF predominately lies west to east within ¹²² the ACC, but turns sharply north between the two plateaus (Crozet and Del Cano Rise), ¹²³ before turning eastward to the north of Crozet as it comes into contact with the Agulhas ¹²⁴ Return Current [*Bakker et al.*, 2007; *Pollard et al.*, 2007]. Over the plateau and to the ¹²⁵ north of the island (bounded by the SAF) is an area of Polar Frontal Zone characterised

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¹²⁶ by weak circulation (15 – 20 Sv), within which iron can accumulate during the winter ¹²⁷ months that can subsequently fuel a bloom [*Planquette et al.*, 2007]. The phytoplankton ¹²⁸ community structure of the Crozet bloom, described by *Poulton et al.* [2007], is made ¹²⁹ up of varying sizes of diatoms, and very small prymnesiophyte *Phaeocystis antarctica*. ¹³⁰ Biomass varies considerably near to the plateau between species, but further away from ¹³¹ the plateau, to the northwest and east, prymnesiophyte *P.antarctica* can dominate.

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South Georgia, and Shag Rocks (northwest of South Georgia), hereafter referred to 133 collectively as South Georgia, are located to the east of Drake Passage, highlighted by the 134 left black box in Figure 1d. The islands form part of the North Scotia Ridge at roughly 135 54°S, 37°W, directly in the path of the ACC. The PF lies north of the islands, and the 136 Southern ACC Front flows to the south, looping anti-cyclonically around South Georgia 137 before flowing east again [Orsi et al., 1995; Meredith et al., 2003]. North of the island, 138 enclosed by the PF and Southern ACC Front, is the South Georgia Basin, within which 139 prolonged blooms exist throughout the growing season [Borrione and Schlitzer, 2013]. 140 This paper will focus on the South Georgia Basin bloom, but there are blooms occurring 141 to the south and west of the islands [Ward et al., 2007], although these blooms are partly 142 subsurface and may not be represented by satellite observations. Furthermore the region is 143 one of the most productive regions across the entire Southern Ocean, with various sources 144 of iron and phytoplankton [Ardelan et al., 2008; Murphy et al., 2013; Thomalla et al., 145 2011]. Consequently, from satellite ocean color data alone, it is not possible to delineate 146 blooms fertilised by iron from South Georgia sediments or from elsewhere within the basin 147 (Antarctic peninsula or ice melt). The South Georgia Basin bloom (hereafter referred to 148

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as the South Georgia bloom) is dominated by large diatom species, but is described as "patchy" over scales of 10 - 20 km, with fragmented diatom colonies occurring alongside a more invariant community of small autotrophs and heterotrophs [*Atkinson et al.*, 2008; *Korb et al.*, 2008].

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Each of the three islands have different characteristics which determine the ecosystem that they support and its functioning, but for a generalized overview of the Southern Ocean ecosystem see *Boyd* [2002]. These islands have been selected for this study as their blooms have been extensively explored in the field [*Blain et al.*, 2008; *Pollard et al.*, 2007; *Korb et al.*, 2008; *Murphy et al.*, 2013], the results from which can be used to support our own analysis.

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2.2. Satellite data

¹⁶¹ 2.2.1. Chlorophyll observations

The ocean color data used in this study comes from the ESA Ocean Colour Climate 162 Change Initiative. Here we use a (level 3 geographically mapped) merged and bias cor-163 rected product from the MERIS, MODIS and SeaWiFS datasets, with a horizontal reso-164 lution of up to 4 km [Storm et al., 2013]. Because of the low solar elevation and sea-ice 165 coverage in winter, data is unavailable in some areas, most visibly the Weddell Sea in 166 Figure 1a, but by averaging over a month, year and decade, we can fill in many of the 167 gaps. This study has utilized monthly chlorophyll–a (chl–a) concentrations over the pe-168 riod 1998 - 2007, the first decade in which we have good satellite coverage across the 169 world. In this study, the chlorophyll data is used to represent phytoplankton biomass, 170

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¹⁷¹ defining the island blooms.

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¹⁷³ 2.2.2. Altimetric sea surface currents

The satellite altimeter data is produced by Ssalto/Duacs and distributed by the Archiving Validation and Interpretation of Satellite Data in Oceanography (Aviso) group, with support from CNES (http://www.aviso.altimetry.fr/duacs/). Here we utilize a merged dataset, from only two satellites at any one time, each having the same ground track and stable sampling which provides a homogenous time series. This along-track, delayed time data product has great stability and therefore is the ideal product for use in inter-annual comparison studies [*Le Bars et al.*, 2014].

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The along-track Absolute Dynamic Topography (ADT) is obtained by adding the Sea Level Anomaly to the Mean Dynamic Topography (Mean Sea Surface Height minus Geoid). A mapping procedure using optimal interpolation with realistic correlation functions is applied to produce ADT maps (MADT or L4 products) onto a Cartesian $1/4^{\circ}$ x $1/4^{\circ}$ grid [*Aviso*, 2014]. Here we use the sea surface geostrophic velocities computed from the ADT over the period of 1998 – 2007.

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2.3. NEMO model and Ariane Lagrangian particle tracking

The NEMO $1/12^{\circ}$ resolution ocean general circulation model has been developed with particular emphasis on realistic representation of fine-scale circulation patterns [*Madec*, 2008], which provides an ideal platform to conduct Lagrangian particle-tracking experiments around the small islands of the Southern Ocean. Full details of the model run,

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¹⁹³ including model setup and configuration, can be found in *Marzocchi et al.* [2015] as only ¹⁹⁴ a brief description will be given here. The model is initialized with World Ocean Atlas ¹⁹⁵ (WOA) 2005 climatological fields and forced with 6–hourly winds, daily heat fluxes, and ¹⁹⁶ monthly precipitation fields [*Brodeau et al.*, 2012]. The run begins in 1978, with output ¹⁹⁷ through to 2010, of which we are interested in 1998 – 2007. Model output is stored offline ¹⁹⁸ as successive 5 day means throughout the model run, of which the velocity fields are used ¹⁹⁹ for the particle tracking in this paper.

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The Ariane package [Blanke & Raynaud, 1997] (available online at: http://stockage. univ-brest.fr/~grima/Ariane) is applied to the NEMO velocity field to track water parcels using point particles that are released into the modeled ocean circulation (cf. *Popova et al.* [2013] and *Robinson et al.* [2014], who used output from the NEMO 1/4° model). These particles are intended here to represent water masses fertilized by iron scoured from the island sediments. Further details about the Ariane package can be found in *Blanke & Raynaud* [1997] and *Blanke et al.* [1999].

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An important caveat to the results is that we do not expect the NEMO 1/12° model to reproduce the detailed mesoscale flows year–by–year due to chaotic dynamics, as the mesoscale eddy field is not initialised to match that of the real world (only possible using data assimilation). Nevertheless, the model does reproduce the larger scale flow field in the vicinity of the islands, which is important for downstream advection (see Figure 1b and c).

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2.4. Experiment design

In order to study the advection of iron from island sources and make a qualitative 215 comparison with the ocean color 1998 – 2007 observations, Lagrangian particles were re-216 leased monthly into the modeled circulation from around the shelf regions of each island, 217 from January 1998 to December 2007. Particles are deployed in every other grid cell of 218 the $1/12^{\circ}$ model grid along the horizontal (latitudinally and longitudinally), and at each 219 level of the NEMO grid depth domain down to a maximum depth of 180 m (30 depth 220 levels, not equally spaced see *Madec* [2008]), around each of the three islands (cf. Srokosz 221 et al. [2015], who used a similar analysis for the Madagascar bloom). Figure 2 shows the 222 starting positions of the particles around each of the islands. The particles had to be 223 spaced at a high enough resolution to resolve the fine scale circulation patterns around 224 each island, but the experiments were limited computationally, as the islands are not of 225 a comparable area, so there could not be a particle within every model grid cell. The 226 particles are released in both the horizontal and vertical extent, to represent iron that is 227 scoured from the shelf sediment (down to 180 m in this experiment) and mixed upwards 228 [Ardelan et al., 2008; Blain et al., 2001; Hewes et al., 2008; Planquette et al., 2007] as well 229 as other island sources, such as river run-off [van der Merwe et al., 2015]. Particles that 230 are subducted deeper than 200 m, i.e. out of the euphotic zone, along their trajectory are 231 removed from the analysis. At the horizontal and vertical grid spacing described, that 232 results in 8240 Lagrangian particles being released each month from the Kerguelen and 233 Heard Island, 465 particles from Crozet, and 2820 particles from South Georgia and Shag 234 Rocks. 235

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237 2.4.1. Assumptions and limitations of method

The main assumption in this study is that surface waters in the Southern Ocean are iron limited, and that the addition of iron to an area, via horizontal advection, would initiate a bloom. However in reality, productivity can be co-limited in the Southern Ocean, with light or silicate for example, and there are also seasonal factors which control phytoplankton growth, which can vary in both time and space [*Boyd*, 2002].

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In the analysis to follow, the advection time over a period of twelve months is discussed. 244 Note that the residence time of bioavailable iron in surface waters is not yet fully under-245 stood, but thought to be relatively short, on the order of only weeks to months [Boyd and 246 Ellwood, 2010; Shaked and Lis, 2012; Schallenberg et al., 2015]. However, studies have 247 also shown that iron can be transported during winter months and remain in the upper 248 ocean to be available to stimulate blooms in the summer months [Mongin et al., 2009; 249 d'Ovidio et al., 2015]. Graham et al. [2015] postulates that this might be possible due to 250 intense biological recycling of iron, or the long-range transport of particulate iron, or even 251 by currently unknown processes. For the time being these questions remain unanswered, 252 and so for the purpose of this study all of the iron from the islands is assumed to remain 253 available throughout the year. A further assumption is that all advective pathways have 254 the potential to be fertilized with iron. 255

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A caveat to this analysis is that, in using satellite ocean color data, it is not possible to detect subsurface chlorophyll maxima, which are known to exist in certain regions of the Southern Ocean [*Holm-Hansen et al.*, 2005; *Tripathy et al.*, 2015]. Therefore we cannot

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use our analysis to draw any conclusions on the location or variability of known subsurface
chlorophyll maxima [*Ward et al.*, 2007], and make the distinction now that only surface
blooms are considered, hereafter just referred to as blooms.

