INVESTIGATING ON-BOARD FE FERTILIZATION EXPERIMENTS USING FAST REPETITION RATE FLUOROMETRY IN THE SOUTHERN OCEAN.

By
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Dissertation submitted in fulfilment of the requirement for the degree of Masters of Science in the Department of Zoology, University of Cape Town
Date: 11 February 2013

Supervisors: Associate Professor Mike Lucas and Dr Sandy Thomalla
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Abstract

In the face of increasing carbon dioxide (CO\textsubscript{2}) accumulation in the atmosphere and associated climate change, an understanding of important global carbon sinks such as the Southern Ocean becomes imperative. Phytoplankton, being the major driver of the biological pump, need to be understood at the level of photosynthetic efficiency, especially in relation to growth limiting factors such as iron (Fe). This study reports both observational and experimental phytoplankton growth rate observations from the summer months of 2009/10 and 2010/11, focusing on the Southern Ocean area between South Africa, Acta Bukta (Antarctica) and South Georgia. The study includes six ship based Fe enrichment incubation experiments to examine the interaction of Fe and light controls on photosynthesis. This study focuses on the phytoplankton photosynthetic responses using Fast Repetition Rate fluorometry (FRRf). Throughout the study area phytoplankton photosynthetic efficiency (F\textsubscript{v}/F\textsubscript{m}) increases in response to Fe alleviation, as is the case in the on-deck Fe-supplemented incubations. Chlorophyll-a (chl-a) increase due to Fe alleviation is not guaranteed, however it is dependent on initial community structure, the light environment and grazing rates. This research implies that the majority of the Southern Ocean is Fe limited, and that an increase in Fe would lead to an increase in photosynthetic efficiency, but not necessarily biomass or carbon export.
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<table>
<thead>
<tr>
<th>Abbreviation</th>
<th>Definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>$\rho_{NO_3}/Total\ \rho_{N}$</td>
<td>f-ratio calculation: (uptake Nitrate/total nitrogen uptake)</td>
</tr>
<tr>
<td>$\sigma_{PSII}$</td>
<td>The functional absorption cross section of photosystem II</td>
</tr>
<tr>
<td>°C</td>
<td>Degrees Celsius</td>
</tr>
<tr>
<td>AASW</td>
<td>Antarctic Surface Water</td>
</tr>
<tr>
<td>ACC</td>
<td>Antarctic Circumpolar Current</td>
</tr>
<tr>
<td>AOML</td>
<td>Atlantic Oceanographic and Meteorological Laboratory</td>
</tr>
<tr>
<td>APF</td>
<td>Antarctic Polar Front</td>
</tr>
<tr>
<td>APZ</td>
<td>Antarctic Polar Zone</td>
</tr>
<tr>
<td>ATP</td>
<td>Metabolic energy</td>
</tr>
<tr>
<td>CaCO$_3$</td>
<td>Calcium carbonate</td>
</tr>
<tr>
<td>CH$_2$O</td>
<td>Glucose</td>
</tr>
<tr>
<td>chl-a</td>
<td>Chlorophyll-a</td>
</tr>
<tr>
<td>CO$_2$</td>
<td>Carbon Dioxide</td>
</tr>
<tr>
<td>CROZEX</td>
<td>The Crozet Natural Iron Bloom and Export Experiment</td>
</tr>
<tr>
<td>CSIR</td>
<td>Counsel for Scientific and Industrial Research</td>
</tr>
<tr>
<td>CTD</td>
<td>Conductivity, Temperature and Depth</td>
</tr>
<tr>
<td>DIC</td>
<td>Dissolved Inorganic Carbon</td>
</tr>
<tr>
<td>DOC</td>
<td>Dissolved Organic Carbon</td>
</tr>
<tr>
<td>$F_v/F_m$</td>
<td>Photochemical efficiency</td>
</tr>
<tr>
<td>Fd</td>
<td>Ferredoxin</td>
</tr>
<tr>
<td>Fe</td>
<td>Iron</td>
</tr>
<tr>
<td>FeSO$_4$</td>
<td>Iron Sulphate</td>
</tr>
<tr>
<td>FIRE</td>
<td>Fluorescence, Induction and Relaxation</td>
</tr>
<tr>
<td>Acronym</td>
<td>Description</td>
</tr>
<tr>
<td>---------</td>
<td>-------------</td>
</tr>
<tr>
<td>FRRf</td>
<td>Fast Repetition Rate fluorometry</td>
</tr>
<tr>
<td>Fuc:Hex</td>
<td>Fucoxanthin:19’-hexanoyloxyfucoxanthin</td>
</tr>
<tr>
<td>GF/F</td>
<td>Glass Fibre Filter</td>
</tr>
<tr>
<td>GtC</td>
<td>Giga tons of carbon</td>
</tr>
<tr>
<td>H₂O</td>
<td>Water</td>
</tr>
<tr>
<td>HNLC</td>
<td>High-Nutrient-Low-Chlorophyll</td>
</tr>
<tr>
<td>HPLC</td>
<td>High Performance Liquid Chromatography</td>
</tr>
<tr>
<td>IPCC</td>
<td>Intergovernmental Panel on Climate Change</td>
</tr>
<tr>
<td>IR</td>
<td>Infra-Red</td>
</tr>
<tr>
<td>Km</td>
<td>Kilometer</td>
</tr>
<tr>
<td>Kg m⁻³</td>
<td>Kilogram per cubic metre</td>
</tr>
<tr>
<td>NADPH</td>
<td>Nicotinamide Adenine Dinucleotide Phosphate</td>
</tr>
<tr>
<td>NH₄</td>
<td>Ammonium</td>
</tr>
<tr>
<td>Nm</td>
<td>Nautical miles</td>
</tr>
<tr>
<td>NO₃</td>
<td>Nitrate</td>
</tr>
<tr>
<td>NO₂</td>
<td>Nitrite</td>
</tr>
<tr>
<td>NOAA</td>
<td>National Oceanic and Atmospheric Administration</td>
</tr>
<tr>
<td>NOC</td>
<td>National Oceanography Centre</td>
</tr>
<tr>
<td>m</td>
<td>Metre</td>
</tr>
<tr>
<td>MATLAB</td>
<td>MATrix LABoratory</td>
</tr>
<tr>
<td>ml</td>
<td>Millilitres</td>
</tr>
<tr>
<td>MLD</td>
<td>Mixed Layer Depth</td>
</tr>
<tr>
<td>MTF</td>
<td>Length of the multi-turnover flash phase in ms</td>
</tr>
<tr>
<td>MTRI</td>
<td>The initial interval between relaxation pulses</td>
</tr>
<tr>
<td>MTRP</td>
<td>The number of pulses in the relaxation sequence</td>
</tr>
</tbody>
</table>
O$_2$ Oxygen
PAR Photosynthetically Active Radiation
Pc Plastocyanin
P$_g$C y$^{-1}$ Petagrams of carbon per year
pH Measure of the acidity of a solution
PO$_4$ Phosphate
POC Particulate Organic Carbon
PON Particulate Organic Nitrogen
ppm Parts per million
Pq Plastoquinone
PS Photosystem
PSU Practical salinity unit
UCTD Underway Conductivity, Temperature and Depth
μg.L$^{-1}$ Micrograms per litre
μmolL$^{-1}$ Micromoles per litre
VNO$_3$ d$^{-1}$ Specific uptake rate of NO$_3$ per day
s Seconds
SACCF Southern Antarctic Circumpolar Current Front
SAF Subantarctic Front
SANAE South African National Antarctica Expedition
SANAP South African National Antarctic Program
SBdy Southern Boundary of the Antarctic Circumpolar Current
Si Silicon
SiO$_4$ Silicate
SOIREE Southern Ocean Iron Release Experiment
<table>
<thead>
<tr>
<th>Acronym</th>
<th>Definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>LSTF</td>
<td>Length of a single turnover flash in μsec</td>
</tr>
<tr>
<td>STF</td>
<td>Subtropical Front</td>
</tr>
<tr>
<td>STRI</td>
<td>Initial interval between relaxation pulses in μsec</td>
</tr>
<tr>
<td>STRP</td>
<td>Number of pulses in the relaxation sequence</td>
</tr>
<tr>
<td>XBT</td>
<td>Expendable Bathythermograph</td>
</tr>
<tr>
<td>UCDW</td>
<td>Upper Circumpolar Deep Water</td>
</tr>
<tr>
<td>WOCE</td>
<td>World Ocean Circulation Experiment</td>
</tr>
<tr>
<td>WW</td>
<td>Winter Water</td>
</tr>
</tbody>
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CHAPTER 1: INTRODUCTION

