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# Regional scale characteristics of the seasonal cycle of chlorophyll in the Southern Ocean

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#### Abstract

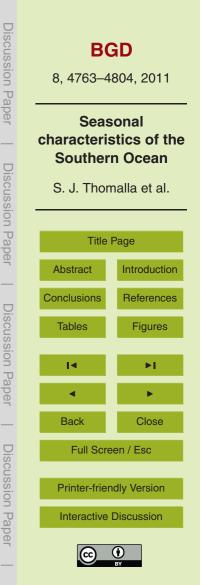
The seasonal cycle is the mode that couples climate forcing to ecosystem production. A better understanding of the regional characteristics of the seasonal cycle addresses an important gap in our understanding of the sensitivity of the biological pump

- to climate change. The regional characteristics of the seasonal cycle of phytoplankton biomass in the Southern Ocean were examined in terms of the timing of the bloom initiation, its amplitude, regional scale variability and the importance of the climatological seasonal cycle in explaining the overall variance. The study highlighted important differences between the spatial distribution of satellite observed phytoplankton biomass
- and the more dynamically linked characteristics of the seasonal cycle. The seasonal cycle was consequently defined into four broad zonal regions; the subtropical zone (STZ), the transition zone (TZ), the Antarctic circumpolar zone (ACZ) and the marginal ice zone (MIZ). Defining the Southern Ocean according to the characteristics of its seasonal cycle provides a more dynamic understanding of ocean productivity based
- on underlying physical drivers rather than climatological biomass. The response of the biology to the underlying physics of the different seasonal zones resulted in an additional classification of four regions based on the extent of interannual seasonal phase locking and the amplitude of the integrated seasonal biomass. This characterisation contributes to an improved understanding of regional sensitivity to climate forcing potentially allowing more rebust predictions of long term climate trends.
- <sup>20</sup> tentially allowing more robust predictions of long term climate trends.

## 1 Introduction

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Biological production and carbon export to the deep ocean, "the biological pump" is considered a major contributor to the Southern Ocean  $CO_2$  sink removing an estimated 3 PgC ( $1 PgC = 10^{15} g$  of carbon) from surface waters south of  $30^{\circ}$  S each year (33 % of the global organic carbon flux) (Schlitzer et al., 2002). The Southern Oceans biological pump also plays an important role in regulating the supply of nutrients to



thermocline waters (Subantarctic Mode Water and Intermediate Water) of the entire Southern Hemisphere and North Atlantic (Sarmiento et al., 2004), which in turn drives low latitude productivity (Sigman and Boyle, 2000). Climate models and decadal data sets suggest an increase in mixed layer stratification of the Southern Ocean through in-

- <sup>5</sup> creased freshening and a shift towards a positive phase of the Southern Annular Mode (SAM) causing a poleward intensification of the westerly winds (Gille, 2008; Böning et al., 2008; Le Queré et al., 2009; Turner et al., 2009). Such trends will impact the Southern Ocean carbon sink and highlights gaps in our understanding about the regional character of the sensitivity of biological production in the Southern Ocean to
   <sup>10</sup> changes in spatial and temporal atmospheric forcing scales (Sarmiento et al., 1998;
- Caldeira and Duffy, 2000; Russell et al., 2006; Arrigo et al., 2008).

A significant part of these gaps lies in the links between physical, physiological and ecological factors that influence the annual cycle of primary production growth and loss rates (Behrenfeld, 2010). The Southern Ocean is an unusual ocean in that it has the

- <sup>15</sup> greatest inventory of unused macro-nutrients in the world ocean (Levitus et al., 1993) but low average phytoplankton standing stocks, with diverse spatial and temporal variations in phytoplankton biomass (Sullivan et al., 1993; Arrigo et al., 2008). Numerous studies in the literature have addressed the factors governing phytoplankton distribution, diversity, biomass and production. These include both bottom-up controls of the
- <sup>20</sup> physiological response of phytoplankton assemblages to physical and biogeochemical forcing (e.g., Martin et al., 1990; Cullen, 1991; Nelson and Smith, 1991; de Baar et al., 1995; Boyd, 2002) as well as top-down controls of grazing (e.g., Cullen 1991; Price et al., 1994; Smetacek et al., 2004; Behrenfeld, 2010). Moreover, given the known influence of temperature on phytoplankton growth (e.g., Raven and Geider, 1988) and
- <sup>25</sup> photosynthesis (e.g., review by Davidson, 1991) it is not surprising that the cold temperatures of the Southern Ocean also effect phytoplankton biomass and the seasonal cycle. However, despite the effects of cold temperatures on biological processes in the Southern Ocean, the key factors exerting bottom up control of phytoplankton production in this, the largest "high nitrate-low chlorophyll (HNLC)" region in the world



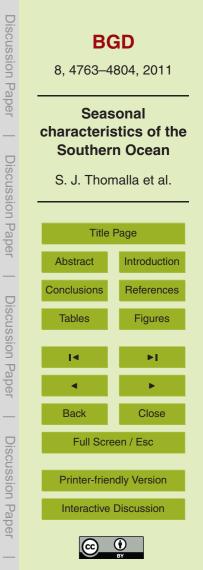
ocean (Minas and Minas, 1992) are considered to be the availability of light, iron (Fe) and silicic acid (Boyd, 2002; Moore and Abbott, 2002; Arrigo et al., 2008). However, the complexity of the interaction between these various regulating factors in the diverse spatial domain of the Southern Ocean is still poorly understood. Part of the lack

- of understanding of the links between upper ocean physics and biological processes controlling production in the Southern Ocean is due to the operational limitations in resolving them at the required in situ spatial and temporal scales. This has necessitated the use of remotely sensed and modelling techniques to further our understanding of this complex environment. Although a current weakness in remotely sensed data is its
- inability to directly link ocean colour with carbon export, it has the added advantage of being able to address the temporal and spatial scale gaps in our knowledge of a hitherto under sampled ocean (e.g., Arrigo et al., 1998; Moore and Abbott, 2000; Korb and Whitehouse, 2004; Park et al., 2010).

The spatial distribution of surface chlorophyll blooms in the Southern Ocean in sum<sup>15</sup> mer is thought to be consistent with an Fe limited regime, and this argument has been well documented (de Baar et al., 1995; Boyd et al., 2000; Gervais et al., 2002; Coale et al., 2004; Blain et al., 2007; Pollard et al., 2009). However, what has not been well understood are the physical control mechanisms responsible for supplying surface waters with Fe and modulating light that are ultimately responsible for controlling the phytoplankton seasonal cycle and spatial distribution. The intricate spatial and seasonal distribution of chlorophyll reflects the complex nature of the factors controlling production in the Southern Ocean, including sensitivity to mesoscale forcing as well as

remarkable regional and basin scale differences. Understanding regional characteristics of variability in bloom dynamics and the associated physical drivers is necessary to understand the sensitivity of the response of ocean productivity to climate change.

One of the strongest modes of variability within the carbon cycle is the seasonal cycle. It is the mode through which the physical mechanisms of climate forcing are coupled to ecosystem responses such as productivity, diversity and ultimately carbon export (Rodgers et al., 2008). The regional characteristics of the seasonal cycle in



the Southern Ocean are however not well understood. In this study we aim to examine the regional scale characteristics of the seasonal cycle of phytoplankton biomass and propose a hypothesis that relates the observed variability in biological response to variability in the physical drivers. This study addresses an important gap in our understanding of the papaitivity of the biological pump to alimpte aborgo in the Southern

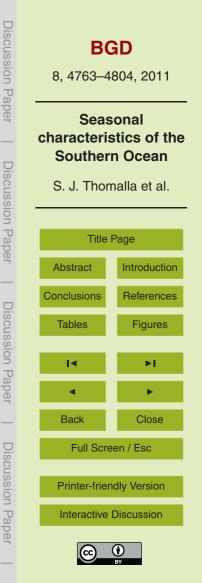
- <sup>5</sup> derstanding of the sensitivity of the biological pump to climate change in the Southern Ocean. Although this study is based exclusively on satellite data in a locale where ocean colour algorithms have well-known deficiencies, we anticipate that our results and hypotheses may guide future studies aimed at these gaps. The aim of this paper is not to investigate further the numerous controls of production in the Southern Ocean
- <sup>10</sup> but rather to use remote sensing data at appropriate temporal and spatial scales to characterise regional differences in the Southern Oceans seasonal cycle.

#### 2 Data and methods

#### 2.1 Satellite-derived surface chlorophyll concentrations

Ocean colour data are used to examine seasonal, intra-seasonal and inter-annual dynamics of phytoplankton blooms in the Southern Ocean. SeaWiFS (Sea-viewing Wide Field of view Sensor, McClain et al., 1998) data used in this study cover the period from January 1998 to December 2007. 8 day mean level 3 standard mapped images of chlorophyll (mg Chl m<sup>-3</sup>) on a global 9 km equidistant cylindrical grid from SeaWiFS were obtained from the Goddard Space Flight Centre (GSFC)
(http://oceancolor.gsfc.nasa.gov). The Southern Ocean domain south of 30° S was extracted and interpolated onto a regular 1/4 degree grid. Chlorophyll concentrations in the ocean tend to be log normally distributed (Campbell, 1995). When computing parametric statistical analyses on the original 8 day composite dataset, and in some cases for display purposes, the decimal logarithm of chlorophyll concentration has been used.

<sup>25</sup> Note that in most cases comparison with raw time-series do not lead to any substantial differences.