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As touched upon in the introduction, the representation of iron and its transport in 264 this method is a simplification. Ideally this study would be performed using tracers in 265 a high-resolution, fully coupled biogeochemical model, but the computational resources 266 required for this would be extreme. Such a study would need a coupled model at a res-267 olution high enough to formally resolve the small-scale circulation features that occur 268 around the islands at the center of this study. As such, the analysis presented in our re-269 sults and discussion is restricted to consider only potential iron advection and consequent 270 fertilization. 271

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3. Results

3.1. Ocean color

Figure 1a is a 10 year average of satellite derived sea surface chl-a concentrations in 273 November, over 1998 to 2007. The islands of interest are highlighted by black boxes, 274 from which it is clear that these island blooms can be more than double the magnitude 275 of productivity anywhere else in the Southern Ocean. Figure 3 is the decadal monthly 276 averages, of surface chl-a concentration, for a single location inside, and a single loca-277 tion outside of the bloom sites for each island. Each location was selected arbitrarily 278 based on persistence either inside or outside (upstream of the ACC) of the annual bloom. 279 The latitude and longitude coordinates of each location inside the bloom are $72^{\circ}E$ and 280

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49°S, 52.5°E and 45.5°S, 38.5°W and 52.5°S; for Kerguelen, Crozet and South Georgia 281 respectively. The coordinates of each location upstream of the bloom are $66^{\circ}E$ and $48^{\circ}S$. 282 45.5°E and 46.5°S, 49.5°W and 52.5°S (cf. Park et al. [2008] their figure 11, Pollard et al. 283 [2009] their figure 1, and Korb et al. [2004] their figure 1, for schematic positioning of the 284 ACC around Kerguelen, Crozet and South Georgia respectively). In this paper, a bloom 285 is defined by chl-a concentrations higher than 0.5 mg m^{-3} , as it is consistently higher 286 than chl-a outside of each islands typical bloom regions [Comiso et al., 1993; Moore and 287 Abbott, 2000]. Also, when 0.5 mg m⁻³ of chl–a is exceeded in Figure 3, it occurs on a 288 steep gradient from one month to the next, indicating the start of a bloom. Addition-289 ally this concentration is low enough to avoid complications with double peaks in chl-a 290 associated with South Georgia, as can be seen in Figure 3c. South Georgia is a region 291 that frequently has two bloom peaks per year [Borrione and Schlitzer, 2013], however it 292 is outside of the scope of this work to analyze peak bloom events. Therefore, in order 293 to focus on inter-annual rather than inter-seasonal variability, we consider the average 294 chl-a concentration over the bloom period. The error bars in Figure 3 are 1 standard 295 deviation in chl–a for each month, over the ten year period. The size of the error bars is 296 an indication of the seasonality across the regions and annual cycles. South Georgia in 297 particular, has large error bars which is due to the range in magnitude of annual blooms. 298 For instance, the average chl-a for January over 1998 - 2007 is $<1 \text{ mg m}^{-3}$, however in 299 January of 2002, the concentration was as high as 15 mg m^{-3} [Korb and Whitehouse, 2004]. 300

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Figure 3 also includes the averaged (decadal) monthly mixed layer depth (MLD) in the bloom' site for each island, calculated online in the NEMO model. Comparing the bloom

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and MLD curves we see that the bloom is likely triggered by the onset of a shallowing mixed layer [Venables and Moore, 2010]. The MLD, specifically its role in terminating the blooms, is considered in further detail in the discussion.

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Figure 4 shows example years of a small and a large averaged bloom period (hereafter referred to as the bloom) for each island, during 1998–2007. Maximum and minimum blooms for Kerguelen occur in 2003 and 2000, Crozet is 2004 and 2001, and South Georgia is 2002 and 2006. Strikingly, Figure 4 demonstrates the strong inter–annual variability in both bloom magnitude and area, which may be explainable by studying the potential iron advection from the islands.

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3.2. NEMO vs. Aviso surface current speed

The ability of the chosen model to accurately represent the circulation in the study area is critical to the quality of the results. In order to assess the performance of the NEMO 1/12° model we can compare with satellite derived sea surface currents (Aviso). The Aviso data is the geostrophic component of the velocity, whereas the NEMO model is the absolute velocity, but this should not impact a comparison between the two as they are near equal at the surface. By comparing the decadal averages of NEMO and Aviso, side by side (Fig. 1), we can assess the models performance.

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Figure 1[b] and 1[c] are a comparison of the decadal (1998 – 2007) average ocean surface current speed, from NEMO and Aviso respectively, across the Southern Ocean. Qualitatively, the model correctly captures the major features, and also their magnitude. Fast

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flowing currents are stronger in the model than Aviso, and also boundaries of fast flowing currents within the modeled circulation are more defined than in the observations. This may be due to data smoothing caused by the correlation function applied to the Aviso data or due to the model under representing sub-mesoscale features. Figures S1 - S3 in the supporting information show the decadal, annual and monthly averaged circulation, of both model and satellite derived velocities, for each island for illustrative purposes.

3.3. Advection of iron towards the bloom site

In this paper we hypothesize that the advection of iron downstream of islands allow blooms to occur in the otherwise high nutrient, low chlorophyll regime of the Southern Ocean. Here we investigate the timescales of fertilization, and the degree to which the circulation can impact inter-annual variability, during the period 1998 – 2007.

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The Kerguelen bloom occurs on decadal average during November to January, as demon-338 strated in Figure 3a. For this analysis, we focus on the average surface chl-a concentration 339 over the bloom period (November to January in Kerguelen's case) for each year, referred 340 to as the bloom. Figure 5 shows the patch around Kerguelen that could potentially be 341 fertilized with iron by the local circulation in the NEMO model. The fertilized patch 342 is depicted by colored markers, which represent the location of trajectories in October 343 for each year, with the different colors indicating the month in which the particles were 344 released from the island. Strikingly, the fertilized patch is much larger than the bloom 345 extent, represented by black contours in each annual subplot. The trajectories propagate 346 east from the island between the latitude band of roughly $45^{\circ}S - 54^{\circ}S$, but then spread 347

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³⁴⁸ both northward and southward in extent from roughly 77°E. However, despite the fertil-³⁴⁹ ized patch reaching as far north as 40°S in Figure 5, we can see from the black contours ³⁵⁰ that the bloom area is never north of 45°S in any of the years.

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Having found that the horizontal advection of iron would be sufficient to fertilize the 352 bloom in principle, a further question arises as to whether the bloom is terminated by the 353 exhaustion of iron in the surface water. This question cannot be addressed directly using 354 the NEMO $1/12^{\circ}$ simulation, as it is not a coupled biogeochemistry model. However, if 355 the bloom is terminated by the exhaustion of iron then a question that can be addressed 356 is: can advection resupply iron in the period between the end of one bloom and the start 357 of the next? As addressed by Mongin et al. [2009], and more recently by d'Ovidio et al. 358 [2015]. For Kerguelen, the location of the fertilized patch was very consistent, however 359 there are temporal differences in the timing of advection. Nevertheless, the results show 360 the maximum advection time for the particles to reach the furthest extents of the bloom is 361 on the order of 5-6 months, suggesting that horizontal advection is sufficient to resupply 362 the bloom area with iron, in agreement with Mongin et al. [2009] and d'Ovidio et al. [2015]. 363

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Figure 6, is the same as 5, but focusing on Crozet. The Crozet bloom occurs one month earlier than the Kerguelen bloom, on decadal average during October to December [*Pollard et al.*, 2007], and so the trajectories shown in Figure 6 represent the fertilized patch in September. Figure 6 suggests there is more inter–annual variability in the circulation around Crozet than Kerguelen, both spatially and temporally. In Figure 6, the fertilized patch tends to be north of the island and to the east, made up of particles released in

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June through to August (light green to orange on the color bar). This indicates that the 371 timescale for fertilization, of water mass being within the immediate vicinity of Crozet 372 (where the particles are released) to outside of the bloom area (the black contours), is 373 on the order of 3-4 months, however Figure 6 clearly shows the inter-annual variability 374 in this timescale. There are some years in Figure 6 where we see the fertilized patch 375 extending to the west of the island, most visibly in the years 2000, 2002, 2003, and 2007. 376 The color of the markers seen to the west of the island in some of the years show the 377 particles were released earlier in the year, ranging from January (2000) to April (2002). 378 Focusing on the black contours in Figure 6, representing chl–a concentrations above 0.5 379 mg m⁻³ during the bloom period, there are years in which the bloom is propagated to the 380 west also, most clearly apparent in 2000 and 2007. 381