1.1) Climate change

The last century has seen a significant warming across the globe, with the global average temperature increasing by 0.76°C between 1850 and 1899 as well as between 2001 and 2005 (IPCC, 2007a). Although there is a natural background climatic variation, a substantial climate change signal has emerged from this natural variability (Field et al., 2002; Sabine et al., 2004b), with the warming trend increasing significantly over the second half of this century (IPCC, 2007a). Further indications of climate change have been observed in the oceans. The latent heat differences existing between water and gas, means that the oceans have heated 20 times more than the atmosphere since 1960 (Levitus et al., 2005). This has been observed specifically in warming of Southern Ocean Mode Waters and the Upper Circumpolar Deep Waters over this time period. Apart from warming, climate change can be seen in changes in global precipitation, which along with melting of the ice caps (Haeberli et al., 2005a; Haeberli et al., 2005b; Kaser et al., 2006) is linked to changes observed in ocean salinity. Between 1955 and 1998, freshening has occurred in subpolar latitudes and in the Pacific Ocean, while the Atlantic, Indian and shallower parts of the tropical and subtropical oceans have become saltier (IPCC, 2007d).

Geological and ice-core data provide our principle records of past climatic cycles, their changes and the long-term effects of climate change (Hughes et al., 2003). These records reveal long-term cooling of Earth over the last 60 million years, while more recently cycles in global climate have resulted in regular shifts between ice ages and eras of warming. The past shifts in our climate are attributed to either natural climate variability or climate change (where the changes occur outside of the natural cycles). The natural climate
variability results from cyclical variations in earth’s orbit around the sun, and in the angle of
tilt of the earth’s axis of rotation (MacCarthy and Rubidge, 2005), this effect on climate by
the earths’ movements are known as Milankovitch cycles which are clear throughout the
earth’s climate history. The less regular and modern variability since the 1750’s is regarded
as human-induced climate change.

Climate change can be (and has been in the past) caused by various factors such as
volcanic eruptions, variability in solar luminosity, changes in the position of land masses,
ocean currents due to plate tectonics, as well as Milankovitch forcing over longer periods
(23 000, 40 000 and 100 000 years). New to this list since the 1750’s is rapid human-forced
change over tens to hundreds of years due to anthropogenic aerosols and greenhouse gas
emissions that cause the corresponding greenhouse effect (Field et al., 2002). There is
mounting evidence (Robock, 1979; Wigley and Raper, 1990; Friis-Christensen and Lassen,
1991; Kelly and Wigley, 1992; Crowley and Kim, 1993; Rind and Overpeck, 1993; Cubasch
et al., 1997; Briffa et al., 1998; Mann et al., 1998; Crowley and Kim, 1999; Damon and
Peristykh, 1999; Free and Robock, 1999; Lean and Rind, 1999) that volcanic activity
significantly contributed to decadal-scale climate variability in the little ice age (Crowley,
2000). A prime example of the influence of volcanic eruptions on the world’s climate was
seen by the effects of the large dust cloud present in the stratosphere in the years following
Mount Pinatubo’s eruption (Bluth et al., 1992; Krueger et al., 1995), which cooled earth by
about 0.5°C. Furthermore, volcanism in the Southern Hemisphere north of 20° S influences
Northern Hemispheres temperatures (Crowley, 2000). Variation in solar luminosity caused by
sun-spots is believed to affect the climate due to variations in the radiation entering the
atmosphere (Hoyt, 1979).
However, neither of these natural processes can explain rapid global warming experienced since the 17th century (Foukall et al., 2006). The tectonic movement over this time is not sufficient to alter the Earth’s climate. Instead, anthropogenic influences provide the most plausible cause of recent global warming. The two principle causes are due to aerosols and greenhouse gases.

Anthropogenic aerosols pollute the atmosphere causing changes in the Earth’s albedo (which alters the fraction of solar radiation reflected both back to earth and space). Aerosols are tiny liquid droplets in the air which cause the scattering of light, and affect cloud formation and hence, rain patterns. This without a doubt is a global problem, as seen by the many Clean Air Acts around the world; United States of America (Clean Air Act of 1963, amended in 1970, 1977 and 1990), South Africa (Air Pollution Prevention Act in 1965 and the National Environmental Management: Air Quality Act 2004), Philippine (Clean Air Act of 1999) and Britain (Clean Air Act of 1956, updated in 1993) all have individual Clean Air Acts. These acts were drafted in response to human health issues with amendments being adjusted to include the health of the atmosphere itself. It should be recognised that while some aerosols cause cooling (e.g. sulphate aerosols), others cause warming (e.g. dark soots). These contradictory effects are due mainly to the type of aerosol, their height in the atmosphere, as well as their light scattering or absorption properties. Either way, anthropogenic aerosols are not the main cause of current global warming trends, though they are responsible for effecting precipitation patterns (Ramanathan et al., 2001; Menon et al., 2002) and possibly enhancing bush encroachment (Wigley et al., 2010).

More significant are the greenhouse gases. An elevated greenhouse effect is caused by elevated greenhouse gas concentrations of (e.g. carbon dioxide [CO₂], methane, nitrous oxide
and fluorocarbons) in the atmosphere, which in turn alters the balance between long-wave infra-red (IR) radiation lost to space, or re-radiated back to Earth (IPCC, 2007a). Where the radiation balance results in more reflected IR, this leads to an accumulation of excess heat within the atmosphere. The greenhouse effect is primarily anthropogenic, due to fossil fuels combustion, land use change (primarily in the tropics), as well as various industrial processes (Field et al., 2002) such as cement production (Hendriks et al., 2003). Although the causes of climate change over the last century are still argued, within scientific circles it is almost unanimously agreed that such changes stem from anthropogenic activities and that “the link between greenhouse gases and observed climate change are now incontrovertible” (Hoegh-Guldberg, 1999; IPCC, 2001).