## 2.2 Mean Absolute Dynamic Topography (MADT)

We adapted the same technique described by Sokolov and Rintoul (2007b) to locate the main frontal branches of the Antarctic Circumpolar Current (ACC); the Subantarctic Front (SAF), the Polar Front (PF) and the Southern Antarctic Circumpolar Front

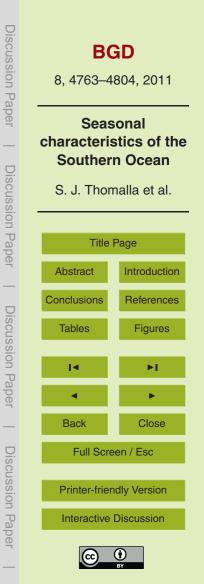
- <sup>5</sup> (SACCF), as well as the Subtropical Front (STF) for the entire circumpolar extent. We did this using the CLS/AVISO MADT dataset from the Data Unification and Altimeter Combination System (DUACS) archive. The global, 1/4 of a degree, weekly averages are obtained through an improved space/time objective analysis method combining TOPEX/Poseidon, ERS-1/2, JASON-1, GFO and ENVISAT data (Le Traon et al., 1998).
- The MADT is the sum of the sea level anomaly data and a mean dynamic topography (Rio05-Combined Mean Dynamic Topography, from Rio and Hernandez, 2004). The original weekly maps have been linearly interpolated in time to match the 8 day composite calendar on which the SeaWiFS datasets are provided.

#### 2.3 Bloom initiation date

We use "bloom" to refer to events of elevated chlorophyll concentration, without reference to a particular threshold. The initiation of the bloom (or the date of bloom onset) is understood here as the period of the year registering a relative increase in chlorophyll concentration, irrelevant of the actual value. The chlorophyll bloom is defined statistically, as in other studies (Henson and Thomas, 2007; Follows and Dutkiewicz, 2002; Siegel et al., 2002). Given the presence of missing values and the large degree of variability in some areas, extra care should be taken to allow the algorithm to accommodate for "aberrant" cases and avoid false detections of the bloom initiation date.

The mean bloom initiation dates were obtained for each pixel (on the 1/4 degree grid) as follows:

1. The time-series running from the 1st week of May 1998 to the last week of April 2007 is extracted.



- 2. The aberrant values (isolated spikes over the 99th percentile) are masked and discarded in the subsequent calculations.
- 3. The mean seasonal cycle is computed over the 9 yr analysed.
- 4. A 1-D Gaussian filter (with sigma = 1) is applied, effectively reducing the degree of intra-seasonal variability.
- 5. The median is calculated.

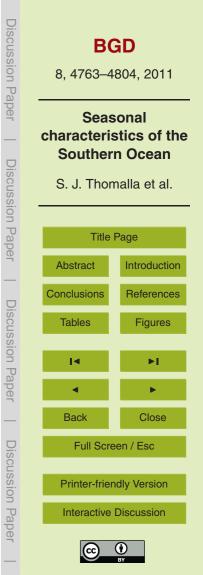
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- 6. The filtered mean seasonal cycle is repeated and wrapped around itself. The bloom initiation date is subsequently constrained to fall between the seasonal minima's.
- 7. The date of the bloom initiation is determined as the first week of the year where the chlorophyll concentration reaches +5% above the median and stays above this value for at least 2 consecutive weeks.

The bloom initiation dates for each year (from 1998/1999 to 2006/2007) have also been calculated. In this case steps 3 and 6 are not applied. It has been verified that
the bloom initiation date obtained from the average of the 9 yr is generally comparable to the one obtained from the mean seasonal cycle, albeit presenting a more noisy field given the sensitivity of the mean to extreme values and the short period considered. Inter-annual variability in the bloom initiation date was calculated as the standard deviation of the bloom initiation dates for each of the 9 yr. Given the short sample, one must be cautious in interpreting this figure, as the presence of some extremes can lead to high standard deviations.

## 2.4 Variance explained by the seasonal cycle

The part of the overall variance that is explained by the seasonal cycle was computed as the variance explained by the regression of log(chlorophyll) onto a repetition of the



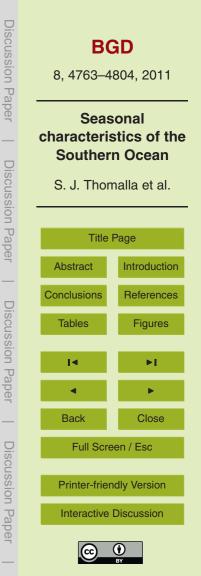
climatological mean seasonal cycle (calculated over the 9 available years) smoothed by a Gaussian filter of  $\sigma = 1$ . A value of 1 (100%) indicates that the time-series is a perfect repetition of the climatological mean seasonal cycle (i.e. there is no intraseasonal or inter-annual variability) while a value of 0 means that there is no yearly <sup>5</sup> cyclic component in the overall variability or no annually reproducible mean seasonal cycle.

# 2.5 Mixed Layer Depth (MLD)

MLD's have been taken from the monthly climatology developed by De Boyer-Montegut (2004). This global gridded product at 2° resolution is based on 4 490 571 individual temperature profiles obtained from the National Oceanographic Data Centre and the WOCE sections datasets, over the period 1941–2002. As the spatial and seasonal distribution of temperature profiles in the Southern Ocean is better than salinity profiles (De Boyer-Montegut et al., 2004) the MLD's were defined according to the optimal temperature criterion as a difference of 0.2 degrees with the temperature at 10 m.

## **2.6** Photosynthetically Active Radiation (PAR)

PAR is defined as the quantum energy flux from the sun in the spectral range 400 to 700 nm. PAR is expressed in Einstein m<sup>-2</sup> day<sup>-1</sup> (where 1 Einstein is equivalent to the energy of 1 mole of photons of monochromatic light). We used here the PAR generated by SeaWiFS and obtained through the GSFC. This dataset is provided on the same 9 km resolution grid as the chlorophyll estimates, for 8 day composites. The same period January 1998–December 2007 has been extracted and the values interpolated onto a regular 1/4 degree grid. SeaWiFS PAR values do not take into account any reflectance incurred at the air-sea interface and may therefore overestimate the amount of radiation entering the water column. Maximum reflectance during winter, when solar
radiation is low, ranges from 11 % at 40° N to 23 % at 70° N. During the summer months however, the zenith angle has little effect on the amount of surface reflectance which

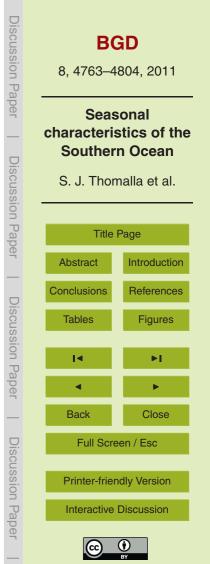


is generally between 4 and 7 % at all latitudes (40–70°) and under all atmospheric and wind conditions (Campbell and Aarup, 1989).

## 3 Results and discussion

# 3.1 Zonal characterization of seasonal biomass variability

- Remote sensing derived chlorophyll distributions in the Southern Ocean south of 30° S 5 reflect a zonal (annular) character of variability such as observed between frontal and pelagic zones as well as remarkable regional contrasts between the Atlantic and Pacific Oceans (Fig. 1a). These spatial zones also display important seasonal contrasts and inter-annual variability (Fig. 1a-c). While technically an estimate of pigment concentrations, we use SeaWiFS surface chlorophyll as a proxy for phytoplankton biomass 10 (Sullivan et al., 1993; Comiso et al., 1993; Moore et al., 1999; Moore and Abbott, 2000, 2002). Satellite surface chlorophyll concentrations do not however take into account deep chlorophyll maxima or changes in the carbon to chlorophyll ratio (C: Chl-a) with depth. Although typical deep chlorophyll maxima in the subtropics may result in underestimated satellite derived chlorophyll, typical chlorophyll profiles in the Southern 15 Ocean are uniform in the upper mixed layer and decreases exponentially at greater depths (Arrigo et al., 2008). The lack of depth integrated chlorophyll distribution is
- thus unlikely to significantly effect satellite derived chlorophyll concentrations in the Southern Ocean. Changes in C:Chl-a are associated with physiological responses
- to changing light, temperature and nutrient conditions and are lowest at high temperatures (25–30 °C), low irradiances (< 20 μmol photons m<sup>-2</sup> S<sup>-1</sup>) and nutrient replete conditions and increase at high irradiances, low temperature and nutrient limiting conditions (Taylor et al., 1997). Although seasonal variability in C: Chl-*a* ratios is significant at lower latitudes, the variations are relatively minor at higher latitudes when compared in the seasonal variability in C is the seasonal variability in C is the variation of the variati
- <sup>25</sup> with the much larger seasonal range in chlorophyll concentration (Taylor et al., 1997). Although changes in C: Chl-*a* ratios need to be considered when using satellite sur-



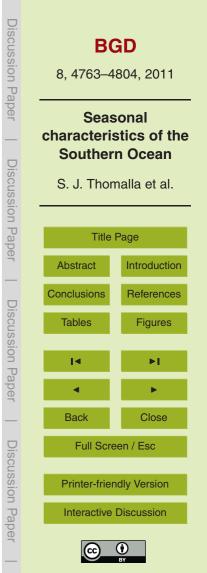
face chlorophyll measurements as a proxy for biomass, existing data suggest that these intra-cellular C: Chl-*a* ratios do not play a significant role in the seasonal cycle in the Southern Ocean (Taylor et al., 1997). In addition, standard ocean colour algorithms typically underestimate chlorophyll in the Southern Ocean by 2–3 times compared to in

situ measurements (Kahru and Mitchell, 2010). Underestimated chlorophyll concentrations are however unlikely to impact the characterisation of the patterns of the seasonal cycle which forms the primary focus of this paper.