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The bloom associated with South Georgia occurs on decadal average, during October to 383 April, however South Georgia experiences the highest seasonality of all the three islands 384 in this study. Here we discuss the South Georgia bloom, although the surrounding area is 385 one of the most productive regions within the Southern Ocean [Ardelan et al., 2008; Young 386 et al., 2014, so separating a bloom associated with iron only advected from South Georgia 387 is not non-trivial. In order to address this issue, we have applied a mask to the ocean color 388 data, to remove chl-a that was most likely fertilized from other iron sources in the region, 389 guided by the surface chl-a climatology around South Georgia produced by Borrione and 390 Schlitzer [2013]. Figure 7 is again, the same as Figures 5 and 6, with the colored markers 391 representing the particle locations in September (preceding the start of the bloom). The 392 extent of the fertilized patch around South Georgia changes annually, although to a lesser 393

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degree than around Crozet. What does remain almost annually consistent, is the north 394 and eastward advection of the particles (with the exception of 2006) and an associated 395 bloom occurring within the Georgia Basin, which is just north of the island. In some of 396 the years, most distinctly in 2004 and 2005, there is a well defined boundary edge to the 397 trajectories on the western side of the fertilized patch. This sloping western boundary 398 edge is also apparent in the average bloom area in almost all years (2006 being the most 399 apparent exception). The trajectories and bloom are restricted to the east of this bound-400 ary due to the eastward flowing PF which acts as a physical barrier [Moore et al., 1999; 401 Korb and Whitehouse, 2004]. The colored markers represent the particles locations in the 402 month of September, and therefore particles that are released at the beginning of Septem-403 ber have only had one month to be advected, and consequently are the closest to South 404 Georgia. Focusing just on the recently released particles, from August and September 405 (orange and red), it is apparent that, for the majority of the years, this western boundary 406 of both the fertilized patch and bloom area is an important route for iron to be advected 407 away from South Georgia, flowing towards Shag Rocks and then along the PF. This cir-408 culation feature was also found by Young et al. [2011] in their higher resolution regional 409 model, described as a unidirectional link between the two land masses (see their Figure 7). 410 411

Table 1 provides the size of both the annual blooms and fertilized patches around Kerguelen. As can also be seen in Figure 5, the fertilized patch is much larger than the bloom, and there is more variability in the bloom size than in the fertilized patch. Consequently, the annual percentage of the bloom area that is within the fertilized patch is consistently very high, with an average of 77% (st dev ± 6.5). As the fertilized patch is

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⁴¹⁷ much larger than the bloom area, it would suggest that iron availability is not the only, or ⁴¹⁸ at least most important, factor controlling the Kerguelen bloom extent and inter–annual ⁴¹⁹ variability. The year 2003 had the largest bloom in our study period, in which the bloom ⁴²⁰ did extend out across and to the southern edges of the fertilized patch. It is possible, ⁴²¹ that in 2003 the primary limiting factor to the Kerguelen bloom was alleviated so the ⁴²² bloom could extend further out into the regions of available iron. This hypothesis will be ⁴²³ considered later in the study.

424

Looking at Table 2, the Crozet bloom is a third of the size of the Kerguelen bloom, with an average bloom size of 242,416 km² compared to the Kerguelen average of 782,455 km². Focusing on the percentage of the bloom site overlapped by the trajectories (*Fertilized patch*) for each year, there is a range of 60% - 32% overlap. This is reflected in the percentage of the fertilized patch overlapped by the bloom, ranging from 67% - 34%. Both the bloom area and fertilized patch around Crozet vary annually, and Crozet has the lowest overlap out of the three islands studied.

432

The average size of the South Georgia bloom over 1998 – 2007 was 618,645 km², smaller than the average size of the fertilized patch at 742,038 km². In Table 3, we can see a large range in the bloom area around South Georgia across the years, the maximum being 946,833 km² in 2002 and the minimum being 414,108 km² in 2006 (see Figure 4). There is also a range in the size of the fertilized patch, although not as large as the range in bloom size. Focusing on the amount of overlap between the bloom and trajectories, we see that the annual bloom overlaps are generally larger than the fertilized patch overlaps (2002

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and 2003 being the exceptions). This is due to the fertilized patch being larger than the bloom area for the majority of the years. However, as with the other two islands, there is a range in the annual overlaps, which can be explained by a combination of inter–annual variability in the sizes and locations of the annual blooms, and also, to differing degrees for each island, the inter–annual variability in the size and locations of the fertilized patches (Kerguelen being the most consistent, and Crozet exhibiting the most variation).

446

Figure 8 shows the overlap of the bloom (bloom period average, chl-a concentration 447 greater than 0.5 mg m^{-3}) by the fertilized patch from each individual monthly release of 448 particles. In the Kerguelen plot, we see a maximum range of around 10 - 25% between 449 years, in the overlap between monthly releases of particles and the average bloom. The 450 cause of this range is a combination of inter-annual variability in both the advection and 451 bloom extent. In comparison with Figure 5 and Table 1, it is apparent that the high-452 est degree of variability comes from the bloom, although the inter-seasonal variation in 453 advection timing and consequently fertilization could also impact bloom development. 454 Particles released in October, just prior to the start of the bloom, cover around 10 - 15%455 of the bloom area, with the maximum bloom coverage from releases in April – June for 456 the majority of the years. This gives an advective fertilization timescale of between 5-7457 months for maximum bloom coverage. The circulation on the Kerguelen Plateau itself is 458 known to be sluggish, certain parts even described as stagnant [Park et al., 2014]. This 459 localized slow moving water on the plateau (where particles start) may account for the 460 low bloom overlap percentage by particles released just prior to the start of the bloom 461

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462 (November).

463

In Figure 8, Crozet shows less inter-annual variability than Kerguelen, of less than 10% difference between years. The most apparent difference between Crozet, and the other two islands, is that the advective fertilization timescale is much shorter, with maximum bloom overlap from particles released in June – August, which is 2 – 4 months prior to the start of the bloom (typically October). However, Crozet has the lowest bloom overlap, with a maximum of 25% from an August release in 2000. For the majority of the years, the maximum percent coverage of the bloom is below 20%.

471

The South Georgia plot of Figure 8 shows a degree of consistency in the timing of fertilization, but high variability in the bloom overlap from each monthly release, across the years. The variability in bloom overlap is on the order of 10 - 15%, and the advective fertilization timescale is roughly April – June, 4 - 6 months prior to the typical start of the bloom. The maximum percentage bloom overlap is 40%.

477

4. Discussion

Here we consider other factors that could impact the bloom, light limitation and nutrient control, before addressing our three main research questions: Can advection explain
the extent of the bloom area? Can advection explain the bloom inter-annual variability?
And what factors could cause bloom termination?

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4.1. Light limitation

In addition to iron limitation in the Southern Ocean, light limitation also plays an im-483 portant role in controlling productivity Wadley et al. [2014]. The light levels encountered 484 by phytoplankton cells is partly determined by the mixed layer depth (MLD), as they are 485 vertically mixed between high surface irradiance and low subsurface irradiance (Venables 486 and Moore [2010] – explanations and references therein). To assess the light availability 487 around the islands during the typical bloom periods, Figure 9 shows the decadal average 488 monthly depths of the mixed layer, calculated online in the NEMO model, over 1998 – 489 2007.490

491

The top row of Figure 9, shows the MLD around Kerguelen which remains in a similar 492 spatial pattern during the bloom period, with a distinct divide between the shallower 493 north and deeper south. During the period 1998 – 2007, the Kerguelen bloom is con-494 strained to the south of this divide where the MLD is deepest. The middle row shows 495 the MLD around Crozet which exhibits the typical shallowing north to south of the MLD 496 from winter into summer. In Figure 9 the bottom row is a two month decadal average of 497 the mixed layer for the South Georgia region. Two months have been averaged together 498 in order to capture the entire bloom period within the plot, from which we can see the 499 typical north to south shallowing of the mixed layer from winter into summer. Both the 500 Kerguelen and Crozet Islands blooms have typically terminated when the mixed layer is 501 shallow enough for there still to be light available, which suggests that neither bloom is 502 terminated by light limitation [Venables et al., 2007; Venables and Moore, 2010]. The 503 South Georgia bloom, however, persists for the entire season and typically ends when the 504

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⁵⁰⁵ mixed layer begins to deepen in winter, strongly indicating that the bloom is terminated ⁵⁰⁶ by diminishing light, and not by the exhaustion of iron.

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4.2. Nutrient control

The depth of the mixed layer is also significant for the amount of nutrients being brought to the surface from the deep, such as nitrate and silicate as well as iron. WOA climatologies show the concentration of nitrate to be high across much of the Southern Ocean, south of the Sub-Antarctic Front, whereas silicate concentrations decrease rapidly north of the Polar Front [*Boyer et al.*, 2013]. At Kerguelen, during the recent KEOPS 2 cruise (October – November 2011) *Lasbleiz et al.* [2014] found higher concentrations of silicate south of the PF at roughly 72°E, close to the plateau.