Present atmospheric CO$_2$ concentration is rising between 10 and 100 times faster than at any other time in the past 420 000 years (Falkowski et al., 2000). This is due to the rapid industrialization that occurred after the 1750’s, when burning fossil fuels provided the energy for great industrial advancements, but with corresponding growth in CO$_2$ emissions. This development has continued into the 21st century with “56.6% of all anthropogenic greenhouse gas emissions coming from 85% of the global economies that are primarily energy driven through the consumption of fossil fuels” (IPCC, 2011). At the end of 2010, atmospheric CO$_2$ exceeded pre-industrial levels by 39%. In the absence of any further climate change policies, climate models are predicting a global average surface temperature rise of 1.5 – 5°C (Field et al., 2002; IPCC, 2007b) over the next century (relative to 1999).

“Anthropogenic caused climate change is already having a significant impact on multiple systems globally” (Rosenzweig et al., 2008). These include melting Arctic ice, shrinking mountain glaciers (Haeberli et al., 2005a; Haeberli et al., 2005b; Kaser et al.,
and more extreme weather patterns, with wet areas receiving more precipitation, and dry areas less, leading to the intensification and spread of deserts (Hillel and Rosenzweig, 2002; IPCC, 2007c).

Because modern societies are non-nomadic, climate change has important social and economic consequences. To reduce the impact of such threats, the Cancun agreement (2010) called for limiting the maximum global average temperature rise to 2°C above pre-industrial values. To achieve this, CO$_2$ would need to stabilize in the atmosphere between 445 - 490 ppm (at the end of 2010, atmospheric CO$_2$ was at 390 ppm) (IPCC, 2011). Current global agreements on CO$_2$ emissions are insufficient to limit us to such a goal (Cuypers et al., 2011).

1.2 ) The carbon cycle

There is a continual carbon cycle that has existed throughout Earth’s history which has varied over time, in response to climate change (Barnola et al., 1987; Petit et al., 1999; IPCC, 2007a; Sundquist and Visser Ackerman, 2014). Industrialization has seen disruption in the natural carbon cycle (IPCC, 2013), with anthropogenic released carbon in to the atmosphere estimated at a net flux of 8.4 PgC y$^{-1}$ (Sabine et al., 2004b) The oceans are naturally both a major source (~70.6 PgC y$^{-1}$) and sink (~70 PgC y$^{-1}$; Sabine et al., 2004b), for atmospheric CO$_2$ (Siegenthaler, 1986). Human perturbations have altered this natural cycle by ~20 PgC y$^{-1}$ and ~21.9 PgC y$^{-1}$ for source and sink respectively, resulting in a net overall sink of ~1.3 PgC y$^{-1}$ (Sabine et al., 2004b). The net uptake of carbon by the ocean since global industrialization had by 1994 increased the oceanic carbon inventory by 118 ± 19 GtC (IPCC, 2007d). Though the oceans continue to take up CO$_2$, the rate at which they succeed in doing so has decreased since the 1980’s (IPCC, 2007d).
Trends in atmospheric CO\(_2\), such as the slowing of emitted CO\(_2\) (from 0.5% per year to 0.3% per year (Hansen et al., 1998) in the late 1990’s (despite a period of worldwide increased economic and industrial growth (Zagha and Nankani, 2005), puzzle the scientific community (Field et al., 2002). Associated changes in the terrestrial carbon sinks such as bush encroachment in grasslands (Bond and Midgley, 2000; Morgan et al., 2007; Kgope et al., 2009; Wigley et al., 2010) occurred at this time (Higgins et al., 1999; Roques et al., 2001; Moleele et al., 2002). The economic and social consequences of such changes (Bond and Midgley, 2000) lead to the necessity of better understanding the carbon sinks, such as within the oceans. It is also key to understand processes that influence the carbon cycle, such as photosynthetic responses (Coale et al., 1996), and how these respond to changing environmental conditions whether due to geo-engineering consequences or to natural changes that occurs with climate change or climate variability.

1.3) The role of the oceans

“Although the intricate two-way relationship existing between the Earth’s oceans and climate is not fully understood” (Field et al., 2002), the oceans have a fundamental influence on the global climate system with regard to both variability and change (Field et al., 2002; IPCC, 2007d). The oceans have already reduced the magnitude of human-driven climate change (Fung et al., 2005; Friedlingstein et al., 2006) through the uptake of CO\(_2\). Over the last two centuries, the oceans have been the primary sink for anthropogenic carbon, taking up ~23% of collective fossil fuel emissions in 2008 (anthropogenic atmospheric carbon 9.9 +/- 0.9 PgC y\(^{-1}\) and CO\(_2\) by ocean sinks 2.3 +/- 0.4 PgC y\(^{-1}\)) (Le Quere et al., 2009), thus successfully slowing the growth rate of atmospheric CO\(_2\) and its effects on the earth’s climate.
The short and long term effects of the oceans on life on Earth can be seen in the transport of heat around the globe via the oceans and the corresponding weather patterns that are regulated by ocean-atmosphere coupling. The long-term effect of the oceans on climate however, is of vital interest. The CO$_2$ flux depicted in figure 1 represents the integrated relationship between the oceans and the atmosphere, which affects climate in the long term. With rising atmospheric CO$_2$ concentrations, the mechanisms affecting CO$_2$ flux, it’s basic limitations and influencing features become key to understanding and making possible mitigation plans as “alterations in these fluxes could buffer or enhance climate change” (Field et al., 2002).

![Figure 1: Ocean carbon flux](http://www.acecrc.org.au/Research/Southern%20Ocean%20Carbon%20Sink)

The oceans are an enormous reservoir for carbon (Field et al., 2002), and are currently a net carbon sink of ~1.3 PgCy$^{-1}$ (Sabine et al., 2004b). CO$_2$ is drawn out of the atmosphere by two basic processes: the “physical solubility pump” and the “biological carbon pump”.
1.3.1) The physical solubility pump

The strength of the physical solubility pump is set primarily by temperature and the degree of warming and cooling that, together with salinity, regulates density and the rate of surface water sinking, taking with it atmospheric CO$_2$ (Watson and Orr, 2003). The expulsion of salt during winter seawater freezing and ice formation in Polar latitudes (Warren and Wunsch, 1981) leads to deep-water formation due to sinking of the resulting cold, dense water. Furthermore, as the waters cool, gases (including CO$_2$) become increasingly soluble in it (Cutnell and Johnson, 2007). Subduction of this water therefore carries with it the CO$_2$ signal of the atmosphere, carrying the CO$_2$ into deep reservoirs within the oceans. However, eventual over-turning of the ocean by the thermohaline circulation (Gordon, 1986; Broecker, 1991; Schmitz, 1995) will result in CO$_2$ ‘outgassing’ around 1 000 years later in the Equatorial Pacific, the largest ‘source’ of CO$_2$ to the atmosphere (Petit et al., 1999; Toggweiler, 2008). Over such a long time scales, sequestered CO$_2$, is isolated from further interactions with the atmosphere (IPCC, 2007d).

Removal of CO$_2$ into the deep ocean lowers the concentration of CO$_2$ in surface layers, which creates an atmosphere-ocean concentration gradient that allows further uptake of CO$_2$ by the oceans. It is this physical process that forms part of the carbon sink within the oceans, termed the physical solubility pump.