In the subtropics, surface chlorophyll concentrations are low year round, except for a band of increased chlorophyll between  $30^{\circ}$  and  $40^{\circ}$ S in winter when nutrients are

- replenished (Fig. 1b). Surface chlorophyll concentrations in the Southern Ocean are low during the austral winter when light levels are low (Fig. 1b) (Mitchell et al., 1991; Veth et al., 1997; Lancelot et al., 2000; Smith et al., 2000; Boyd et al., 2001). Given the importance of Fe for photo-adaptation, Fe-light co-limitation is also likely (Sunda and Huntsman, 1997; Smith et al., 2000; Boyd et al., 2001). With the onset of a higher importance of a higher in environ adaptation of the sector of the sector of the sector.
- <sup>15</sup> irradiance-mixing regime in spring, chlorophyll concentrations increase rapidly forming blooms (Fig. 1a). Basin-scale differences are also evident with chlorophyll concentrations in the Pacific being noticeably lower than the Atlantic and Indian, in particular in the zonal band of 40–50° S which disrupts the almost symmetrical distribution around Antarctica. This band shows very little seasonal difference between summer and winter
   <sup>20</sup> chlorophyll concentrations (Fig. 1a and b).

Lowest summer chlorophyll concentrations are associated with the pelagic waters north of the sea ice zone (Fig. 1a) where production rates are limited by deep mixing of the upper mixed layer and trace metal limitation (Mitchell and Holm-Hansen, 1991; Boyd et al., 2000; Sokolov and Rintoul, 2007a). An exception to these low production <sup>25</sup> waters is found at oceanic frontal zones (Fig. 1a). Enhanced chlorophyll concentrations associated with the major fronts of the ACC have been well documented and attributed to a number of processes that include cross-frontal mixing of macronutrients, an improved light environment through enhanced stratification and increased Fe



concentrations through upwelling and interaction of the fronts with shallow topography

(e.g., Lutjerharms et al., 1985; Perissinotto et al., 1992; Pollard and Regier et al., 1992; Laubscher et al., 1993; De Baar et al., 1995; Moore et al., 1999; Moore and Abbott, 2002). Although traditionally, enhanced chlorophyll concentrations have been associated with mesoscale activities at the position of the fronts, Sokolov and Rintoul (2007b) more recently revealed that multiple frontal branches delimit regions with similar el-

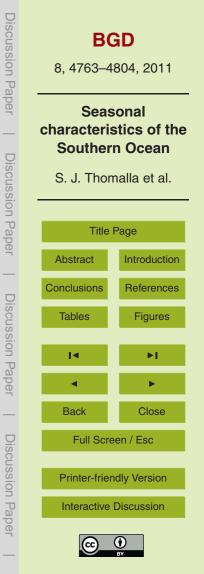
evated chlorophyll concentrations and seasonality rather than the fronts themselves being associated with enhanced productivity, at least where fronts are distant from topography.

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- Another exception to low production Southern Ocean waters is found over regions of shallow bathymetry; around and downstream of subantarctic islands, over mid-ocean ridges and large plateaus (Fig. 1a). In these regions of shallow bathymetry, current flow through relative vorticity (Hogg and Blundell, 2006; Moore et al., 1999) and/or bottom pressure torque (Sokolov and Rintoul, 2007a) is thought to increases the flux of Fe into surface waters (Korb and Whitehouse, 2004; Park et al., 2010; Venables and Meredith, 2010; Venables, and Mered-
- ith, 2010; Venables and Moore, 2010) accounting for the generally inverse correlation between depth and chlorophyll in the Southern Ocean (Comiso et al., 1993). Highest chlorophyll concentrations are generally associated with the marginal ice zone (MIZ) and the continental shelf (Fig. 1a) (Smith and Gordon, 1997; Moore and Abbott, 2000; Arrigo and Van Dijken, 2004), through enhanced irradiance from increased vertical
   stratification when ice melts (Smith and Nelson, 1986), through Fe input from melting
- ice (Sedwick and DiTullio, 1997; Gao et al., 2003; Grotti et al., 2005) and mixing of Fe rich sediments along the continental shelf (Schoemann et al., 1998; Johnson et al., 1999).

## 3.2 The seasonal cycle of chlorophyll in the Southern Ocean

<sup>25</sup> The characteristics of the seasonal cycle of phytoplankton biomass in the Southern Ocean are examined in terms of the timing of the bloom initiation, its amplitude, interannual and intra-seasonal variability and the importance of the climatological seasonal cycle in explaining the overall variance. The spatial distribution of the seasonal cycle

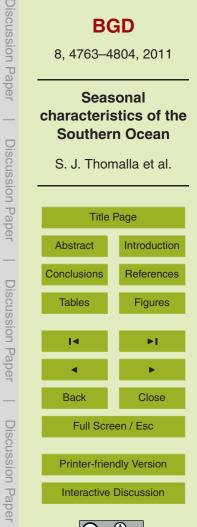


is subsequently defined to four zonal regions. The subtropical zone (STZ), the transition zone (TZ), the Antarctic circumpolar zone (ACZ) and the marginal ice zone (MIZ). The following section will describe the foundation for these zonal definitions based on seasonal characteristics rather than the distribution of chlorophyll concentrations, which has generally been the case in previous Southern Ocean studies. Defining the

Southern Ocean according to the characteristics of its seasonal cycle provides a more dynamic understanding of the characterisation of production based on underlying physical drivers rather than climatological biomass.

The chlorophyll bloom is considered a key phase in the seasonal cycle of phytoplankton biomass. The circumpolar mean bloom initiation dates are depicted in Fig. 2. Unlike in the Arctic, where significant trends towards earlier blooms were detected in 11 % of the area of the Arctic Ocean (Kahru et al., 2010), no homogenous regions showing distinct trends in either earlier or later bloom initiation dates were detected in the Southern Ocean, thus justifying the use of a 9 yr mean. In this study, the bloom initiation date has been associated with high biomass standing stocks and classified

- <sup>15</sup> initiation date has been associated with high biomass standing stocks and classified statistically according to an increase in chlorophyll concentration relative to the annual median (Henson and Thomas, 2007; Follows and Dutkiewicz, 2002; Siegel et al., 2002). This classification of a phytoplankton bloom is different to that of a recent study by Behrenfeld (2010), which classifies the bloom initiation according to the time when
- <sup>20</sup> phytoplankton population net growth rate becomes positive (see also Boss and Behrenfeld, 2010). Their results, based on chlorophyll concentrations in the North Atlantic, found that bloom initiation occurred in mid-winter when light levels are minimal and near-surface mixing is deepest. Behrenfeld's dilution-recoupling hypothesis challenges Sverdrup's Critical Depth Hypothesis (Sverdrup, 1953) by de-emphasizing the role of
- <sup>25</sup> light and instead suggests a much greater role for the balance between phytoplankton growth and losses through grazing. Although net population growth may increase in mid-winter, which according to Behrenfeld (2010) marks the start of the seasonal bloom, chlorophyll concentrations and specific growth rates are at their seasonal minimum. Our definition of the chlorophyll bloom on the other hand coincides with peaks

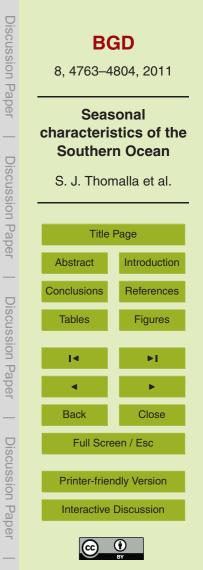




in phytoplankton productivity,  $CO_2$  uptake and carbon export, all of which are products of phytoplankton biomass and specific growth rates (Boss and Behrenfeld, 2010). The annual cycles of phytoplankton biomass in our results emphasize the role of bottomup controls (light and nutrients) on increases in phytoplankton specific growth rates for determining bloom initiation (Sverdrup, 1953). The lack of information on growth

- for determining bloom initiation (Sverdrup, 1953). The lack of information on growth rates does not allow us to quantify the role of grazing. In this study, we do not assume that the role of grazing is negligible, but rather that statistically significant seasonal increases in biomass can only occur when specific growth rates exceed loss terms and net population growth remains positive, despite potential increases in grazing pressure associated with increased encounter rates when the seasonal mixed layer shallows.
  - According to Fig. 2, bloom initiation in the STZ ( $\sim$  30–40° S), is characterized by an onset taking place in autumn (April–June) when macro-nutrients are replenished with the deepening of the seasonal mixed layer. This STZ is well defined in its latitudinal extent. North of  $\sim$  30° S, insufficient heat loss in winter limits the breakdown of stratification and deep winter overturning to below the nutricline preventing winter increases
- <sup>15</sup> cation and deep winter overturning to below the nutricline preventing winter increases in chlorophyll biomass. South of ~ 40° S, low winter PAR limits net primary productivity, preventing concomitant increases in winter chlorophyll concentrations, forcing phytoplankton populations towards spring bloom initiations (September–November). The transition between autumn and spring bloom initiations is remarkably abrupt (with the
- exception of the central and eastern Pacific where the transition is extended in latitude). The abrupt transition is indicative of a rapid shift from one seasonal regime to another, rather than a monotonic progression in the timing of the bloom initiation with increasing latitude. Further south is the MIZ where bloom initiation dates are in summer (December to February), reflecting the delay in seasonal PAR availability as well as the time it takes for phytoplankton blooms to fully respond to the power seasonal regime.
- <sup>25</sup> it takes for phytoplankton blooms to fully respond to the newly created ice-free waters associated with the region (Arrigo et al., 2008).