515

One possible hypothesis is that silicate is the primary limiting factor controlling the 516 large and highly variable Kerguelen offshore bloom (i.e. longitudinally far away from the 517 plateau), both in spatial extent and inter-annual variability. This could explain why the 518 bloom is contained to the south (bloom northern limit of 44°S), where a deeper MLD can 519 mix silicate to the surface, despite the iron potentially being advected and available as far 520 north as 40°S (see Figure 5). Many previous Southern Ocean iron fertilization studies, 521 both artificial and natural, have reported the development of a large diatom bloom in 522 the fertilized patch [Blain et al., 2001; de Baar et al., 2008; Mongin et al., 2008] and 523 consequently, in the region of Kerguelen, depletion of silicate over the plateau Mosseri 524 et al., 2008]. The absence of a non-diatom bloom is explained by the efficient grazing of 525 microbial communities by copepods and salps as suggested by Banse [1996] and Smetacek 526

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et al. [2004]. However, the majority of studies have been focused on the bloom above 527 the plateau rather than further downstream, whereas the 2011 KEOPS II cruise focused 528 mainly on the bloom located just northeast of the Kerguelen Islands above the abyssal 529 plain. Their results suggest that the majority of diatom silica production during the bloom 530 event is sustained by 'new' silica, supplied primarily from pre-bloom winter water and also 531 vertical supply. As the bloom progresses, the silicon pump is strengthened by the sinking 532 of biogenic silica, and consequently the standing stock of available silica diminishes over 533 time. Estimates for the duration of the high productivity bloom period is on the order 534 of 85-86 days, after which the bloom declines [Mongin et al., 2008; Closset et al., 2014]. 535 These conditions could also be true of the far offshore bloom, which in some years extends 536 further east than 95°S (2003, in Figure 5), but further in-situ observations, of both silicate 537 concentrations and bloom composition, would be necessary to either prove or disprove this. 538 539

4.3. Can advection explain the extent of the bloom area?

Focussing now on the circulation around each island, we discuss if the modeled advection 540 can explain the spatial extent of the island blooms. Our results suggest that iron advected 541 from the Kerguelen and Heard Islands could fertilize an area which overlaps the annual 542 bloom extent, but is actually much larger than the area of the bloom. Figure 10 shows 543 that the bloom which occurs over the plateau (southeast of the Kerguelen island) is 544 predominately fertilized by iron advected from Heard island [Zhang et al., 2008]. This 545 is in agreement with a water mass path way study on the plateau using radium isotopes 546 during the 2005 KEOPS cruise, which also found the water mass on the plateau to have 547 originated from Heard island [van Beek et al., 2008]. 548

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We find that the larger bloom event, which extends as far as 100°E in some years, is mostly fertilized by iron advected from Kerguelen island (see Figure 10), in agreement with *Mongin et al.* [2009] who also performed a modeled advection study on the Kerguelen bloom.

During the recent KEOPS II cruise, iron budgets were calculated focusing on blooms 553 occurring on the plateau, and also offshore in the "plume", which show the importance 554 of a horizontal supply of iron particularly, for the offshore bloom [Bowie et al., 2015]. 555 This separation, in the fertilization of the plateau bloom and offshore bloom, is due to 556 the PF which occurs between the two islands, and flows close to the southern and eastern 557 edge of Kerguelen. Using the definition described in *Park et al.* [2014], the thick black 558 contour in Figure 10 represents the modeled location of the PF for the year 2003. The 559 general position and shape of the PF is fairly consistent each year, however the modeled 560 PF does exhibit small annual variations. In Figure 10, it is apparent that the extent of 561 the Kerguelen island trajectories, and also in 2003 the bloom, is strongly bounded (in the 562 south) by the location of the PF. 563

564

The location of the Crozet bloom was different annually, in some years propagating north west, but most frequently to the north east of the island. The fertilized patch is also predominately to the north east of Crozet, but there are exceptions in some years when small narrow currents flow north west from the island. Meridionally, both the fertilized patch and consequently bloom area occur northwards of the islands, due to the formation of a Taylor Column around the island vicinity [*Popova et al.*, 2007]. Zonally, the majority of particles are advected by water which has detrained from a branch of the

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SubAntarctic Front (SAF), and are advected eastward which corresponds with the orange 572 Argo float trajectories in Figure 5 of *Pollard et al.* [2007]. Lagrangian particles which are 573 advected westward are entrained into a secondary branch from the main SAF, which flows 574 anticyclonic around Del Cano Rise (blue drifter trajectories in Figure 5 [Pollard et al., 575 2007), before eventually turning eastward at roughly 44deg. In Figure 6, we see that 576 the years in which the fertilized patch is propagated to the west (2000, 2002, 2003 and 577 2007), the particle trajectories are from releases earlier in the year, roughly from January 578 to April. This is due to the water mass north of the island (but south of the SAF) being 579 very sluggish, resulting in particle entrainment into the anticyclonic component of the 580 SAF around Del Cano Rise, taking several months. 581

582

The model does show potential iron advection extending into all regions of the Crozet 583 annual bloom areas, however the fertilized patch in the north west was never as large as 584 the blooms which occurred in the north west. Read et al. [2007] found that sub-mesoscale 585 features were important in the development and duration of the Crozet bloom, and ac-586 counted for the bloom's "patchiness". Though the physical model used in this study 587 is at a very high resolution $(1/12^{\circ})$, it cannot reproduce the exact eddy field behavior 588 year-on-year, and therefore we do not expect the annual bloom to match the annual 589 fertilized patch. Considering the stochastic nature of eddies, we believe the model to 590 have demonstrated that the local Crozet advection is sufficient to disperse iron into all 591 annual extents of the bloom. Furthermore, the NEMO modeled Lagrangian pathways are 592 in general agreement with drifter data and also altimetry-based Lagrangian model results 593

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⁵⁹⁴ [Pollard et al., 2007; Sanial et al., 2014].

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A bloom associated specifically with iron advected from South Georgia is impossible to 596 delineate in this study, as the surrounding region is one of the most productive areas of 597 the Southern Ocean (Figure. 1a), due to various other sources of iron e.g. the Antarctic 598 Peninsula [Ardelan et al., 2008; Murphy et al., 2013]. Advection from South Georgia is 599 predominately northwards and then eastwards, joining with the ACC, overlapping with 600 the annual blooms that occur in the north easterly region of South Georgia [Korb et al., 601 2004]. There is a striking sloped western edge to both the bloom area and fertilized patch 602 in most years, caused by the position of the PF, which is bounded by the local topography 603 [Moore et al., 1999]. Between South Georgia and the PF and ACC, the modeled advection 604 was annually consistent and likely to fertilize the annually occurring bloom in this area. 605 606

4.4. Can advection explain the bloom inter-annual variability?

The area that could potentially be fertilized with iron via advection around Kerguelen 607 annually extends into a fairly consistent spatial coverage, although there are significant 608 inter-seasonal variations. Despite this, the fertilized patch was much larger than the 609 bloom area in all years of the study period, suggesting that advection alone cannot ex-610 plain the blooms inter-annual variability. Focussing on 2003, in Figure 10, we see the 611 open ocean bloom extending as far south as 60°S between a southward and then north-612 ward deviation of the PF (creating a v-shape). However in most years, the bloom area 613 does not closely match the fertilized patch, and in no years does the bloom propagate 614 as far north (bloom northern limit of 44°S) as the Lagrangian particles. Assuming that 615

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the modeled spread of Lagrangian particles is correct, this would suggest that another factor is limiting the spread of the bloom into all areas of available iron, a factor which could be the predominate driver of the inter-annual variability. This would support the theory of silicate limiting the Kerguelen bloom, but without more silicate concentration observations in the far offshore area we can only speculate.

621

The Crozet blooms during 1998 – 2007 show a high degree of inter-annual variability, 622 most frequently extending far to the east, but in some years to the west and on occasion 623 extending further north than typical. Our results find a similar degree of inter-annual 624 variability in the modeled local circulation around Crozet, both in the timing of fertil-625 ization (i.e. the speed of advection) and the extent of the fertilized patch (i.e. size and 626 direction of patch). Although the fertilized patch in our model does not closely match the 627 observed blooms, our results do suggest that iron advection could predominately control 628 the inter-annual variability seen in the Crozet bloom. 629

630

The advection of iron from South Georgia annually covers a similar region (a predominately north, then eastward flow), although there are exceptions. The timescale for fertilization is highly variable (distance travelled from the iron source out into the bloom area per month), which could have an impact on the bloom.

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4.5. Factors controlling bloom termination

As the NEMO model offers a range of diagnostics, we can also propose possible bloom termination mechanisms for each island. In the modeled MLD data, the region surround-

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ing Kerguelen does not clearly follow the north to south shallowing of the MLD in summer 638 typical of the Southern Ocean (Figure 9a). There is a very clear divide in the depths of 639 the mixed layer between the shallower north and deeper south at roughly $45^{\circ}S - 50^{\circ}S$. 640 The Kerguelen bloom is always to the south of 45° S, however in the WOA dataset there is 641 nitrate available north of this, as well as iron according to our advection results (Figure. 642 5). Additionally, the WOA nitrate concentration is still high in February, so it is unlikely 643 to be nitrate exhaustion that terminates the bloom. We conjecture that as the bloom is 644 constrained to regions with a deep mixed layer, it is dependent on a deep supply of silicate 645 as suggested by Mongin et al. [2008], and found to be the case by Closset et al. [2014]646 in the bloom just offshore of the plateau. There is partial evidence from the WOA that 647 the surface silicate concentration downstream of Kerguelen is lower in February, than in 648 the previous three months, however this is based upon very few data. Looking at data 649 from the first KEOPS cruise, figure 1 in the supplementary material of Blain et al. [2007] 650 shows the concentrations of both nitrate and silicate from locations inside and outside of 651 the bloom. It shows that inside the bloom there is no silicate but there is nitrate, whereas 652 outside the bloom there is plenty of both, suggesting silicate to be the limiting nutrient. 653 This suggests that the sampling conducted during the KEOPS II expedition close to the 654 plateau [Closset et al., 2014], needs to be repeated further downstream in future field work 655 in order to determine whether the offshore bloom has similar dynamics longitudinally. 656

The modeled monthly MLD around the Crozet region does exhibit some inter–annual variability, but typically, shallows north to south from winter into summer, and is shallower than 50 m by the end of the Crozet bloom. However the WOA climatology suggests

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there is still nitrate available in January, which suggests that iron exhaustion most likely terminates the bloom. This is supported by an experiment performed on the CROZEX cruise, where the addition of iron to an area of bloom decline resulted in the stimulation of further phytoplankton growth [*Moore et al.*, 2007].