The oceans have warmed since the 1950’s, with a rise in the average global surface (to 700m) ocean temperature of 0.10°C (IPCC, 2007b; IPCC, 2007d). This is a matter of concern, as the solubility of CO$_2$ in the ocean is an inverse function of temperature (Field et al., 2002). Warming of the oceans therefore threatens the efficiency of the “physical solubility pump”. A thirty-year study currently by Rintoul et al., (in press) further supports
this concern as it reveals that Antarctic Bottom Water formation has declined by up to 60% over that period.

Rising temperatures are less of a concern for phytoplankton communities (Chisholm et al., 2001), which are affected more by other factors associated with temperature, rather than by small temperature changes themselves. However, as warming oceans lead to stronger stratification, which reduces upward nutrient flux, this will negatively influence the biological carbon pump.

1.3.2) The biological pump

The biological pump (Volk and Hoffert, 1985; Longhurst, 1991; Falkowski and Raven, 2007) refers to the transfer of CO$_2$, fixed during photosynthesis, into the deep ocean through the sinking of particulate detritus. Phytoplankton are responsible for approximately half of the carbon fixation on earth even though they make up less than 1% of the earth’s photosynthetic biomass (Falkowski et al., 2000). The fraction of this particulate organic carbon (POC) that is not respired back into surface waters as CO$_2$, sinks below the euphotic zone (Chisholm et al., 2001) into the deep ocean, where it too no longer interacts with the atmosphere. But apart from POC, photosynthesis also produces dissolved organic carbon (DOC) that does not itself sink, but represents a principal store of organic carbon (Tranvik and Jansson, 2002). Where subduction occurs, DOC is exported into the deep ocean, where together with POC; both forms are re-mineralised by bacteria into dissolved inorganic carbon (DIC) (Orr et al., 2005). The deep ocean contains the largest store of mobile carbon on Earth, amounting to about 38,110 PgC, or about 50 times more carbon than is present in the atmosphere (Sabine et al., 2004b; Sabine and Tanhua, 2010), and 10 times more than the earth’s plant and soil carbon stores (Sabine et al., 2004b).
The vast expanse of the oceans, along with the presence of phytoplankton in surface waters ensures that the oceans play a key role in the global carbon cycle and hence climate regulation (Chisholm et al., 2001; Sabine et al., 2004a, 2004b; Orr et al., 2005; Canadell et al., 2007). Each year, about 45% of photosynthesis on Earth occurs in aquatic environments (Falkowski, 1994; Field et al., 1998), the rate of which places an upper bound on the overall biomass and productivity of ecosystems (Falkowski and Raven, 2007). This in turn controls the potential of CO$_2$ uptake by photosynthetic organisms. The ability to influence the rate of photosynthesis thus becomes a relevant interest. As 23% of anthropogenic CO$_2$ is absorbed by the oceans (Le Quere et al., 2009), this makes the oceans a key area of study as a CO$_2$ sink.

The geological record of past climate change (Hughes et al., 2003) is limiting because it cannot predict what future changes in ocean chemistry, due to higher atmospheric CO$_2$, may have on the photosynthetic ability of phytoplankton. Current systems, their physical properties, the phytoplankton communities found within them, and their photosynthetic abilities provide different scenarios for predicting this response. Past climates suffered from unexpectedly rapid climatic shifts over decades or less. These changes were especially prominent at high latitudes, with ice-age transitions being linked to abrupt changes in the North Atlantic circulation (Broecker, 2000; Hughes et al., 2003). As current climate change is approached with unprecedented interference in the atmosphere, there is uncertainty in relating the speed of predicted climate change to the past (Hughes et al., 2003). As marine life is reliant on the biogeochemical status of the ocean, it is heavily influenced by changes in the physical state and circulation (IPCC, 2007d), this provides a further area of uncertainty that one must be aware of when considering climate change and oceanic research.
As the biological pump faces climate change the most pressing concern is the effects that changing surface temperatures will have on the system. Increases in surface temperatures affect the mixed layer depth (MLD) and stratification, which in turn affects the light environment, as well as upward nutrient supply. The interconnected relationship between light and nutrient supply controls the phytoplankton bloom, which regulates CO$_2$ export. Most studies of climate change on phytoplankton production are model based (Cox et al., 2000; Bopp et al., 2001; Fung et al., 2005; Taucher and Oschlies, 2011), showing the same main result: an overall decrease in export production in response to global warming (Taucher and Oschlies, 2011). The metabolic sensitivity of individual phytoplankton species to temperature is also an important component to consider. However more research is needed in this field before conclusive conclusions can be drawn.

One area of concern is potential degradation in the functioning of the biological pump due to changes in ocean pH. Additional absorption of CO$_2$ by the ocean leads to an increase in ocean acidity, while the concentration of carbonate ions decreases. The exact effect of this on marine biology is poorly documented; however, there are two emerging effects. Firstly, as the saturation state of calcite and aragonite in the oceans fall, a community shift is seen away from the abundance of the most pH sensitive calcifying species. For example, the rate at which coral reefs and coccolithophores form Calcium carbonate (CaCO$_3$) decreases and the dissolution rate may increase. As coccolithophores act as ‘ballast’ for other less dense decaying matter, this may reduce carbon export into deeper waters (IPCC, 2007d; Archer, 2009); thus, decreasing the strength of the biological pump.
1.4) The Southern Ocean

The Southern Ocean is found south of 30° S - 40º S, uniquely connecting the Pacific, Atlantic and Indian Oceans. A combination of the active subduction of high latitude surface waters rich in CO$_2$ into the deeper ocean (IPCC, 2007d), and the region’s strong biological carbon pump estimated to be 3 PgC (Schlitzer, 2002), accounting for 20% of the global annual phytoplankton production (Orr et al., 2001; Hassler et al., 2012), entitles the Southern Ocean to be viewed as one of the most important ocean sinks of anthropogenic CO$_2$ (Berger and Wefer, 1991; Caldeira and Duffy, 2000; Sigman and Boyle, 2000) with a disproportionate global impact (Boyd, 2002) on biogeochemical cycles, biodiversity and climate regulation (Hassler et al., 2012).

Of all the world’s oceans, the Southern Ocean contains the highest inventory of unused macronutrients (Levitus et al., 1993; Boyd et al., 2002), yet there is only low and varied phytoplankton biomass and productivity throughout the ocean (Sullivan et al., 1993; Seeyave et al., 2007). It is this high-nutrient-low-chlorophyll (HNLC) nature of the Southern Ocean that makes it so interesting. The limited biomass accumulation and hence limited export of CO$_2$ into deeper waters (de Baar et al., 1997; Blain et al., 2007; Pollard et al., 2009) despite high residual nitrate (NO$_3$) concentrations has made the Southern Ocean a centre for many studies on primary production. One factor known to limit phytoplankton growth in this HNLC ocean is iron (Fe) (Pollard et al., 2009). The absence of a continental land-masses in the Southern Hemisphere makes the Southern Ocean one of the most Fe-impoverished of the world’s oceans (Duce and Tindale, 1991; de Baar et al., 1995; de Baar et al., 1997; de Baar and Boyd 2000; Mahowald et al., 2005; Wagener et al., 2008).
1.5) The role of Fe in the Southern Ocean

The ‘iron hypothesis,’ originally proposed by Martin (1990), espoused elevated phytoplankton production and enhanced carbon export during glacial-interglacial transitions (Sigman and Boyle 2000) due to Fe-stimulated growth. Antarctic and Greenland ice-cores dating from the Last Glacial Maximum revealed a 30x increase in Fe-rich dust flux (Yung et al., 1996; Aumont et al., 2008; Mackie et al., 2008), prompting detailed studies on the role of Fe in the control of phytoplankton growth rates.