While the bloom onset in the STZ is spatially homogeneous, the Southern Ocean (south of 40° S) is characterized by large spatial variability with bloom initiation dates that can occur as early as May (e.g., Crozet, ~ 45° S, 50° E) or as late as January (south



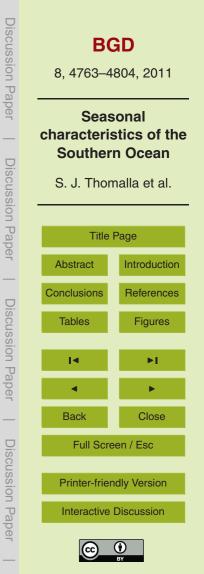
of Tasmania, ~ 42° S, 147° E). The transition between zonally comparable regions of early and late bloom onset can be relatively abrupt: for example the region of early (September, 100–140° E) and late (December, 130–170° E) bloom development found south of Australia (50–55° S) and the region of early onset evident north east of Crozet where the spring bloom starts particularly early in the year (May).

Regions with high inter-annual variability (std. dev. 10–16) in bloom initiation date (Fig. 3) correspond to transition regions between different seasonal regimes. In particular the transition between autumn and spring bloom initiation around ~ 40° S. The boundaries of this region of high variability in bloom initiation define what we term the

- <sup>10</sup> TZ. The remaining region of spring bloom initiation between the TZ and MIZ is what we term the ACZ. Other regions of high variability in bloom initiation similarly coincide with transition regions between different seasonal regimes, e.g. the region of late bloom initiation south of Australia (130–170° E) and the transition between spring and summer bloom initiation at the confluence of the ACZ and MIZ which is particularly
- extended in the Atlantic and the western Indian Ocean sector (20° W–60° E). Although the shift in mean bloom initiation date appears abrupt (Fig. 2), the TZ of high variability in bloom initiation dates is extended in latitude highlighting the discrepancy between climatological (Fig. 2) and annual time series (Fig. 3).

The percentage of the variance explained by the mean seasonal cycle (Fig. 4) de-<sup>20</sup> fines how well the mean climatological seasonal cycle (from 9 yr) represents the evolution of chlorophyll over each year. Immediately apparent from this figure are the sharp gradients between strongly contrasting regions where the seasonal cycle for each year is coherent with the 9 yr mean (i.e. high seasonality; > 70%) compared to regions where there is large variability from year to year in the timing and amplitude of the <sup>25</sup> bloom and only a low percentage of the variance is explained by the mean seasonal cycle (i.e. low seasonality; < 30%) (Fig. 4).

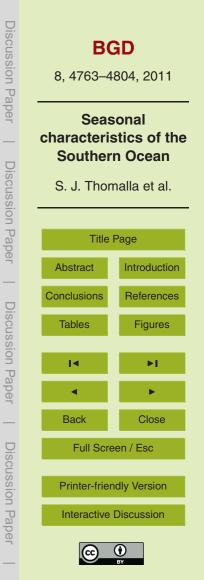
Despite the amplitude of the seasonal cycle in the STZ being weak (~ 0 to  $0.5 \,\text{mg}\,\text{m}^{-3}$ ) between summer minima and winter maxima, (see Fig. 1a and b) the overall variance of the chlorophyll signal is strongly phase locked to the mean seasonal



cycle (> 70 % of variance explained) (Fig. 4). An example of the time series of chlorophyll compared to the climatological mean seasonal in the STZ is shown in Fig. 5a. In this nutrient limited region, phytoplankton blooms occur in a predictable manner (std. dev. < 4, Fig. 3;  $R^2 = 91$ , Fig. 5a) when seasonal net heat loss and overturning in late autumn/early winter deepens the mixed layer to below the nutricline, relieving nutrient stress in surface waters and allowing phytoplankton production and biomass to increase. This finding agrees well with Dandonneau et al. (2004), who found low ratios of inter-annual to total variance from monthly SeaWiFS chlorophyll concentrations in the subtropical band between 1998 and 2001.

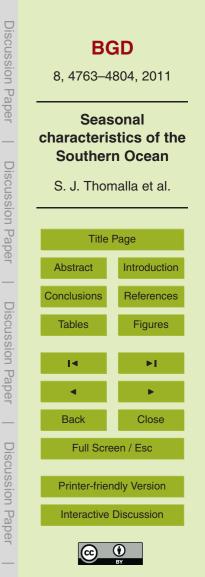
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- In the ACZ and MIZ, large regions of relatively high seasonality are also found, e.g. in the Pacific south of 50° S (~ 50 %), east of Kerguellen (~ 49° S, 70° E) (60–80 %) and a finer scale banded structure associated with the SAF zone and PF zone in the eastern Atlantic (50–70° S, 0–10° E) (Fig. 4). In regions with a high degree of seasonality (e.g. Fig. 5b and c;  $R^2$  = 66 and 64, respectively) one would expect that intra-seasonal
- forcing does not play a significant role in the phytoplankton seasonal cycle and that the annual time series would be almost entirely explained by the seasonal forcing of light, heat flux and seasonal MLD (as in the subtropics, e.g. Fig. 5a). This does not mean that Fe or light is not limiting but merely that it does not vary sufficiently on an intra-seasonal time scale to influence the inter-annual variability of the phytoplank-
- ton seasonal expression. In such instances one would expect there to be sufficient winter preconditioning of the water column with limiting nutrients (notably Fe) to allow a bloom initiation when the seasonal PAR threshold for increased primary production and biomass accumulation is met in spring. The amplitude of the seasonal bloom would depend on the supply of Fe via physical forcing mechanisms that include a)
- the seasonal re-supply of Fe through winter overturning, b) the depth of the summer mixed layer relative to the ferricline (deeper MLD's having higher Fe reserves), c) the amount of lateral advection of Fe into surface waters (e.g. downstream advection from continents and subantarctic islands) and d) the amount of Fe supplied by upwelling at fronts (e.g. Fig. 5b and c for low (~ 0.25 mg m<sup>-3</sup>) versus high (~ 1 mg m<sup>-3</sup>) summer



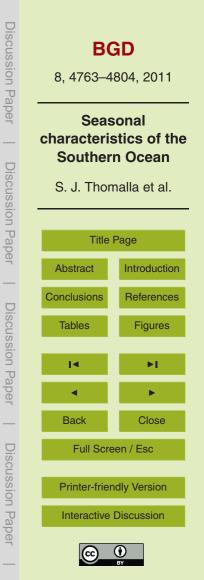
blooms, respectively). In these instances, the intra-seasonal re-supply of Fe to surface waters through wind mixing or mesoscale variability plays a less significant role in promoting phytoplankton growth, hence the low degree of intra-seasonal and inter-annual variability ( $R^2 = 66$  and  $R^2 = 66$ , Fig. 5b and c, respectively).

- <sup>5</sup> Regions of low seasonality (Fig. 4) generally correspond to regions of high variability in the phasing of the bloom initiation (Fig. 3). The zone of low variance explained by the mean seasonal cycle (< 20 %) marks the TZ (~ 40° S). This region is particularly extensive in the Pacific (~ 40–50° S), where chlorophyll concentrations are so low that the seasonal cycle is indistinguishable from intra-seasonal noise (e.g. Fig. 5d;
- $R^2 = 27$ ). The transition between spring and summer bloom initiation at the ACZ and MIZ confluence also marks a region of low seasonality. At a smaller scale, transitions between regions of contrasting bloom initiation dates within the ACZ are also characterized by weak seasonality e.g. the region of low variance explained by the seasonal cycle corresponding to the region of late bloom initiation south of Australia.
- Low seasonality related to the transition between regions of different bloom initiation is potentially related to the diversity of conditions encountered at the confluence between contrasting seasonal regimes. Transition regions of low seasonality are thought to be driven by a combination of multiple limiting factors and forcing mechanisms from both seasonal regimes. If there are too many factors simultaneously limiting phytoplankton production (e.g. nutrients and light), phytoplankton are not able to optimise to
- their environment, restricting the ability of the system to build up biomass. The expression of this will manifest as low annual chlorophyll concentrations with high inter-annual and intra-seasonal variability in bloom characteristics such as are found at the TZ in the Pacific (Figs. 4 and 5d).
- Low seasonality (Fig. 4) does not however necessarily coincide with low chlorophyll concentrations (Fig. 1a). The TZ of low seasonality in the Atlantic and Indian has relatively high summer chlorophyll concentrations (e.g. Fig. 5e;  $R^2 = 44$ ; chlorophyll max = ~ 1 mg m<sup>-3</sup>). Likewise, low seasonality regions surrounding the continental margins of Antarctica, America and Africa (Fig. 4) reflect high summer chlorophyll



concentrations (Fig. 1a). If our hypothesis for the low chlorophyll concentrations in the Pacific TZ holds true then there has to be a reduction in the number of factors simultaneously limiting production in regions of low seasonality but high chlorophyll. We propose that in these regions intra-seasonal physical forcing mechanisms are sequen-