665

The South Georgia bloom is the most variable in this study, varying in both timing and extent. The decadal average bloom period is from October to April, although it can last longer and also start earlier in some years. The nitrate concentration remains high throughout the bloom period, which suggests a deepening mixed layer being the limiting factor for the otherwise persistent South Georgia bloom. This is supported by *Korb et al.* [2008], who found evidence of a persistent supply of both macronutrients and iron, by physical processes, to the area throughout the growing season.

673

5. Conclusions

In the high nutrient, low chlorophyll Southern Ocean [Martin et al., 1990; de Baar et 674 al., 1995; Boyd et al., 2007, blooms are observed in satellite ocean color data occurring 675 annually downstream of Kerguelen, Crozet and the South Georgia islands. It is generally 676 accepted that the iron limitation prevailing across the Southern Ocean is locally overcome 677 by the horizontal advection of iron from island sources [Blain et al., 2001; Murphy et al., 678 2013; Sanial et al., 2014]. In this study, Lagrangian particle tracking, with the NEMO 679 $1/12^{\circ}$ ocean general circulation model, was used to assess whether potential iron advec-680 tion can explain the extent of the blooms, and also their inter-annual variability over 681 the period 1998 - 2007. We also use the modeled circulation and diagnostic variables to 682

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⁶⁸³ consider possible causes of bloom termination for each of the islands.

684

We find that lateral advection downstream of the Southern Ocean islands is sufficient to fertilize all areas where annual blooms can occur. The patch fertilized by iron-rich water from Kerguelen is much larger in extent than the area of the bloom, whereas the patch fertilized around Crozet is comparable in size, taking into account inter-annual variability, to the size of the bloom. The patch of water fertilized by iron-rich South Georgia sediments also closely matches with the annual bloom, however delineating a bloom associated only with South Georgia proved problematic.

692

The advection around Kerguelen was consistent in spatial extent annually, however the 693 timing of potential fertilization varied inter-seasonally across the years. This could con-694 tribute to the blooms inter-annual variability, however the results suggest that the far 695 offshore Kerguelen bloom (in some years occurring as far east as 100°E), has another 696 primary factor controlling its inter-annual variability, and we offer the hypothesis of sil-697 icate being the ultimate limiting factor on a diatom dominated Kerguelen bloom. This 698 hypothesis could be tested with in-situ nutrient sampling of the area, similar to the recent 699 KEOPS II expedition [Closset et al., 2014], or alternatively by a high-resolution coupled 700 biogeochemical model to properly resolve the key biogeochemical and physical processes. 701 The results suggest that the inter-annual variability seen in the Crozet bloom can be 702 explained by variations in the advected iron supply. The fertilized patch around South 703 Georgia was fairly consistent spatially, however it did have variations in the timing of 704 advection from the island out to the bloom site. This potentially could account for the 705

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⁷⁰⁶ inter–annual variability seen in the South Georgia bloom.

707

In assessing the possible causes of bloom termination, we find that nutrient exhaustion 708 is most likely to cause the Kerguelen and Crozet blooms to collapse (silica and iron, re-709 spectively). Whereas winter convection causing the mixed layer to deepen is most likely 710 the terminating factor of the South Georgia bloom, as physical processes maintain a con-711 tinual supply of macronutrients and iron to the area, these are unlikely to be limiting 712 [Korb et al., 2008]. Typically, both the Kerguelen and Crozet blooms end well before the 713 mixed layer begins to deepen in winter, whereas the South Georgia bloom persists for the 714 entire season until the mixed layer deepens. 715

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728 the work.

References

- ⁷³⁰ Abraham, E. R., C. S. Law, P. W. Boyd, S. J. Lavender, M. T. Maldonado, and A.
 ⁷³¹ R. Bowie (2000), Importance of stirring in the development of an iron-fertilized
 ⁷³² phytoplankton bloom, *Nature*, 407 (6805), 727-730, doi:10.1038/35037555.
- 733
- ⁷³⁴ Ardelan, M. V., O. Holm-Hansen, C. D. Hewes, C. S. Reiss, N. S. Silva, H. Dulaiova,

E. Steinnes, and E. Sakshaug (2010), Natural iron enrichment around the Antarctic Peninsula in the Southern Ocean, *Biogeosciences*, 7 (1), 11–25.

- 737
- Atkinson, A., M. J. Whitehouse, J. Priddle, G. C. Cripps, P. Ward, and M. A. Brandon
 (2001), The pelagic ecosystem of South Georgia, Antarctica, *Marine Ecology Progress Series*, 216, 279 308.
- 741

SSALTO/DUACS User Handbook: (M)SLA and (M)ADT Near-Aviso (2014) 742 Real Time and Delayed Time Products, CLS-DOS-NT-06-034, SALP-743 MU–P–EA–21065–CLS, CNES, Ramonville St. Agne, France, ,59pp, (http: 744 //www.aviso.altimetry.fr/fileadmin/documents/data/tools/hdbk_duacs.pdf). 745 746

⁷⁴⁷ Bakker, D. C., M. C. Nielsdottir, P. J. Morris, H. J. Venables, and A. J. Watson (2007),
⁷⁴⁸ The island mass effect and biological carbon uptake for the subantarctic Crozet

DRAFT

⁷⁴⁹ Archipelago, Deep Sea Research II, 54, 2174 – 2190, doi:10.1016/j.dsr2.2007.06.009.

750

⁷⁵¹ Banse, K. (1996), Low seasonality of low concentrations of surface chlorophyll in
 ⁷⁵² the Subantarctic water ring: underwater irradiance, iron, or grazing?, *Progress in* ⁷⁵³ Oceanography, 37 (3-4), 241-291.

754

⁷⁵⁵ Blain, S., et al. (2001), A biogeochemical study of the island mass effect in the context
⁷⁵⁶ of the iron hypothesis: Kerguelen Islands, Southern Ocean, *Deep–Sea Research Part I–*⁷⁵⁷ Oceanographic Research Papers, 48 (1), 163–187, doi:10.1016/S0967–0637(00)00047–9.

⁷⁵⁹ Blain, S., et al. (2007), Effect of natural iron fertilization on carbon sequestration in the
⁷⁶⁰ Southern Ocean, *Nature*, 446 (7139), 1070–1074, doi:10.1038/nature05700.

761

⁷⁶² Blain, S., B. Quguiner, and T. Trull (2008), The natural iron fertilization exper⁷⁶³ iment KEOPS (KErguelen Ocean and Plateau compared Study): An overview,
⁷⁶⁴ Deep Sea Research Part II: Topical Studies in Oceanography, 55 (5–7), 559–565,
⁷⁶⁵ doi:10.1016/j.dsr2.2008.01.002.

766

⁷⁶⁷ Blanke, B., and S. Raynaud (1997), Kinematics of the Pacific Equatorial Undercurrent:
 ⁷⁶⁸ An Eulerian and Lagrangian approach from GCM results, *Journal of Physical Oceanog-*

 $_{769}$ raphy, 27 (6), 1038–1053, doi:10.1175/1520–0485.

770

Х - З	38
-------	----

⁷⁷¹ Blanke, B., M. Arhan, G. Madec, and S. Roche (1999), Warm water paths in the
⁷⁷² equatorial Atlantic as diagnosed with a general circulation model, *Journal of Physical*⁷⁷³ Oceanography, 29, 2753?2768.

774

Borrione, I., and R. Schlitzer (2013), Distribution and recurrence of phytoplankton
blooms around South Georgia, Southern Ocean, *Biogeosciences*, 10 (1), 217–231,
doi:10.5194/bg-10-217-2013.

778

Bowie, A. R., et al. (2015), Iron budgets for three distinct biogeochemical sites around
the Kerguelen Archipelago (Southern Ocean) during the natural fertilization study,
KEOPS-2, *Biogeosciences*, 12 (14), 4421-4445, doi:10.5194/bg-12-4421-2015.

782

Boyd, P. W., et al. (2000), A mesoscale phytoplankton bloom in the polar Southern Ocean
stimulated by iron fertilization, *Nature*, 407 (6805), 695-702, doi:10.1038/35037500.

Boyd, P. W. (2002), Environmental Factors Controlling Phytoplankton Processes in
the Southern Ocean, Journal of Phycology, 38 (5), 844-861, doi:10.1046/j.15298817.2002.t01-1-01203.x.

789

Boyd, P. W. (2007), Biogeochemistry: iron findings, Nature, 446 (7139), 989-991,
 doi:10.1038/446989a.

792

- Boyd, P. W., et al. (2007), Mesoscale iron enrichment experiments 1993–2005: syn thesis and future directions, *Science*, *315* (5812), 612–617, doi:10.1126/science.1131669.
- ⁷⁹⁶ Boyd, P. W., and M. J. Ellwood (2010), The biogeochemical cycle of iron in the ocean, ⁷⁹⁷ Nature Geoscience, 3 (10), 675-682, doi:10.1038/ngeo964.
- 798
- Boyd, P. W., K. R. Arrigo, R. Strzepek, and G. L. van Dijken (2012), Mapping phytoplankton iron utilization: Insights into Southern Ocean supply mechanisms, *Journal of Geophysical Research: Oceans*, 117 (C6), C06009, doi:10.1029/2011JC007726.
- 802
- Boyd, P. W., D. C. E. Bakker, and C. Chandler (2012), A New Database to Explore the
 Findings from Large–Scale Ocean Iron Enrichments Experiments, *Oceanography*, 25
 (4), 64–71, doi: http://dx.doi.org/10.5670/oceanog.2012.104.
- 806
- Boyer, T.P., J. I. Antonov, O. K. Baranova, C. Coleman, H. E. Garcia, A. Grodsky, D. R.
 Johnson, R. A. Locarnini, A. V. Mishonov, T.D. O'Brien, C.R. Paver, J.R. Reagan, D.
 Seidov, I. V. Smolyar, and M. M. Zweng, 2013: World Ocean Database 2013, NOAA
 Atlas NESDIS 72, S. Levitus, Ed., A. Mishonov, Technical Ed.; Silver Spring, MD,
 209, http://doi.org/10.7289/V5NZ85MT.
- 812
- Brodeau, L., B. Barnier, A.-M.Treguier, T. Penduff, and S. Gulev (2010) An ERA40–
 based 461 atmospheric forcing for global ocean circulation models. *Ocean Model. 31*,
 88–104.