The HNLC status of the Southern Ocean is thought to be consistent with a Fe limited regime and this argument has been well documented (Martin and Fitzwater 1988; Martin, 1990; Martin, 1992; de Baar et al., 1995; Coale et al., 1996; Boyd et al., 2000; de Baar and Boyd, 2000; Blain et al., 2001; Gervais et al., 2002; de Baar et al., 2005; Blain et al., 2008; Pollard et al., 2009; Smetacek et al., 2012).

Fe-limitation affects the efficient functioning of phytoplankton, both in terms of their photo-physiology and nitrogen metabolism (Raven, 1990; Cochlan et al., 2008). Fe limitation impairs pigment synthesis, so reducing the efficiency of the electron transport system in photosystem (PS) I and PS II (Behrenfeld et al., 1996). Fe is critical for the assimilation of NO$_3^-$ and is also needed in the biosynthesis of chlorophyll (Miller et al., 1984), without which chlorosis occurs (Lawrence, 2005), substantially decreasing the photosynthetic energy conversion efficiency of the phytoplankton (Laws and Bannister, 1980; Kolber et al., 1988), and so impairs the plants ability to fix carbon. The combination of the above severely affects Fe limited cells, particularly where there is potential light limitation (Sunda and Huntsman, 1997; Lindley and Barber, 1998; Timmermans et al., 2001; Moore et al., 2007a, 2007b).
the high latitude South Ocean, with its low light levels and cloudy conditions, this is of particular importance.

Now, in the face of climate change, where current predictions forecast an increase in atmospheric dust (Hillel and Rosenzweig, 2002) and hence changes in Fe-flux, as well as deepening the MLD (Sarmiento et al., 1998; Boyd, 2002) there are concerns that phytoplankton bloom productivity may decline (Smetacek and Nicol, 2005) concurrent with shifts in species composition; both affecting the efficiency of the “biological carbon pump”.

1.6) Experimental evidence supporting the importance of Fe

To determine whether Fe was indeed a limiting factor in HNLC oceans (Martin 1991, 1990), two forms of experiments began to occur: small-scale open ocean Fe fertilization experiments (Martin et al., 1994; Coale et al., 1996; Boyd et al., 2000; Gervais et al., 2002; Hoffmann et al., 2006; Smetacek et al., 2012), and on-board Fe enrichment experiments (Martin, 1990; Moore et al., 2007a, 2007b). These experiments, summarised by De Baar et al., (1995) and Boyd et al., (2007) have confirmed that Fe limits phytoplankton growth in HNLC oceans.

The first open-ocean Fe enrichment experiment (IronEx1- Martin, et al., 1994) occurred in 1993, in the eastern equatorial Pacific Ocean, showing a “clear unambiguous physiological response to the addition of Fe, resulted in the doubling of biomass, tripling of chl-a [chlorophyll-a] - and quadrupling primary productivity”. Unexpectedly, there was however, no observed NO$_3$ drawdown. Fe limitation of algae blooms in HNLC waters was further confirmed by Coale et al., (1996) (IronEx II), however the link between NO$_3$ uptake
and Fe was found to be contradictory; with nutrient measurements by Martin et al., (1994) indicating “little or no systematic difference in nitrate” while Coale et al., (1996) reported “a strong drawdown of approximately 5 µM nitrate as the biological response developed”.

The success of these experiments, was followed by two small-scale Fe fertilisation experiments in the Southern Ocean (Boyd and Law, 2001; Pollard et al., 2007). Though both showed an increase in phytoplankton productivity in response to the addition of Fe (Chisholm et al., 2001), these experiments differ fundamentally. The Southern Ocean Iron Release Experiment (SOIREE) (Boyd and Law, 2001) was the first in situ Fe fertilization experiment in the Southern Ocean (1999), which showed an increase in chlorophyll and a 10% increase in CO₂ draw down. The Crozet Natural Iron Bloom and Export Experiment (CROZEX) (Pollard et al., 2007, 2009) differed in that it was the first planned natural Fe fertilization experiment, studying a natural Fe-enriched bloom off the Crozet Islands. These two studies are fundamentally important, as together they show an undeniable response, with increased chlorophyll and increased photosynthetic efficiency (Boyd and Abraham, 2001; Moore et al., 2007a, 2007b) in response to both artificial and natural Fe fertilization in the Southern Ocean.

On board Fe enrichment experiments, such as those run in the Ross Sea by Martin, (1990), and on RRS Discovery during CROZEX (Moore et al., 2007a, 2007b) and on MV SA Agulhas in the austral summer of 2010/11, allows phytoplankton responses to Fe and macronutrient additions to be carefully assessed.

1.7) Fe and light co-limitation in the Southern Ocean

The ‘iron hypothesis’ in the Southern Ocean, though strongly supported, is not sufficient to fully explain the dynamics of chlorophyll blooms that occur in this ocean. The
Southern Ocean, being so far South, and often under cloud cover is exposed to low yearly light. Additionally, the development of phytoplankton blooms increases light attenuation with depth, suggesting that light limitation becomes important in areas of deep MLD’s, or near the bottom of the euphotic zone in stratified waters (Sunda and Huntsman, 1997). The co-limitation of Fe and light in the Southern Ocean was first proposed by Raven (1990), who anticipated that at low irradiances, the Fe requirement of phytoplankton would increase with the light-harvesting requirements (Boyd, 2002). This is due to the increased chlorophyll:carbon ratio needed to capture sufficient photons (Venables and Moore, 2010). This co-limitation was confirmed by Sunda and Huntsman, (1997), through culture experiments.

The variation of phytoplankton growth between night and day, highlights the importance of light (Hassler et al., 2012). Using naturally Fe fertilized areas such as downstream of South Georgia, the Crozet and Kerguelen Islands as examples, Venables and Moore, (2010) conclude that although blooms in these areas only begin in spring when light is sufficient, they are not limited by light for the three months of summer.

Satellite imagery coupled with in situ measurements advocate that the Southern Ocean contains more phytoplankton then the available Fe can theoretically sustain, suggesting that Fe is recycled during bloom events. The development of small cells in Fe limited areas, and the resulting increased grazing by microzooplankton, is conducive to material and Fe recycling (Sunda and Huntsman, 1997). This yet unmeasured dynamic complicates the understanding of the Fe-light integration on phytoplankton (Strzepek et al., 2005; Hassler et al., 2012).
Southern Ocean phytoplankton blooms are still poorly understood, doubtless due to the complex and dynamic interplay between Fe availability, chemistry (whether Fe is biologically available or not) and biology in surface waters (Raven, 1990; Boyd et al., 1999; Boyd, 2002; Hassler et al., 2012). However, as current climate projections involving the Southern Ocean predict warming, causing “stratification and an alteration of the MLD” (Sarmiento et al., 1998; Boyd, 2002). By contrast, MLD’s in mid-latitude regions of the Southern Ocean appear to be deepening because of increased wind stress. These projections highlight the importance of understanding the effect of Fe-light limitation in phytoplankton photosynthetic processes in the Southern Ocean.

1.8) Understanding photosynthetic processes

Knowledge of photosynthetic processes in marine organisms provides an understanding of how they might respond to changes in the light environment they experience. Such interpretations are essential to understand the influence of community structure on global biogeochemical cycles in marine environments (Falkowski and Raven, 2007), and hence enhance predictions of possible changes in the future.