- tially relieving light and/or nutrients at a frequency that is long enough to allow phytoplankton blooms to fully develop. At the latitudes of the continental margins of Africa and America for example, nutrients rather than light are considered the dominant limiting factor in summer. The upwelling of nutrients, particularly on eastern boundaries, through periodic wind events relieves nutrient stress on a sub-seasonal time scale.
- The period of sub-seasonal forcing must optimise growth in biomass at rates that exceed losses to account for the high integrated summer chlorophyll concentrations found here. The sub-seasonal wind events responsible for the upwelling and mixing of nutrients are likely to be responsible for the high intra-seasonal and inter-annual variability expressed as low seasonality. Along the continental margins of Antarctica on the other
- <sup>15</sup> hand, elevated Fe associated with the continental shelf and ice melt (Fitch and Moore, 2007) makes light the dominant limiting factor. Intra-seasonal forcing of the interaction between a deepening of the MLD through wind mixing and the re-establishment of stratification through fresh water buoyancy, at adequate time scales, similarly accounts for the high chlorophyll concentrations but low seasonality found in this region.
- The Supplement accompanying Figs. S1–4 elaborates on intra-seasonal and interannual variability (that result in either high or low seasonality) and complements the zonal characterisation of the Southern Ocean's seasonal cycle. Inter-annual and intraseasonal variability in the STZ is low and consistent with a high degree of seasonality (Fig. 4), whereas in the TZ, inter-annual and intra-seasonal variability is particularly high and seasonality consequently weak. A well defined but short lived bloom is found in the ACZ, and although there is low variability in bloom initiation dates, inter-annual variability in the amplitude of the bloom is high. In the MIZ, variability in bloom initiation dates is low but both inter-annual and intra-seasonal variability in the chlorophyll seasonal cycle is high, leading to low seasonality.



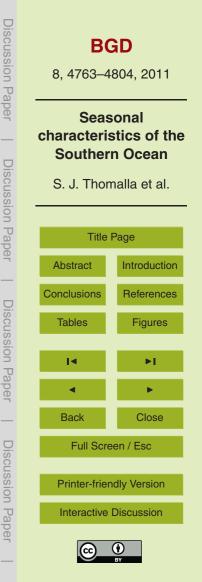
Although the fronts of the STF and ACC clearly influence the regional characterisation of the seasonal cycle of chlorophyll in the Southern Ocean (Figs. 1a, 2, 4), the physical forcing mechanisms responsible for enhanced chlorophyll (large scale flow versus small scale instabilities) are still unclear and need to be investigated further.

- Sokolov and Rintoul (2007) propose that mean chlorophyll distribution (and seasonal cycle) is best explained by upwelling where the ACC interacts with topography, followed by downstream advection where flow-topography interactions drive upwelling of nutrients, independent of mesoscale instabilities. Fronts however are known to provide a source of buoyancy through mesoscale and sub-mesoscale instabilities which influ-
- ence phytoplankton production through nutrient supply (upwelling) and light availability (stratification) (Swart and Speich, 2010; Kahru et al., 2007) as well as being zones of seasonal convergence. High resolution satellite imagery and model simulations now show that incorrect representation of sub-mesoscale frontogenesis can result in errors of up to 50 % in primary production estimates (Levy et al., 2001; Glover et al.,
- <sup>15</sup> 2008). Despite the significant contribution of fronts to characterising both the spatial and seasonal distribution of chlorophyll in the Southern Ocean, the contrasting view points in the literature highlights the indeterminate role of the fronts in controlling phytoplankton production. These differences seem to depend on the light/Fe co-limitation regime that they are superimposed upon which in effect determines the physical forcing <sup>20</sup> mechanism that is ultimately responsible for the observed variability in the chlorophyll
- seasonal cycle.

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## 3.3 Mixed layer depth, irradiance and the phytoplankton seasonal cycle

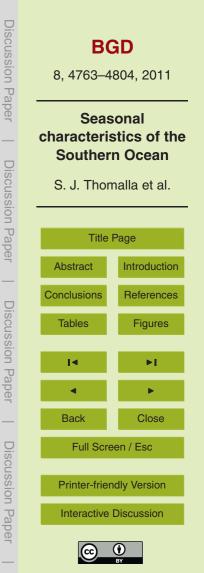
As expected, mean MLD's for winter (August) are deeper than in summer (February) and present large spatial variability (Fig. 6a and b). In the Atlantic ( $\sim 20^{\circ}$  E– $20^{\circ}$  W) and eastern Indian Ocean ( $\sim 60-180^{\circ}$  E), MLD's of > 200 and > 300 m respectively are found between  $\sim 40$  and 50° S in winter (Fig. 6a). Whereas in the Pacific in winter, ( $\sim 150-70^{\circ}$  W) MLD's of > 400 m are found further south between  $\sim 50$  and  $60^{\circ}$  S. In the 40– $50^{\circ}$  S latitudinal band, MLD's are relatively shallow ( $\sim 80-140$  m) in winter



compared to similar latitudes in the Atlantic and Indian (Fig. 6a). MLD's in summer present a more zonally coherent distribution, with MLD's shallower than  $\sim$  50 m observed north of 40° S and south of 60° S, with MLD's of 80–120 m in between (Fig. 6b).

The chlorophyll concentration at the sea surface generally responds to the seasonal

- <sup>5</sup> cycle of solar radiation which strongly impacts vertical stability through net heat flux, influencing vertical nutrient supply and the timing and intensity of phytoplankton blooms (Dandonneau et al., 2004). A first order estimate of the large-scale relationship between the seasonal cycles of MLD and chlorophyll (Fig. 7) can therefore be used to improve our understanding of the transition between different seasonal regimes.
- <sup>10</sup> Consistent with our understanding of the subtropics as a nutrient-limited regime, the correlation in the STZ is uniformly positive (> 0.8), with increased chlorophyll associated with of the deepest winter MLD's. High chlorophyll concentrations coinciding with deep MLD's in the subtropics are likely enhanced by photoadaptation of light limited cells in the deep winter mixed layer (Letelier et al., 1993).
- The transition between a positive and negative correlation (Fig. 7) corresponds remarkably well to the transition between winter and summer centred seasonal cycles (Fig. 2) reflecting the sharp climatological change in physical control mechanisms. Negative correlations are evident in the ACZ and MIZ, consistent with a light limited regime where increased chlorophyll concentrations coincide with shallow MLD's.
- A shoaling of the MLD provides the required light environment for phytoplankton production favouring increases in specific growth rates that exceeds export and losses through grazing and results in biomass accumulation (Sverdrup, 1953; Mitchell et al., 1991a; Comiso et al., 1993). However, in these high latitudes, low light (PAR) environments, the relationship is more convoluted than in the subtropics, with areas of low
- and in some instances slightly positive correlations (e.g., Crozet) interspersed among the general trend of a negative correlation. This "patchy" appearance is indicative of maximum seasonal chlorophyll concentrations that are not uniformly phased with the timing of minimum MLD's. Unlike in the subtropics, where a simple overturning threshold initiates the seasonal cycle, the physical forcing mechanisms of the mixed layer



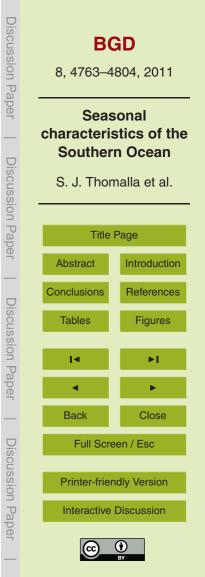
dynamics in the Southern Ocean are more varied and complex, as are the requirements to promote phytoplankton growth. Although the prevailing negative correlation between MLD and surface chlorophyll suggests that light is more important in limiting production in the subantarctic (relative to the subtropics), the Southern Ocean is

- <sup>5</sup> also nutrient limited, in particular with regards to Fe and Si (Boyd et al., 2002). The prolonged persistence of a shallow mixed layer can in some cases lead to nutrient limitation of primary production such that a periodic intra-seasonal deepening of the mixed layer (allowing Fe re-supply), at time scales of the same order as the phytoplankton growth rate, is necessary to maintain production rates at high levels throughout the summary production at all 2005). In such instances, white plantage and the production rates at high levels throughout the summary production and the production rates at high levels throughout the summary production and the production rates at high levels throughout the summary productio
- <sup>10</sup> summer season (Pasquero et al., 2005). In such instances, phytoplankton population growth rates are able to exceed their losses resulting in a well defined seasonal phytoplankton bloom (the integrated effect of localised sub-seasonal blooms) that does not coincide with the shallowest MLD's.
- Changes in the mixed layer result from the interaction of turbulent mixing, through <sup>15</sup> wind stress, and buoyancy forcing, through air/sea heat fluxes, fresh water fluxes, entrainment and advection (geostrophic and Ekman). These two drivers (turbulent mixing and buoyancy forcing) compete to either strengthen stratification or destroy it. We propose that the complex nature of the controls on stratification and production in the Southern Ocean plays an instrumental role in the expression of seasonal, sub-<sup>20</sup> seasonal and regional variability in chlorophyll concentrations and the seasonal cycle.