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X - 40

⁸¹⁷ Closset, I., M. Lasbleiz, K. Leblanc, B. Quguiner, A. J. Cavagna, M. Elskens, J. Navez, ⁸¹⁸ and D. Cardinal (2014), Seasonal evolution of net and regenerated silica production ⁸¹⁹ around a natural Fe-fertilized area in the Southern Ocean estimated with Si isotopic ⁸²⁰ approaches, *Biogeosciences*, *11* (20), 5827-5846, doi:10.5194/bg-11-5827-2014.

821

816

⁸²² Comiso, J. C., C. R. McClain, C. W. Sullivan, J. P. Ryan, and C. L. Leonard (1993),
⁸²³ Coastal zone color scanner pigment concentrations in the Southern Ocean and rela⁸²⁴ tionships to geophysical surface features, *Journal of Geophysical Research*, 98 (C2),
⁸²⁵ 2419, doi:10.1029/92jc02505.

826

de Baar, H. J., J. de Jong, T. M., D. C. Bakker, B. M. Loscher, C. Veth, U. Bathmann, and V. Smetacek (1995), Importance of iron for plankton blooms and carbon
dioxide drawdown in the Southern Ocean, *Nature*, 373, 412 – 415, doi:10.1038/373412a0.

de Baar, H. J. W., L. J. A. Gerringa, P. Laan, and K. R. Timmermans (2008), Efficiency
 of carbon removal per added iron in ocean iron fertilization, *Marine Ecology Progress Series*, 364, 269–282, doi:10.3354/meps07548.

834

d'Ovidio, F., A. Della Penna, T. W. Trull, F. Nencioli, M. I. Pujol, M. H. Rio, Y. H.
Park, C. Cott, M. Zhou, and S. Blain (2015), The biogeochemical structuring role
of horizontal stirring: Lagrangian perspectives on iron delivery downstream of the
Kerguelen Plateau, *Biogeosciences*, 12 (19), 5567-5581, doi:10.5194/bg-12-5567-2015.

DRAFT

April 13, 2016, 2:40pm

8	3	9

- Gille, S. T., M. M. Carranza, and R. Cambra (2014), Wind-induced upwelling in the
 Kerguelen Plateau region, *Biogeosciences*, 11 (22), 6389-6400, doi:10.5194/bg-11-63892014.
- 843
- Graham, R. M., A. M. De Boer, E. van Sebille, K. E. Kohfeld, and C. Schlosser (2015),
 Inferring source regions and supply mechanisms of iron in the Southern Ocean from
 satellite chlorophyll data, *Deep Sea Research Part I: Oceanographic Research Papers*,
 104, 9-25, doi:10.1016/j.dsr.2015.05.007.
- 848
- Hewes, C. D., C. S. Reiss, M. Kahru, B. G. Mitchell, and O. Holm–Hansen (2008), Control
 of phytoplankton biomass by dilution and mixed layer depth in the western Weddell–
 Scotia Confluence, *Marine Ecology Progress Series*, *366*, 15–29, doi:10.3354/meps07515.
- Holm-Hansen, O., M. Kahru, and C. D. Hewes (2005), Deep chlorophyll a maxima
 (DCMs) in pelagic Antarctic waters. II. Relation to bathymetric features and dissolved
 iron concentrations, *Marine Ecology Progress Series*, 297, 71–81.
- 856

Korb, R. E., M. J. Whitehouse, and P. Ward (2004), SeaWiFS in the southern
ocean: spatial and temporal variability in phytoplankton biomass around South
Georgia, *Deep Sea Research Part II: Topical Studies in Oceanography*, 51, 99–116,
doi:http://dx.doi.org/10.1016/j.dsr2.2003.04.002.

861

Korb, R. E., and M. Whitehouse (2004), Contrasting primary production regimes around
South Georgia, Southern Ocean: large blooms versus high nutrient, low chlorophyll
waters, *Deep Sea Research Part I: Oceanographic Research Papers*, 51 (5), 721–738,
doi:10.1016/j.dsr.2004.02.006.

866

Korb, R. E., M. J. Whitehouse, A. Atkinson, and S. E. Thorpe (2008), Magnitude and
maintenance of the phytoplankton bloom at South Georgia: a naturally iron-replete
environment, *Marine Ecology Progress Series*, 368, 75-91, doi:10.3354/meps07525.

870

Lasbleiz, M., K. Leblanc, S. Blain, J. Ras, V. Cornet–Barthaux, S. Hlias Nunige, and
B. Quguiner (2014), Pigments, elemental composition (C, N, P, and Si), and stoichiometry of particulate matter in the naturally iron fertilized region of Kerguelen in
the Southern Ocean, *Biogeosciences*, 11 (20), 5931–5955, doi:10.5194/bg-11–5931–2014.

Le Bars, D., J. V. Durgadoo, H. A. Dijkstra, A. Biastoch, and W. P. M. De Ruijter (2014), An observed 20-year time series of Agulhas leakage, *Ocean Science*, 10 (4), 601–609, doi:10.5194/os–10–601–2014.

879

Madec, G., (2008), NEMO reference manual, ocean dynamic component: NEMO–OPA,
Notes du pole de modelisation, Tech. Rep. 27, Institut Pierre Simmon Laplace (IPSL),
Paris, France.

883

- Martin, J. H., R. M. Gordon, and S. E. Fitzwater (1990), Iron in Antarctic waters,
 Nature, 345 (6271), 156–158, doi:10.1038/345156a0.
- 886
- Martin, J. M. (1990), Glacial-interglacial CO2 change: the iron hypothesis, *Paleoceanography*, 5 (1), 1–13.
- 889

893

- Marzocchi A., Hirschi J. J.–M., Holliday N.P., Cunningham S.A., Blaker A.T. and Coward
 A.C. (2015) The North Atlantic subpolar circulation in an eddy–resolving global ocean
 model, *Journal of Marine Systems*, 142, 126–143, doi:10.1016/j.jmarsys.2014.10.007.
- Meijers, A. J. S., A. Klocker, N. L. Bindoff, G. D. Williams, and S. J. Marsland (2010),
 The circulation and water masses of the Antarctic shelf and continental slope between
 30 and 80degE, *Deep Sea Research Part II: Topical Studies in Oceanography*, 57 (9–10),
 723–737, doi:10.1016/j.dsr2.2009.04.019.
- 898
- Meredith, M. P., J. L. Watkins, E. J. Murphy, N. J. Cunningham, A. G. Wood, R. Korb,
 M. J. Whitehouse, S. E. Thorpe, and F. Vivier (2003), An anticyclonic circulation above the Northwest Georgia Rise, Southern Ocean, Geophysical Research Letters, 30 (20), doi:10.1029/2003gl018039.
- 903
- Mongin, M., E. Molina, and T. W. Trull (2008), Seasonality and scale of the Kerguelen
 plateau phytoplankton bloom: A remote sensing and modeling analysis of the influence
 of natural iron fertilization in the Southern Ocean, *Deep Sea Research Part II: Topical*

DRAFT

X - 44

- ⁹⁰⁷ Studies in Oceanography, 55 (5–7), 880–892, doi:10.1016/j.dsr2.2007.12.039.
- 908

Mongin, M. M., E. R. Abraham, and T. W. Trull (2009), Winter advection of iron can explain the summer phytoplankton bloom that extends 1000 km downstream of the Kerguelen Plateau in the Southern Ocean, Journal of Marine Research, 67, 225–237.

- ⁹¹³ Moore, J. K., M. R. Abbott, and J. G. Richman (1999), Location and dynamics of ⁹¹⁴ the Antarctic Polar Front from satellite sea surface temperature data, *Journal of* ⁹¹⁵ *Geophysical Research*, 104 (C2), 3059, doi:10.1029/1998jc900032.
 - 916

912

⁹¹⁷ Moore, J. K., and M. R. Abbott (2000), Phytoplankton chlorophyll distributions and ⁹¹⁸ primary production in the Southern Ocean, *Journal of Geophysical Research*, 105 ⁹¹⁹ (C12), 28709, doi:10.1029/1999jc000043.

920

Moore, C. M., S. Seeyave, A. E. Hickman, J. T. Allen, M. I. Lucas, H. Planquette,
R. T. Pollard, and A. J. Poulton (2007), Iron–light interactions during the CROZet
natural iron bloom and EXport experiment (CROZEX) I: Phytoplankton growth and
photophysiology, *Deep Sea Research Part II: Topical Studies in Oceanography*, 54
(18–20), 2045–2065, doi:10.1016/j.dsr2.2007.06.011.