Photosynthesis is the process by which plants convert the sun’s energy into chemical energy, which is stored as sugars or other organic molecules. Species of phytoplankton require these organic compounds for phytoplankton growth and reproduction, as well as cellular tissue. The simplified equation of photosynthesis shows the carbon fixation process:

\[ \text{CO}_2 + 2\text{H}_2\text{O} \leftrightarrow [\text{CH}_2\text{O}] + \text{H}_2\text{O} + \text{O}_2 \]
Photosynthesis is divided into two separate phases - the so-called ‘light reactions’ and the ‘dark reactions’. In the ‘light reactions’, light is captured by plant pigments, notably by chl-a, a green pigment located in the chloroplasts. This ‘light reaction’ can be affected by light limitation as it converts solar energy into chemical energy in the form of ATP:

\[ 2\text{H}_2\text{O} + \text{light} \rightarrow 4[\text{H}^+] + \text{ATP} + \text{O}_2 \]

It is this reaction that allows fast repetition rate fluorometry (FRRf) to be used as a measure of the photosynthetic efficiency of phytoplankton. In the ‘dark reactions’, the ATP is used to fix inorganic CO\(_2\) within the Calvin Cycle (Campbell and Reece, 2005), synthesising the production of sugars.

\[ 4[\text{H}^+] + \text{ATP} + \text{CO}_2 \rightarrow \text{CH}_2\text{O} + \text{H}_2\text{O} \]

The effect of Fe limitation effects photosynthesis in two ways. Firstly, it is necessary in PS II and PS I. Secondarily, it is needed in NO\(_3\) assimilation.

1.8.1) The effect of Fe on PS II and PS I

PS II and PS I are concerned with the light phase of photosynthesis. As the majority of cellular Fe is contained in PS II and PS I, this is heavily affected by Fe-limitation (figure 2).
Ultimately, light drives the synthesis of NADPH (Nicotinamide adenine dinucleotide phosphate) and ATP by energizing PS II and PS I, which are both found in the thylakoid membrane of chloroplasts. Non-cyclic flow is the primary pathway of energy transformation in light reactions (includes both PS’s) and is shown in figure 3. This reaction begins with a light photon striking a pigment molecule in the light-harvesting complex of PS II. This photon is passed along the pigment molecules to the P680 chl-a molecule, exciting its electrons to a higher energy state. The excited electron is both captured by the primary acceptor, and replaced in the P680 molecule by splitting a water (H$_2$O) molecule into oxygen (O$_2$) and two hydrogen ions. The excited electron is passed from PS II’s primary acceptor to PS I via an electron transfer chain consisting of the electron carrier plastoquinone (Pq), a cytochrome complex and a plastocyanin (Pc) protein. This transfer reduces the energy level of the electron, hence providing the energy needed for ATP synthesis. This electron replaces an electron captured from P700 by PS I’s primary acceptor (P700 loses its electron through a pathway mirroring that of P680). Similarly, the new photoexcited electron moves down the second transport chain from PS I’s primary acceptor through the ferredoxin (Fd) protein to be

Figure 2: Light and Fe dependency of photosynthesis and NO$_3$ assimilation (Lucas, 2009).
one of the two electrons to connect with NADP$^+$ to form NADPH (Campbell and Reece, 2005).

The effect of Fe limitation on the photosynthetic apparatus means that not all available light can be used. When Fe deficient, phytoplankton not only decrease their absorption of light, but also dissipate (during the electron transfer chains) a large part of the light absorbed by the PS II antenna (Morales et al., 1998).

1.8.2) The effect of Fe in nitrate assimilation

Fe is necessary in the enzymes involved in NO$_3$ reduction by plants (Raven, 1990; Sunda and Huntsman, 1997; Boyd et al., 1999; De Baar et al., 2005; Lucas et al., 2007) as seen in figure 2.
It has been estimated that 80% of Fe required by phytoplankton is used in photosynthesis (Raven et al., 1999). Fe is an essential element required in diverse metabolic pathways: Fe functions as a catalyst in electron transfer reactions, is needed in chlorophyll synthesis and is critically involved in the assimilation of nitrogen (the enzymes nitrate and nitrite reductase both contain Fe and are necessary to reduce NO$_3$ to ammonium (NH$_4$). These combined processes limit the photosynthetic yield under Fe limited conditions (Hassler et al., 2012).

1.8.3) Fast Repetition Rate Fluorometry

FRRf allows one to investigate the photosynthetic efficiency of phytoplankton in terms of PS I and PS II’s electron transport system, indirectly measuring the phytoplankton’s ability to utilize CO$_2$ in carbon fixation during photosynthesis. Thus it measures changes in the basic photosynthetic processes, which can loosely be related to carbon fixation by phytoplankton.

Changes in photochemical reactions and photosynthetic parameters measured by FRRf have been widely used as a diagnostic tool for nutrient-related changes in photosynthetic efficiency (Kolber et al., 1988; Kolber and Falkowski, 1993). The FRRf protocol allows the simultaneous assessment of the parameters in phase two (light reactions) of photosynthesis: i.e. \( \sigma_{\text{PSII}} \) (the functional absorption cross section) and \( \frac{F_v}{F_m} \) (photochemical efficiency, as described by the relationship between variable fluorescence \( [F_v] \) relative to the maximum theoretical fluorescence \( [F_m] \)) (Suggett et al., 2009). This method uses active chl-a fluorescence measurements to evaluate the efficiency by which absorbed light is utilized during photosynthesis (Suggett et al., 2009). Chl-a fluorescence is a
biophysical bi-product re-emitting light not utilized in photosynthesis (Suggett et al., 2009), and so provides a measure of how efficient phytoplankton are at using light.

FRRf measurements are strongly dependent on prior light exposure, thus after a period of dark acclimation, PS II photochemical efficiency is at its maximum, as the quinones (the primary acceptor molecules of PS II) are then fully oxidised. Since the maximum photosynthetic efficiency of phytoplankton decreases under stressful growth conditions (Kolber et al., 1988), this concept has led to the use of FRRf to assess the large-scale photosynthetic condition of entire photosynthetic communities (Behrenfeld et al., 1996; Moore et al., 2005, 2006; Suggett et al., 2006). High values of Fv/Fm and corresponding low values of σPSII indicate a high photosynthetic efficiency, whereas the inverse indicates that phytoplankton are experiencing physiological stress (Holeton et. al., 2005; Suggett et al., 2006).