#### 3.4 Basin scale meridional controls of the seasonal cycle

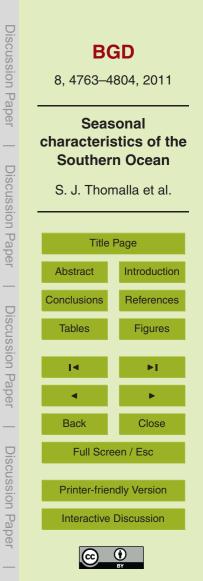
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The zonally asymmetric nature of the seasonal cycle of chlorophyll in the Southern Ocean provides the impetus for basin scale comparisons in addition to zonal characterisation. This asymmetry was similarly noted in the response of the MLD to the SAM by Salle et al. (2009). From a heat budget of the mixed layer they conclude that meridional winds associated with the departure of the SAM from zonal symmetry causes anomalies in heat flux and MLD which has consequences for biological productivity. In



to more accurately characterise regional differences in the phasing of the chlorophyll bloom with respect to MLD's. The seasonal progression of chlorophyll anomalies with respect to the annual mean were plotted with mean MLD's for three transects, one in each of the ocean basins (Fig. 8a–c).

- For each of the three transects a similar latitudinal progression in seasonal characteristics of the chlorophyll bloom is evident (Fig. 8a–c). In the STZ to the north maximum chlorophyll concentrations are in winter spanning August to October in the Atlantic (Fig. 8a), July to September in the Indian (Fig. 8b) and June to September in the Pacific (Fig. 8c), coincident with maximum winter MLD's (50–30 m). Minimum chlorophyll concentrations occur in late summer (January to March) when MLD's are shallow
- phyll concentrations occur in late summer (January to March) when MLD's are shallow (20–50 m). There is a southward extension of the STZ in the Pacific (~ 30–40° S) relative the Atlantic and Indian (~ 30–35° S). Further south, in the ACZ, the phasing of the chlorophyll bloom switches to one in which maximum concentrations are centred around the summer months (November to January), following the shoaling of the winter
- <sup>15</sup> mixed layer (< 70 m). In the MIZ bloom initiation occurs one month later (Fig. 2) with maximum concentrations extend into autumn (March) coinciding with shallowest MLD's (30–40 m). The presence of the Weddell Gyre in the Atlantic transect (Fig. 8a) broadens the latitudinal extent of the MIZ (~ 57–70° S) compared to the Indian (~ 60–65° S) and Pacific (~ 70–75° S) (Fig. 8b and c).</p>
- Besides the comparable characteristics in latitudinal progression of the seasonal cycle, inter-ocean basin differences are also evident. The Atlantic and Indian transects (Fig. 8a and b) are characterised by a relatively rapid TZ (~35–40° S), the northern boundary of the summer bloom region of the ACZ occurring at ~37° S and ~43° S, respectively. In the Pacific transect (Fig. 8c); the TZ between winter and summer centered seasonal regimes is extended in latitude to ~10 degrees (40–50° S), with distinct summer bloom characteristics only evident south of 49° S. The latitudinal transition to
- a summer bloom regime coincides with the northernmost expression of the deepest winter MLD's i.e. 130 m in the Atlantic (42°S), 300 m in the Indian (43°S) and 250 m in the Pacific (49°S). Although the mechanism responsible for the absence of a deep



winter mixed layer in the 40–50° S latitudinal band in the Pacific (90–100 m) is unclear (see also Fig. 6a), it is likely that an additional sustained input of buoyancy (e.g. additional heat source from southward penetrations of subtropical water) prevents the deepening of the mixed layer in winter to depths comparable to those of the Atlantic and Indian (130 and 250 m, respectively).

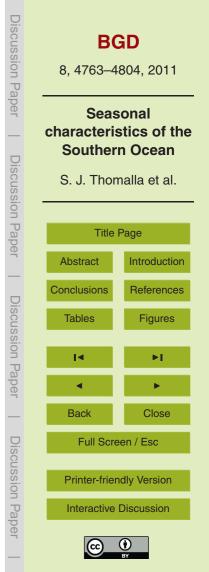
Intuitively, the Pacific, when compared to the Atlantic and Indian, has a potentially lower Fe supply to surface waters through the lack of continental or subantarctic island and sub-ocean land mass and lower mesoscale eddy activity from a lack of bathymetry to interact with the mean flow. Increased eddy kinetic energy associated with such fea-

- <sup>10</sup> tures, enhances the sub-surface flux of Fe into surface waters, potentially enhancing phytoplankton growth (Hense et al., 2000; Moore et al., 1999; Sokolov and Rintoul, 2007a; Park et al., 2010). If the main source of Fe to the surface waters (73–99%) of the Southern Ocean is through upwelling (Johnson et al., 1997; Lefevre and Watson, 1999; Archer and Johnson, 2000), one can deduce that in the Pacific, where Fe sup-
- <sup>15</sup> ply to surface waters through deep water entrainment is limited, shallow winter mixed layers are unlikely to be deep enough to entrain sufficient Fe into the surface waters for stimulating and maintaining high production rates through the summer. Hence, the extended (35–49° S) low chlorophyll TZ in the Pacific, with high variability and low seasonality likely results from Fe limitation.

#### 20 4 Synthesis

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Characterising the Southern Ocean according to the variability of the seasonal cycle provides a more dynamic understanding of the spatial heterogeneity of production based on underlying physical drivers rather than mean climatological biomass. Based on our analysis of the timing, variability and seasonality of seasonal chlorophyll distributions, we defined the spatial distribution of the seasonal cycle into four zonal regions: (1) the STZ (~ 30–35 or 40° S), characterised by an autumn bloom initiation (Fig. 2), low inter-annual and intra-seasonal variability and high seasonality (Fig. 4); (2) The TZ

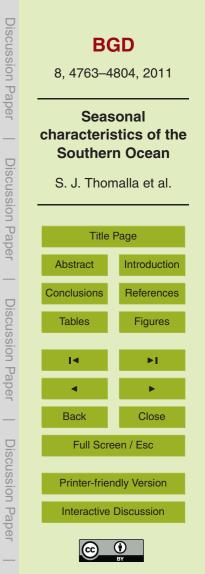


of high variability in bloom initiation dates (centred around ~  $40^{\circ}$  S) (Fig. 3), where interannual and intra-seasonal variability is also high and seasonality consequently weak (Fig. 4); (3) The ACZ (south of the TZ) where bloom initiations are generally in spring (Fig. 2), inter-annual variability in bloom initiation dates is low (Fig. 3) and large areas of both high and low seasonality are found (Fig. 4), and (4) The (MIZ) where bloom initiations are in summer (Fig. 2), variability in bloom initiation dates is low (Fig. 3) but inter-annual and intra-seasonal variability in the chlorophyll seasonal cycle is high, leading to low seasonality (Fig. 4).

This zonal classification effectively characterises the variability of the seasonal cycle in the Southern Ocean. However, when the response of the biology to the underlying physics of the different seasonal regimes was taken into consideration (i.e. low versus high seasonal chlorophyll maxima), an additional classification system was required. In order to summarise the varying responses of the phytoplankton community to the different seasonal regimes, we created a schematic that divides the Southern Ocean

- into a montage of four regions in addition to the four seasonal zones (Fig. 9). This schematic is created from a composite of the mean summer chlorophyll concentration from November to January (1998–2007) and the variance explained by the seasonal cycle (Fig. 4). The four regions result from a combination of high or low chlorophyll concentration and high or low seasonality. No allowance has been made for interme-
- diate chlorophyll concentrations or seasonality, and as a consequence the boundaries between regions are not definitive. This needs to be taken into consideration when interpreting this figure which is intended as a support to a conceptual framework on the nature of the controls of variability in the phytoplankton seasonal cycle.

Region A is representative of a nutrient limited regime with low chlorophyll concentrations but high seasonality. In Region A, in the STZ (30–40° S), nutrients (in particular NO<sub>3</sub>) are limited throughout the summer months and it is only when the mixed layer deepens in autumn that nutrient stress in surface waters is relieved, allowing phytoplankton production and biomass to increase. Similarly, Region A in the low chlorophyll pelagic waters of the Southern Ocean (north of the MIZ, away from fronts and shallow



topographic features) is also nutrient limited. Winter overturning re-supplies limiting surface nutrients (in particular Fe) to support an increase in phytoplankton growth, but only in spring when light levels are sufficient. In these regions, the available nutrients are rapidly used up by the phytoplankton community and the bloom subsequently de-

<sup>5</sup> clines when community losses outweigh growth rates. Despite the amplitude of the seasonal cycle in these regions being weak, the overall variability of the chlorophyll signal is strongly phase-locked to the annual seasonal cycle. In these regions, no sub-seasonal forcing of the mixed layer to below the nutricline is replenishing surface nutrients on intra-seasonal time scales, which accounts for the low intra-seasonal and inter-annual variability and a predictable seasonal cycle (high seasonality).

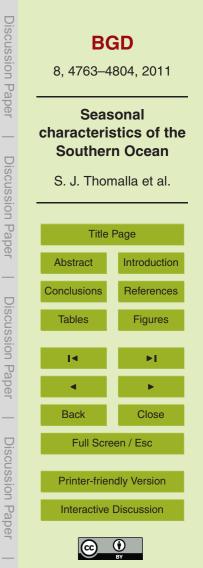
The second region with low chlorophyll concentrations (Region B) is also characterised by low Fe concentrations but this region displays low seasonality. This region is particularly extensive in the Pacific where increased positive buoyancy forcing potentially prevents the deepening of the winter mixed layer to below the ferricline, thus failing to re-set the seasonal Fe supply. If the winter re-supply of Fe to surface waters

failing to re-set the seasonal Fe supply. If the winter re-supply of Fe to surface waters is insufficient and/or the depth of the ferricline is below the deepest summer MLD's then phytoplankton lack the potential to significantly increase their biomass during the summer season through Fe limitation. These regions are characterised by chlorophyll concentrations that are so low and intra-seasonal variability so high that the seasonal cycle is indistinguishable from intra-seasonal noise.