926

⁹²⁷ Mosseri, J., B. Quguiner, L. Armand, and V. Cornet–Barthaux (2008), Impact of iron ⁹²⁸ on silicon utilization by diatoms in the Southern Ocean: A case study of Si/N cycle ⁹²⁹ decoupling in a naturally iron–enriched area, *Deep Sea Research Part II: Topical*

DRAFT

Murphy, E. J., et al. (2013), Comparison of the structure and function of Southern Ocean

- ⁹³⁰ Studies in Oceanography, 55 (5–7), 801–819, doi:10.1016/j.dsr2.2007.12.003.
- 931

932

- regional ecosystems: The Antarctic Peninsula and South Georgia, Journal of Marine Systems, 109–110, 22–42, doi:10.1016/j.jmarsys.2012.03.011.
 Orsi, A. H., T. Whitworth III, and W. D. Nowlin Jr (1995), On the meridional extent and fronts of the Antarctic Circumpolar Current, Deep Sea Research I, 42 (5), 641-673.
 Park, Y.-H., J.-L. Fuda, I. Durand, and A. C. Naveira Garabato (2008), Internal tides
- and vertical mixing over the Kerguelen Plateau, Deep Sea Research Part II: Topical
 Studies in Oceanography, 55 (5-7), 582-593, doi:10.1016/j.dsr2.2007.12.027.
- 942
- Park, Y.-H., F. Roquet, I. Durand, and J.-L. Fuda (2008), Large-scale circulation over
 and around the Northern Kerguelen Plateau, *Deep Sea Research Part II: Topical*Studies in Oceanography, 55 (5–7), 566–581, doi:10.1016/j.dsr2.2007.12.030.
- 946

Park, Y.-H., I. Durand, E. Kestenare, G. Rougier, M. Zhou, F. d'Ovidio, C. Cott,
and J.-H. Lee (2014), Polar Front around the Kerguelen Islands: An up-to-date
determination and associated circulation of surface/subsurface waters, *Journal of Geophysical Research: Oceans*, 119, doi:10.1002/2014jc010061.

951

X - 46

ROBINSON ET AL.: A TALE OF THREE ISLANDS

- Planquette, H., et al. (2007), Dissolved iron in the vicinity of the Crozet Islands, Southern
 Ocean, Deep Sea Research II, 54, 1999 2019, doi:10.1016/j.dsr2.2007.06.019.
- 954

Pollard, R., R. Sanders, M. Lucas, and P. Statham (2007), The Crozet Natural Iron
Bloom and Export Experiment (CROZEX), *Deep Sea Research Part II: Topical Studies in Oceanography*, 54 (18–20), 1905–1914, doi:10.1016/j.dsr2.2007.07.023.

958

Pollard, R. T., H. J. Venables, J. F. Read, and J. T. Allen (2007), Large–scale
circulation around the Crozet Plateau controls an annual phytoplankton bloom in
the Crozet Basin, *Deep Sea Research II*, 54, 1915 – 1929, doi:10.1016/j.dsr2.2007.06.012.

962

Pollard, R. T., et al. (2009), Southern Ocean deep-water carbon export enhanced by
natural iron fertilization, *Nature*, 457 (7229), 577-580, doi:10.1038/nature07716.

Popova, E. E., R. T. Pollard, M. I. Lucas, H. J. Venables, and T. R. Anderson
(2007), Real-time forecasting of ecosystem dynamics during the CROZEX experiment and the roles of light, iron, silicate, and circulation, *Deep Sea Research Part II: Topical Studies in Oceanography*, 54 (18-20), 1966-1988, doi:10.1016/j.dsr2.2007.06.018.

Popova, E. E., A. Yool, Y. Aksenov, and A. C. Coward (2013), Role of advection in
Arctic Ocean lower trophic dynamics: A modeling perspective, *Journal of Geophysical Research*, 118, 1571–1586, doi:10.1002/jgrc.20126.

974

- Poulton, A. J., C. Mark Moore, S. Seeyave, M. I. Lucas, S. Fielding, and P. Ward (2007),
 Phytoplankton community composition around the Crozet Plateau, with emphasis on
 diatoms and Phaeocystis, *Deep Sea Research Part II: Topical Studies in Oceanography*,
 54 (18-20), 2085-2105, doi:10.1016/j.dsr2.2007.06.005.
- 979
- Read, J. F., R. T. Pollard, and J. T. Allen (2007), Sub-mesoscale structure and the
 development of an eddy in the Subantarctic Front north of the Crozet Islands, *Deep-Sea Research Part Ii-Topical Studies in Oceanography*, 54 (18–20), 1930–1948, doi:Doi
 10.1016/J.Dsr2.2007.06.013.
- 984
- Robinson, J., E. E. Popova, A. Yool, M. Srokosz, R. S. Lampitt, and J. R. Blundell
 (2014), How deep is deep enough? Ocean iron fertilization and carbon sequestration in the Southern Ocean, *Geophysical Research Letters*, 41 (7), 2489–2495,
 doi:10.1002/2013gl058799.
- 989
- Sanial, V., P. van Beek, B. Lansard, F. d'Ovidio, E. Kestenare, M. Souhaut, M. Zhou,
 and S. Blain (2014), Study of the phytoplankton plume dynamics off the Crozet Islands
 (Southern Ocean): A geochemical-physical coupled approach, *Journal of Geophysical Research-Oceans*, 119 (4), 2227–2237, doi:10.1002/2013jc009305.
- 994
- Shaked, Y., and H. Lis (2012), Disassembling iron availability to phytoplankton, Frontiers
 in Microbiology, 3 (123), doi:10.3389/fmicb.2012.00123.
- 997

X - 48

Schallenberg, C., P. van der Merwe, F. Chever, J. T. Cullen, D. Lannuzel, and A. R.
Bowie (2015), Dissolved iron and iron(II) distributions beneath the pack ice in the
East Antarctic (120E) during the winter/spring transition, *Deep Sea Research Part II:*Topical Studies in Oceanography, doi:10.1016/j.dsr2.2015.02.019.

1002

- Smetacek, V., P. Assmy, and J. Henjes (2004), The role of grazing in structuring
 Southern Ocean pelagic ecosystems and biogeochemical cycles, *Antarctic Science*, 16
 (4), 541-558, doi:10.1017/s0954102004002317.
- 1006
- Srokosz, M. A., J. Robinson, H. McGrain, E. E. Popova, and A. Yool (2015), Could the
 Madagascar bloom be fertilized by Madagascan iron?, *Journal of Geophysical Research: Oceans*, 120, doi:10.1002/2015JC011075.
- 1010
- Storm, T., M. Boettcher, M. Grant, M. Zuhlke, F. N., T. Jackson, and S. Sathyendranath
 (2013), Product User Guide, in *Ocean Colour Climate Change Initiative (OC_CCI) – Phase One*, edited by S. Groom, Plymouth Marine Laboratory, Plymouth, UK.
- Tagliabue, A., J.-B. Salle, A. R. Bowie, M. Lvy, S. Swart, and P. W. Boyd (2014),
 Surface-water iron supplies in the Southern Ocean sustained by deep winter mixing,
 Nature Geoscience, 7 (4), 314-320, doi:10.1038/ngeo2101.
- 1018
- ¹⁰¹⁹ Takahashi, T., et al. (2009), Climatological mean and decadal change in surface ¹⁰²⁰ ocean pCO2, and net sea-air CO2 flux over the global oceans, *Deep Sea Research Part*

DRAFT

¹⁰²¹ *II: Topical Studies in Oceanography*, *56* (8-10), 554-577, doi:10.1016/j.dsr2.2008.12.009.

- ¹⁰²³ Thomalla, S. J., N. Fauchereau, S. Swart, and P. M. S. Monteiro (2011), Regional ¹⁰²⁴ scale characteristics of the seasonal cycle of chlorophyll in the Southern Ocean, ¹⁰²⁵ *Biogeosciences*, 8 (10), 2849–2866, doi:10.5194/bg-8–2849–2011.
- 1026
- Tripathy, S. C., S. Pavithran, P. Sabu, H. U. K. Pillai, D. R. G. Dessai, and N. Anilkumar
 (2015), Deep chlorophyll maximum and primary productivity in Indian Ocean sector
 of the Southern Ocean: Case study in the Subtropical and Polar Front during austral
 summer 2011, Deep Sea Research Part II: Topical Studies in Oceanography, 118,
 240-249, doi:10.1016/j.dsr2.2015.01.004.
- 1032

Tyrrell, T., A. Merico, J. J. Waniek, C. S. Wong, N. Metzl, and F. Whitney (2005),
 Effect of seafloor depth on phytoplankton blooms in high-nitrate, low-chlorophyll
 (HNLC) regions, *Journal of Geophysical Research*, 110 (G2), doi:10.1029/2005jg000041.

1036

van Beek, P., M. Bourquin, J. L. Reyss, M. Souhaut, M. A. Charette, and C. Jeandel
(2008), Radium isotopes to investigate the water mass pathways on the Kerguelen
Plateau (Southern Ocean), *Deep Sea Research Part II: Topical Studies in Oceanogra- phy*, 55 (5–7), 622–637, doi:10.1016/j.dsr2.2007.12.025.

1041

van der Merwe, P., et al. (2015), Sourcing the iron in the naturally fertilised bloom
around the Kerguelen Plateau: particulate trace metal dynamics, *Biogeosciences*, 12

DRAFT

X - 50

(3), 739-755, doi:10.5194/bg-12-739-2015.