Variability of these parameters has in the past been attributed to two very different controls; the first being NO₃ or Fe stress (Kolber et al., 1988; Greene et al., 1991; Boyd and Abraham, 2001), and the second being the species composition of phytoplankton (Suggett et al., 2004, 2009; Moore et al., 2005). The high macronutrient concentrations found within the Southern Ocean (Levitus et al., 1993) makes the direct effect of NO₃ limitation in this region obsolete. However, as Fe is required for the efficient uptake and utilization of NO₃, in Fe limited seas, Fe limitation can directly influence photosynthetic efficiency.
1.9) **Community structure**

Southern Ocean phytoplankton communities are mostly dominated by diatoms and haptophytes, frequently *Phaeocystis antarctica* (Boyd, 2002; Hassler et al., 2012). Diatoms dominate mainly near frontal zones (Sakshaug et al., 1991; Laubscher et al., 1993; Smetacek et al., 1997) corresponding with relatively higher Fe concentrations found in these zones (Laubscher et al., 1993; de Baar et al., 1995; Boyd, 2002). There is contradictory information regarding the seasonal shifts in diatom-*Phaeocystis* community structure. In the Ross Sea, haptophytes are particularly noticeable during spring (Arrigo et al., 1999) as *Phaeocystis antarctica*’s photo-physiology is efficient at low irradiance (Boyd, 2002), thus allowing these blooms to appear before those of diatoms. However, off sub Antarctic islands such as Crozet, a spring diatom dominated community is succeeded by smaller taxa in summer as Fe concentrations decline (de Baar and Boyd, 2000; Smetacek et al., 2004; Seeyave et al., 2007).

Limiting nutrient concentrations have a profound effect on the community structure of phytoplankton. Small phytoplankton thrive in Fe-limiting areas because their lower surface:volume ratio allows Fe to be scavenged at low concentrations. Even so, different phytoplankton taxa have various Fe requirements and are known to adapt to environmental changes (Strzepek et al., 2011; Hassler et al., 2012), for example, when facing depleting Fe conditions, diatoms shrink in size (Sunda and Huntsman, 1995). Modification of the light-harvesting antenna can also occur to maximize light photosynthetic efficiency (Michel and Pistorius, 2004). And as a balance between Fe and light efficiency, a decrease in pigment concentration can occur in low Fe conditions (Timmermans et al., 2001).
A dominance by small phytoplankton classes is suggested (Price et al., 1994) to indicate an Fe-limited ecosystem further controlled by microzooplankton grazers. Conversely, Fe alleviation promotes a disproportionate response from larger phytoplankton that can also escape the pressures of grazing (Cullen, 1991; Morel et al., 1991; Price et al., 1994; Hoffmann et al., 2006; Moore et al., 2007a).

“Diatoms appear to be primarily limited by iron supply, because in virtually all iron enrichments there has been a floristic shift toward this algal group” (de Baar and Boyd, 2000). This is further confirmed in a Pacific Ocean study by Coale et al., (1996), where Fe enrichment favoured diatom production and a corresponding draw down of NO$_3$ and silicate (SiO$_4$). The response of *Phaeocystis antarctica* to Fe fertilization is however unknown (Boyd, 2002), although it may encourage colony formation (Lucas et al., 2007).

As few Fe enrichment experiments have been carried out in relation to the vast expanse of the Southern Ocean, let alone the world’s oceans, current knowledge on different community responses to Fe alleviation is relatively sparse. Moore et al., (2007a) stated that the outcome of Fe alleviation experiments was strongly influenced by the initial community structure.

1.10) Rational for this investigation

FRRf allows one to investigate the photosynthetic efficiency of phytoplankton in terms of PS I and II’s electron transport system, indirectly measuring the phytoplankton’s ability to fix CO$_2$ during photosynthesis. Thus it allows measurement of changes in the basic
photosynthetic responses to light and Fe co-limitation and hence indirectly, assess the carbon fixation potential.

1.11) Research hypotheses:

The research carried out in this study strives to test three principal hypotheses:

1) The photosynthetic efficiency of phytoplankton within the Southern Ocean will increase with the addition of Fe. However, the degree to which it increases will differ, depending on the area within the Southern Ocean.

2) The nutrient uptake and chl-concentrations of phytoplankton within the Southern Ocean will increase with the addition of Fe. However, the degree to which it increases will differ, depending on the area within the Southern Ocean.

3) The response of phytoplankton in the Southern Ocean to Fe-fertilization is depended on the initial community structure.
CHAPTER 2: METHODS

2.1) General

All fieldwork was carried out on the South African polar resupply vessel, *MV SA Agulhas*. Data were collected over two summer cruises in 2009/10 and again in 2010/11. The first cruise occurred between Acta Bukta (Antarctica) and South Georgia (figure 4).

Figure 4: Chart of the research area occupied during SANAE 49 in the austral summer of 2009/10. The following stations appear on the map; CTD (blue), productivity CTD stations (blue *), underway UCTD stations (red) and XBT stations (green). Legs are divided into southward (V), north westward (A), south eastward (○) and northward (+). Background map data and script courtesy of Wessel and Smith (1996).
This cruise (AGU 148) was part of the SANAE (South African National Antarctica Expedition) 49 cruise from December - February, 2009/10. The cruise consisted of a Southward leg from Cape Town, South Africa to Acta Bukta, Antarctica (9 December - 22 December), a north westward leg from Acta Bukta to South Georgia (16 - 4 January) and back (25 January - 2 February), followed by a return journey to Cape Town (13 - 23 February).

**Figure 5:** Chart of the research area occupied during SANAE 50 in the austral summer of 2010/11. With sites of the CTD (blue), underway UCTD (red) and XBT stations (green) marked on the map. Fe enrichment experiments are labelled and appear in (yellow *). Legs are divided into, into southward (∇), north westward (∆), south eastward (○) and northward (•). Background map data and script courtesy of Wessel and Smith (1996).
The second cruise (AGU 153) was part of the SANAE 50 cruise from December 2010 - February 2011 (figure 5). This cruise consisted of four legs: a southward leg from Cape Town to Acta Bukta (8 - 19 December), a north westward leg from Acta Bukta to South Georgia (1 January - 10 January), a south eastward Leg from South Georgia to Akta Bukta (10 January - 20 January) and the Northward Good-hope line (5 February - 16 February), from Acta Bukta to Cape Town, South Africa.

2.2) \textit{In situ} measurements and sample collection

2.2.1) \textit{In situ} measurements and sample collection: SANAE 49

On the north westward and south eastward legs of SANAE 49, CTD (conductivity, temperature and [pressure] depth) profiles were conducted (using a Sea-Bird 911+ CTD) every day at 09:00 and at 21:00 for water column sampling to 500m. Water was collected for chemical and biological analysis on the upward cast using 6 x 12 litre Niskin bottles. Two \textit{in situ} samples from the clean seawater inflow to the engine room were collected in-between each CTD cast. A Sippican Deep Blue Expendable Bathythermograph (XBT) and UCTD (underway CTD; produced by Ocean Science) were alternatively deployed every hour (at ~10nm intervals), with XBT deployments increasing to every hour (run concurrently with the UCTD) over main frontal and topographical features. Neither instrument could be used during sea ice conditions. The use of the XBT’s formed part of the long term GoodHope program (funded NOAA’s Office of Global Programs as part of their High Density XBT project at NOAA/AOML).

On the Northward leg of SANAE 49, \textit{in situ} sampling occurred for chemical and biological analysis every four hours. Physical data were collected by alternate UCTD and
XBT’s every hour, except over fronts where an XBT was also discharged concurrently with the UCTD.

On SANAE 49’s Southward leg, *in situ* sampling or chemical and biological analysis took place every four hours.

2.2.2) *In situ measurements and sample collection: SANAE 50*

On SANAE 50, the sampling strategy varied greatly depending on the leg.

The north westward leg consisted of underway stations only. Biological and UCTD stations occurred every 40nm and 20nm respectively. Between 69° S - 71° S and around 62° S, UCTD deployments were interrupted by sea ice, while over the South Georgia shelf UCTD deployments ceased due to the shallow bathymetry.