The last two regions refer to areas of high chlorophyll but with either a low (Region C) or high (Region D) seasonality. Regions of high chlorophyll in the Southern Ocean result from integrated seasonal primary production that exceeds community losses and according to the literature results from the relief of Fe stress at ocean fronts, over

shallow bathymetry, along continental margins and in the MIZ (e.g., Boyd et al., 2002). The different seasonal expressions of high and low seasonality, however implies distinct physical supply mechanisms of Fe and light to the surface waters.

In the high chlorophyll, high seasonality regions (Region C), intra-seasonal forcing of the nutrient and light regime is not the dominant mechanism responsible for enhanced



summer chlorophyll. In these regions, the annual time series is almost entirely explained by the seasonal forcing of light, heat flux and seasonal MLD (as in the subtropics). This is not to say that Fe or light is not a limiting factor but merely that it does not vary sufficiently on an intra-seasonal time scale to influence the inter-annual variability

of the phytoplankton seasonal expression. In such instances sufficient winter preconditioning of the water column with limiting nutrients is required in order to allow a spring bloom initiation. The integrated seasonal amplitude of the bloom would depend on the amount of Fe made available through winter overturning, the depth of the summer mixed layer relative to the nutricline, lateral advection of Fe into surface waters and upwelling at fronts.

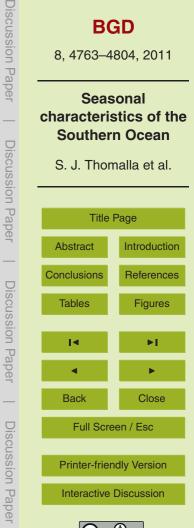
In regions of high chlorophyll but low seasonality (Region D), the seasonal characteristics of high inter-annual and intra-seasonal variability are controlled by sub-seasonal forcing of the nutrient and light supply. In these regions, we hypothesize that high integrated summer chlorophyll concentrations are a direct consequence of high intra-

- <sup>15</sup> seasonal physical forcing of the MLD at appropriate time scales (Pasquero et al., 2005). In these regions, shallow MLD's likely lead to short term depletions of surface nutrients such that a periodic deepening of the ML (to below the nutricline) is necessary for phytoplankton population growth to occur. This high intra-seasonal variability culminates from a combination of high wind stress variability at appropriate time scales and upper water column stabilisation through positive buoyancy forcing via mesoscale dynamics
- and fresh water (ice melt) fluxes.

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Although the fronts of the STF and ACC clearly influence the regional characterisation of the seasonal cycle of chlorophyll in the Southern Ocean, the physical forcing mechanisms responsible for enhanced chlorophyll (large scale flow versus small scale instabilities) are still unclear and need to be investigated further.

The relevance of this more dynamic characterisation of Southern Ocean production is that it may contribute towards an improved understanding of the variability of regional sensitivity to climate forcing. The seasonal cycle is the most important mode of variability in climate change making it likely that long term trends will be reflected sensitively



through changes in the characteristics of the seasonal cycle. A better understanding of the regional sensitivities of the Southern Oceans biological pump to climate change will allow us to make more robust predictions of long term trends.

Supplementary material related to this article is available online at: http://www.biogeosciences-discuss.net/8/4763/2011/ bgd-8-4763-2011-supplement.pdf.

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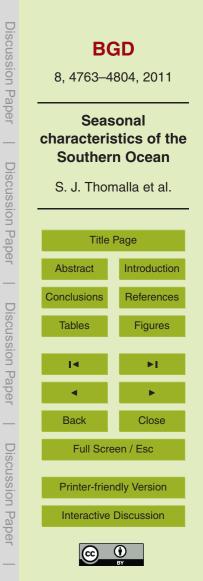
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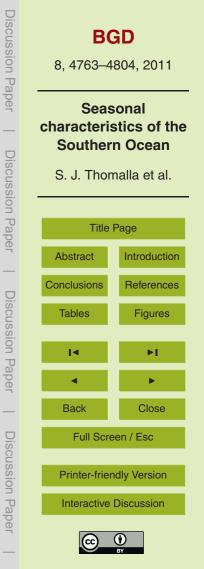
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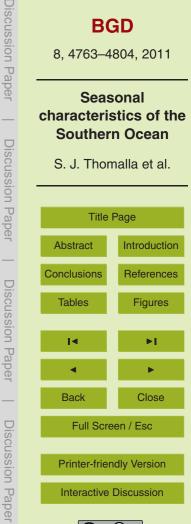
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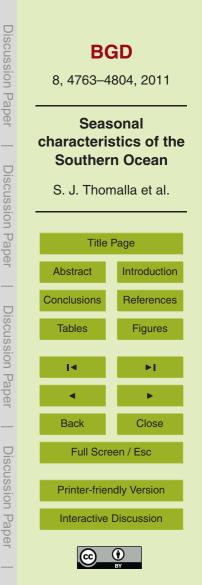
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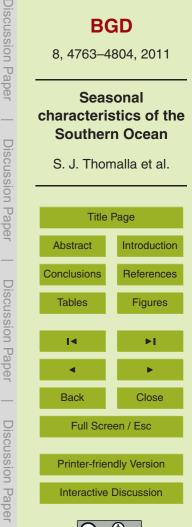
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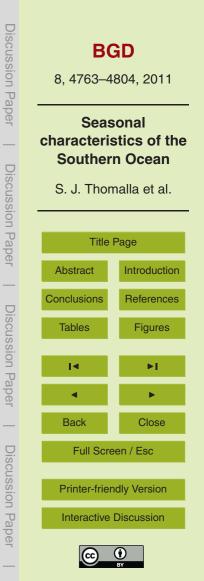
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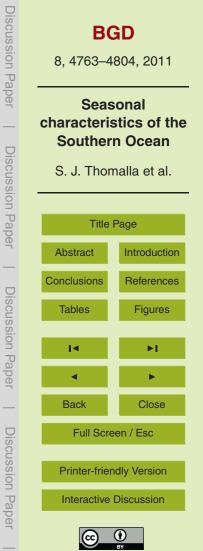
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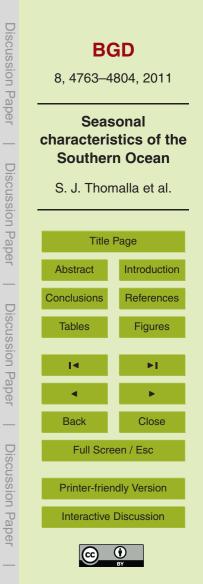
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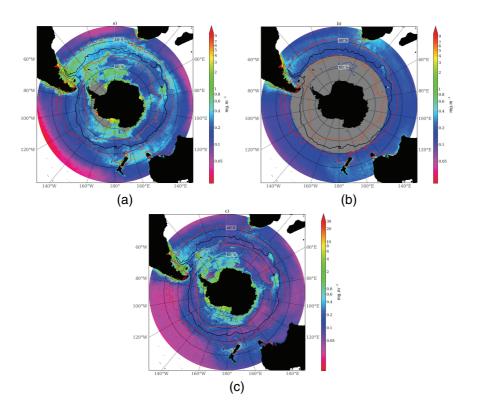
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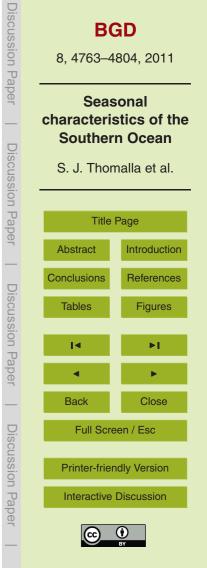
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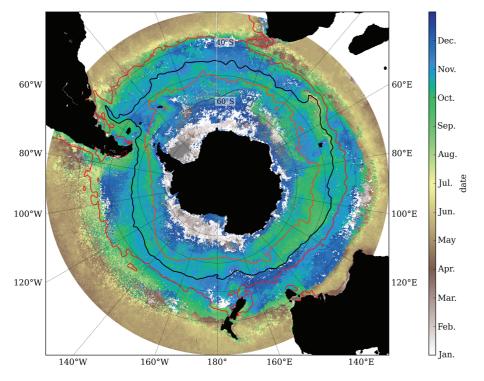
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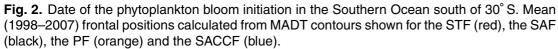


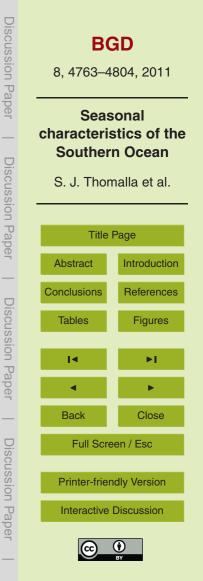


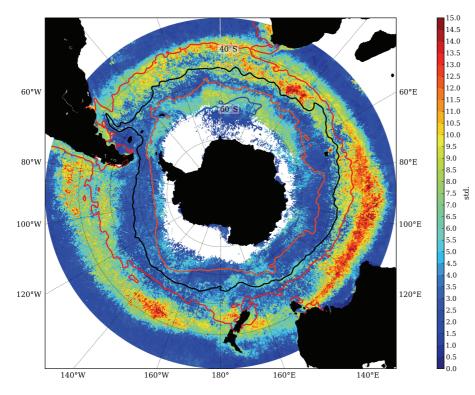


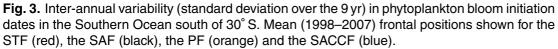
**Fig. 1.** Spatial distribution of mean chlorophyll concentrations for the Southern Ocean south of 30°S for **(a)** peak summer – January, **(b)** peak winter – July and **(c)** the standard deviation of weekly chlorophyll concentrations over the period January 1998–December 2007. Mean July and mean January (1998–2007) frontal positions calculated from MADT contours are shown for the STF (red), SAF (black), PF (orange) and SACCF (blue).