1045

1044

- Venables, H. J., R. T. Pollard, and E. E. Popova (2007), Physical conditions controlling
 the development of a regular phytoplankton bloom north of the Crozet Plateau,
 Southern Ocean, *Deep Sea Research II*, 54, 1949 1965, doi:10.1016/j.dsr2.2007.06.014.
- 1049
- Venables, H., and C. M. Moore (2010), Phytoplankton and light limitation in the
 Southern Ocean: Learning from high–nutrient, high–chlorophyll areas, Journal of
 Geophysical Research, 115 (C2), doi:10.1029/2009jc005361.
- 1053
- Wadley, M. R., T. D. Jickells, and K. J. Heywood (2014), The role of iron sources and
 transport for Southern Ocean productivity, *Deep Sea Research Part I: Oceanographic Research Papers*, doi:10.1016/j.dsr.2014.02.003.
- 1057
- Ward, P., M. Whitehouse, R. Shreeve, S. Thorpe, A. Atkinson, R. Korb, D. Pond, and
 E. Young (2007), Plankton community structure south and west of South Georgia
 (Southern Ocean): Links with production and physical forcing, *Deep Sea Research I*,
 54, 1871–1889, doi:10.1016/j.dsr.2007.08.008.
- 1062
- Young, E. F., M. P. Meredith, E. J. Murphy, and G. R. Carvalho (2011), High–resolution
 modelling of the shelf and open ocean adjacent to South Georgia, Southern Ocean,
 Deep Sea Research II, 58, 1540–1552, doi:10.1016/j.dsr2.2009.11.003.

1066

Young, E. F., S. E. Thorpe, N. Banglawala, and E. J. Murphy (2014), Variability
 in transport pathways on and around the South Georgia shelf, Southern Ocean:
 Implications for recruitment and retention, *Journal of Geophysical Research*, 119, 241–252, doi:10.1002/2013JC009348.

- 1071
- ¹⁰⁷² Zhang, Y., F. Lacan, and C. Jeandel (2008), Dissolved rare earth elements tracing
 ¹⁰⁷³ lithogenic inputs over the Kerguelen Plateau (Southern Ocean), *Deep Sea Research Part*
- ¹⁰⁷⁴ II: Topical Studies in Oceanography, 55 (5–7), 638–652, doi:10.1016/j.dsr2.2007.12.029.
- 1075



Figure 1. A Southern Ocean overview of satellite ocean color, satellite and modeled surface current speed, and the bathymetry in the model. Panel [a] is a decadal average, 1998 - 2007, of the chlorophyll-a concentration [mg m⁻³] in the month of November. Panels [b] and [c] are the decadal averages (1998 - 2007) of surface current speed [m^{-s}], from the NEMO model, at $1/12^{\circ}$, and the Aviso data, at $1/4^{\circ}$ resolution, respectively. Panel [d] is the Southern Ocean bathymetry D R A F T April 13, 2016, 2:40pm D R A F T within the NEMO 1/12° model, contours are in meters below the sea surface. Black boxes denote the study areas: South Georgia left, Crozet islands middle, Kerguelen right.



Figure 2. The starting positions of the Lagrangian particles around the islands. Particles are placed over shallow bathymetry (< 180 m), around Kerguelen and Heard Island, Crozet Islands, and South Georgia and Shag Rocks; plots [a], [b], and [c] respectively. In plots [a] and [c], only every other particle is plotted for clarity. The plot also includes contours of 500 m (dark blue), 1500 m (green) and 3000 m (red) isobaths. Note that the axis for each panel are not consistent.



Figure 3. The average chlorophyll–a concentration $[mg m^{-3}]$ (satellite ocean color) of each month over the 10 year period. Concentrations are from two locations, one inside (thick green line with markers) and one outside of the bloom region (dashed green line), for each island. The data points from inside the bloom region include error bars which are plus and minus one standard deviation in chlorophyll–a for each month, over the ten year period. The blue line represents the decadal average of the mixed layer depth of each location inside the bloom. Panels [a], [b], and [c] are Kerguelen, Crozet and South Georgia. Note the x axis, 'Month', begins from June through to May.



Figure 4. Example years of satellite ocean color plots of each island. Chlorophyll–a concentrations have been averaged over the bloom period for each year. The top row is Kerguelen (bloom period: Nov – Jan), years 2000 ([a]) and 2003 ([b]); the middle row is Crozet (bloom period: Oct – Dec), years 2001 ([c]) and 2004 ([d]); and the bottom row is South Georgia (bloom period: Oct – Apr), years 2006 ([e]) and 2002 ([f]). Panels [a], [c], and [e] are examples of a small bloom extent during the 1998 – 2007 year period, and panels [b], [d], and [f] are years with a large bloom extent.



Figure 5. Extent of Lagrangian trajectories around Kerguelen. 8240 particles were released monthly from their starting positions, denoted in blue, however only every second particle is shown here for clarity. Particle trajectories in October (preceding the start of the bloom), are depicted by colored markers. The color of the trajectory relates to the month in which it was released as indicated by the color bar. The black contour represents the averaged bloom area, over November – January, of chlorophyll–a concentrations above 0.5 mg m⁻³. Only trajectories that are shallower than 200m are included in this plot.



Figure 6. Extent of Lagrangian trajectories around Crozet. 465 particles were released monthly from their sing positions, denoted in blue, however only every second particle is shown here for clarity. Trajectories in September (preceding the start of the bloom), are depicted by colored markers. The color of the trajectory relates to the month in which it was released as indicated by the color bar. The black contour represents the averaged bloom area, over October – December, of chlorophyll–a concentrations above 0.5 mg m⁻³. Only trajectories that are shallower than 200m are included in this plot.



Figure 7. Extent of Lagrangian trajectories around South Georgia. 2820 particles were released monthly from their starting positions, denoted in blue, however only every second particle is shown here for clarity. Trajectories, in October (preceding the start of the bloom), are depicted by colored markers. The color of the trajectory relates to the month in which it was released as indicated by the colorbar. The black contour represents the averaged bloom area, over October – April, of chlorophyll–a concentrations above 0.5 mg m⁻³. Only trajectories that are shallower than 200m are included in this plot.

	Bloom Area	Fertilized patch	Bloom Overlap	Fertilized Overlap
Year	(km^2)	(km^2)	(%)	(%)
1998	674572	2731476	74	18
1999	803847	2676533	64	19
2000	339783	2693883	78	10
2001	832765	2593000	84	27
2002	613450	2621786	75	18
2003	1602173	2435464	73	48
2004	729515	2637034	72	20
2005	532218	2568354	84	17
2006	1056154	2539699	82	34

Size of the annual bloom and fertilized patch around Kerguelen, and the percent of Table 1 eac

 $^{\mathrm{a}}$ Bloom Area is the total area of the average (November to January) chl-a concentration above 0.5 mg m⁻³; Fertilized patch is the extent of particle trajectories in October (prior to the start of the bloom); Bloom Overlap is the percent of the bloom area overlapped by the fertilized patch; Fertilized Overlap is the percent of the fertilized patch overlapped by the bloom

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Table 2. Size of the annual bloom and fertilized patch around Crozet, and the percent of each area that is overlapped by the other. ^a

	Bloom Area	Fertilized patch	Bloom Overlap	Fertilized Overlap	
Year	(km^2)	(km^2)	(%)	(%)	
1998	223784	216357	35	36	
1999	300810	251913	52	62	
2000	168707	259734	59	38	
2001	70586	231342	60	18	
2002	195589	238834	42	34	
2003	299167	232525	41	52	
2004	355097	252176	48	67	
2005	258354	246721	54	56	
2006	209982	244683	47	40	
2007	342084	282999	32	39	
^a Bloom Area	is the total are	a of the average (October to Decen	nber) chl–a concentration a	bove

 0.5 mg m^{-3} ; Fertilized patch is the extent of particle trajectories in September (prior to the start of the bloom); Bloom Overlap is the percent of the bloom area overlapped by the fertilized patch; Fertilized Overlap is the percent of the fertilized patch overlapped by the bloom

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	Bloom Area	Fertilized patch	Bloom Overlap	Fertilized Overlap
Year	(km^2)	(km^2)	(%)	(%)
1998	527788	801679	79	52
1999	671337	864382	70	54
2000	453359	810443	79	44
2001	560929	798532	74	52
2002	946833	704231	50	67
2003	854692	750085	56	64
2004	587281	713427	69	57
2005	626285	700466	66	59
2006	414108	703676	56	33
2007	543833	573456	61	58

Size of the annual bloom and fertilized patch around South Georgia, and the percent Table 3 0

^a Bloom Area is the total area of the average (October to April) chl–a concentration above 0.5 mg m $^{-3}$; Fertilized patch is the extent of particle trajectories in September (prior to the start of the bloom); Bloom Overlap is the percent of the bloom area overlapped by the fertilized patch; Fertilized Overlap is the percent of the fertilized patch overlapped by the bloom



Figure 8. The percent of the bloom area overlapped by Lagrangian trajectories from each monthly release for each year. For each monthly release of particles, trajectories that were within the bloom area, in the month that is prior to the start of the bloom, were recorded and used to calculate the percentage area coverage of the bloom by Lagrangian trajectories. Any particles deeper than 200 m were not included. The y axis, % bloom overlap, indicates the percentage of the bloom area overlapped by particles from each monthly release shown on the x axis, *Particle release month*. Each colored line represents an individual year.



Figure 9. Monthly climatologies (decadal, 1998 – 2007) of the modeled mixed layer depth, calculated online by the NEMO model, around Kerguelen, Crozet and South Georgia. The color scale is m below the surface, with warm colors indicating shallow depths and cold colors representing deeper depths.



Figure 10. Lagrangian trajectories originating from the Kerguelen and Heard Islands in 2003. Collectively, 8240 particles were released monthly from their starting positions, denoted in blue, however only every second particle is shown here for clarity. Particle trajectories are depicted by colored markers, with the color of the marker relating to the month in which it was released as indicated by the color bar. Gray hatching represents the bloom area, averaged over November – January, where chlorophyll–a concentrations are above 0.5 mg m⁻³. The thick black contour represents the approximate location of the Polar Front in the model for 2003. Only trajectories that are shallower than 200m are included in this plot.