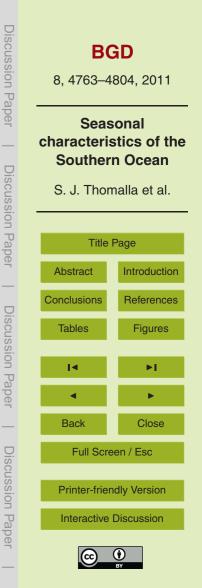


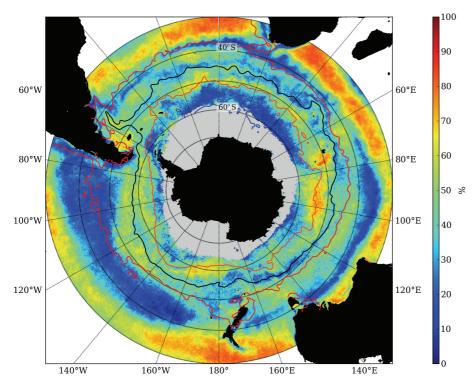


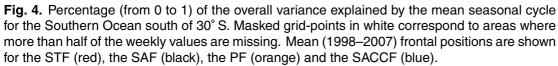


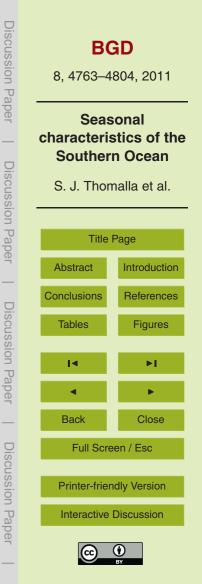


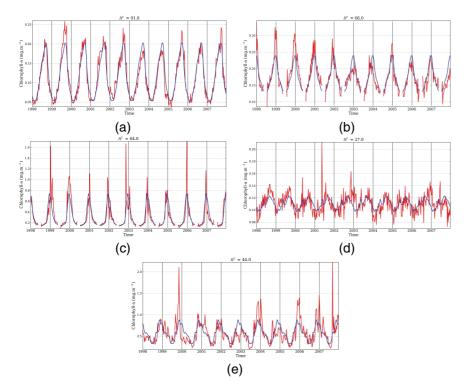




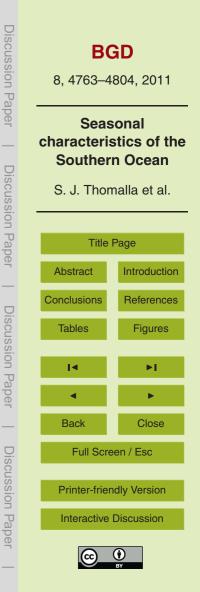


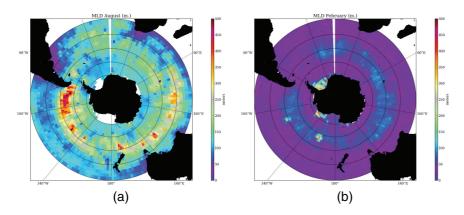




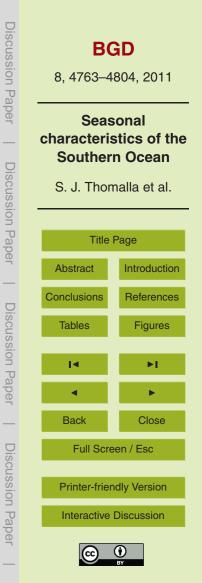


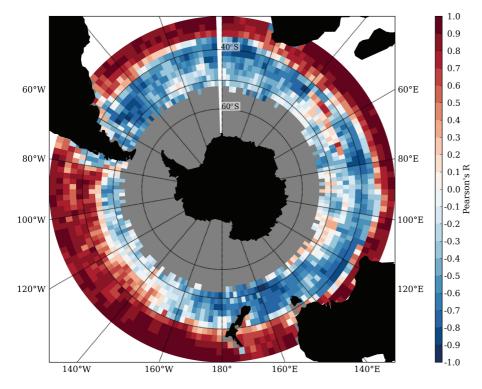
**Fig. 5.** Time series of chlorophyll concentrations from 1998 to 2007 (in red) compared to the climatological mean seasonal cycle (calculated over the 9 yr) (in blue) for  $5 \times 5$  degree blocks in (a) a high seasonality region in the low chlorophyll STZ (30–35° S, 0–5° W), (b) a high seasonality region in the low chlorophyll ACZ west of Kerguellen (50–55° S, 55–60° E), (c) a high seasonality region of high chlorophyll in the ACZ, downstream of Kerguellen (50–55° S, 70–75° E), (d) a low seasonality region in the low chlorophyll TZ in the Pacific (40–45° S, 95–100° W) and (e) a low seasonality region of high chlorophyll in the TZ off the east coast of South America (40–45° S, 50–55° W). The percentage of variance explained by the regression of the time-series (red) onto the mean seasonal cycle (blue) is shown on the figure ( $R^2$ ).

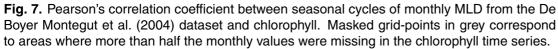


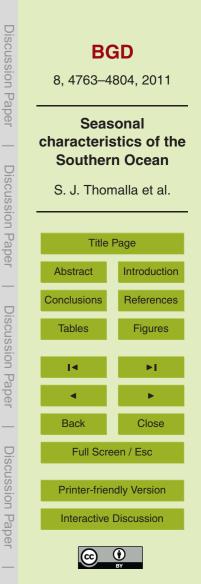


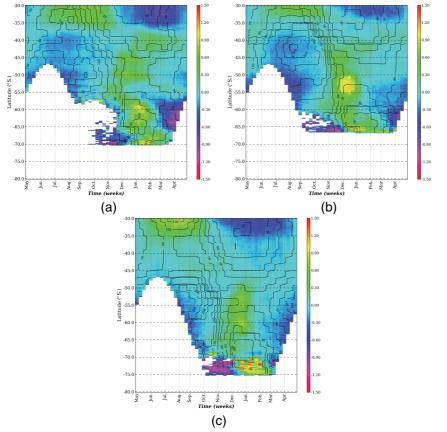
**Fig. 6.** Average MLD in **(a)** winter (August) and **(b)** summer (February) in the Southern Ocean from the De Boyer Montegut et al. (2004) dataset.



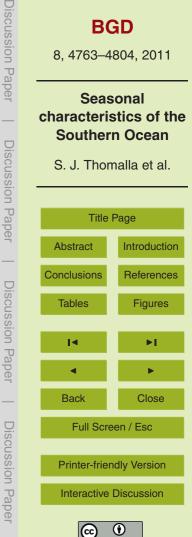


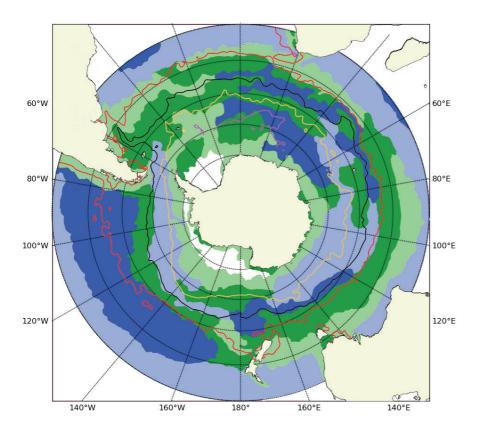


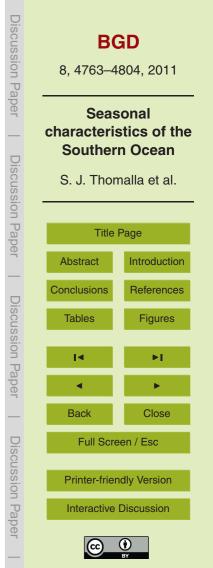




(c) **Fig. 8.** Mean seasonal cycles (May to April – *x*-axis) of chlorophyll (colour scale) and MLD (contours) as a function of latitude  $(30-80^{\circ} \text{ S} - y$ -axis) for a 10° longitudinal transect in (a) the Atlantic  $(0-10^{\circ} \text{ E})$ , (b) the Indian  $(85-95^{\circ} \text{ E})$  and (c) the Pacific  $(110-100^{\circ} \text{ W})$  Oceans. Chlorophyll concentrations are expressed as the anomalies in log(chlorophyll) with respect with the annual mean, MLD is in meters.







**Fig. 9.** A schematic summarising the response of phytoplankton biomass to the underlying physics of the different seasonal regimes. Regions in blue represent regions of low chlorophyll concentration with either high seasonality (light blue) or low seasonality (dark blue). Regions in green represent regions of high chlorophyll concentration with either high seasonality (dark green) or low seasonality (light green). Mean (1998–2007) frontal positions are shown for the STF (red), the SAF (black), the PF (yellow) and the SACCF (pink).