The south eastward leg was divided into two sampling strategies. The first half of this leg consisted of underway stations and second half (from 58.5° S) consisted of CTD stations. For the CTD leg, CTD’s were spaced 20nm apart with UCTDs deployed between each CTD station. CTDs sampled the water column to 500m, sampling a maximum of 13 depths. Water was collected during the upwards cast of the CTD with full biological and chemical sampling occurring at each CTD. Course changes seen south of 55° S were due to poor weather conditions.

For the northward leg, XBT and UCTD deployments were alternated and occurred every 10nm, increasing to every 5nm over main frontal regions. Biological stations occurred every 40nm, except over fronts where the resolution increased to 20nm. Due to winch
problems, UCTD measurements were only conducted to 50° S, after which XBT’s replaced the use of UCTD’s in this region.

2.2.3) Biological stations

All biological stations included samples for chl-a, High Performance Low Chromatography (HPLC), FRRf and nutrients.

*Underway*

*Underway Biological: SANAE 49*

Discrete underway biological samples on SANAE 49 were collected from the engine room clean water supply pump (approximately 5m below the ocean surface). These water samples were analysed for chl-a, HPLC and nutrients.

*Underway Biological: SANAE 50*

Water for discrete underway biological samples on SANAE 50 was collected from an uncontaminated surface sea water supply using the towed Fe-fish (approximately 1 -5m below the ocean surface, depending on swell conditions). When ice and bad weather prevented deployment of the Fe-fish, samples were collected from the clean engine room supply.

Sample variables from the two water sources were compared to see if there were any statistical differences. For chl-a there was no statistical difference, but there was however a
statistical difference between FRRf results. Thus, when the Fe-fish could not be deployed, FRRf samples were obtained from bucket samples, with no statistical difference from the Fe-fish supply.

Underway biological samples for SANAE 50 were analysed for chl-a, HPLC, and FRRf and nutrients. Nutrients sampled at night were filtered and frozen for analysis the following day.

**CTD**

**CTD Biological: SANAE 49**

Biological CTD stations consisted of the top 6 depths collected in the Niskin bottles. These depths included surface, thermocline and chl-a max depth determined on the downward cast. All biological depths were analysed for chl-a, FRRf and HPLC.

**CTD Biological SANAE 50**

The SANAE 50 CTD biological stations consisted of the top 6 depths collected in the Niskin bottles as before. These depths included surface, thermocline and chl-a max (top and bottom), as well as additional depths below the euphotic layer. All biological depths were analysed for chl-a, FRRf and HPLC.
2.3) **On-deck Fe-light enrichment experiments**

2.3.1) **Setting up enrichment experiments**

Six on-board Fe light incubation experiments were carried out during SANAE 50 (for positions see table 3, under results) to test phytoplankton responses to Fe and light. Experiment 8 and 9 were cut after day 3 and 2 respectively after the sea water supply cooling the incubations was compromised.

For each experiment, 13 x 2L clean, uncontaminated polycarbonate bottles were randomly filled in the Fe-free tent with water obtained from the towed Fe-fish). Six of these bottles were spiked with 100µl of 2µmol FeSO\(_4\) (iron sulphate), while a further six were not spiked with Fe, so were control bottles. Three of the Fe spiked bottles and three of the controls were covered with a neutral density filter to provide 50% shading. A final (13\(^{th}\)) bottle was filled with seawater, without any Fe supplement, and not reopened until day 5 of the experiment. This acted as a control to test for long-term contamination during subsampling.

All bottles were placed in an on-deck incubator covered in 50% neutral density filter and cooled by running surface (5m) seawater. Thus, on-deck incubations were performed for 5 days at two different irradiances as six bottles were exposed to 50% light and six bottles to ~25% light.
2.3.2) Sampling enrichment experiments

Every 24 hours, water was removed from 12 bottles for chl-a, nutrient and FRRf determinations. Samples for nutrient analysis were filtered, frozen and analysed back in Cape Town for NO$_3$, SiO$_4$ and phosphate (PO$_4$). Triplicate samples were taken for time zero chl-a and FRRf analysis. Sub-sampling of the incubations occurred between midnight and dawn to avoid light-shock.

Chl-a and FRRf samples were analysed immediately using the methods described later. As the incubation stations were set up concurrently with full biological stations, the starts of each experiment corresponded with HPLC, chl-a, nutrient and FRRf measurements taken from the CTD bottles.

2.4) Analytical methods

2.4.1) Water masses determined from CTD-profiles

The SeaBird 911plus CTD sensors measured temperature and salinity with each cast. Using Ocean Data View (2008), water densities were determined to establish water mass characteristics based on T-S (temperature - salinity) plots (see Appendix A). Water masses were determined according to Park et al., (1998), Orsi et al., (1995) and Veth et al., (1997).

The CTD’s auxiliary sensors also included a SBE 43 dissolved oxygen and underwater PAR (photosynthetically active radiation) sensor. Dissolved oxygen was calibrated against water samples from selected depths from the CTD casts. These samples were run on a SiS Sensoren Instrumente automated dissolved oxygen system following the
Winkler method for dissolved oxygen measurements in discrete water samples (Carpenter, 1965) and in accordance with World Ocean Circulation Experiment (WOCE) standards.

2.4.2) Phytoplankton photo-physiology (FRRf)

A bench top Satlantic FIRe (Fluorescence, Induction and Relaxation) System was used in discrete mode to measure a comprehensive suite of fluorescence parameters (Fv/Fm and σPSII). These photosynthetic physiological parameters can be used to provide highly sensitive and well-resolved data on phytoplankton community responses to light and nutrients. The system is well described in the Satlantic manual, and in several publications (e.g. Kolber et al., 1998).

The parameters used specifically in this study are: Fv/Fm and σPSII. Fv/Fm represents the photochemical efficiency. This looks at the relationship between the variable fluorescence (Fv) and the maximum theoretical fluorescence (Fm). Calculations for Fv/Fm are as follows:

\[ \frac{F_v}{F_m} = \frac{(F_m - F_o)}{F_m} \]

Where: Fv = variable fluorescence

Fm = Maximum theoretical fluorescence

Fo = minimum fluorescence yield

σPSII represents the functional absorption cross section, and is calculated as the slope between Fo and Fm. Figure 6 is an example of the FIRe measurement protocol, showing the mathematical position so of above mentioned parameters.
Each sample for FRRf measurements was collected in a clean dark plastic bottle and placed in a dark environment for half an hour prior to FRRf measurements in the FRRf cuvette (Behrenfeld et al., 1996; Moore et al., 2005, 2006; Suggett et al., 2006). Just prior to placing the sample in the cuvette, the sample was inverted to ensure an even distribution of phytoplankton in each sample. These procedures ensured completion of the photosynthetic cycle at the time of collection so as not to interfere with the FRRf readings. The FRRf readings were also conducted in minimum light conditions to reduce outside light interference.

The Satlantic FIRe instrument was set at the following settings for each site:

Number of automatic samples: 16 (for SANAE 49) or 25 (for SANAE 50)
LSTF 100, STRP 60, STRI 60, MTF 600, MTRP 60, MTRI 100.

Values of $F_v/F_m$ and $\sigma_{PSII}$ were calculated by fitting a best-fit line to the measured saturation curves and by running the results through a MATLAB (MATrix LABoratory...
R2008a) script code (see Appendix B) to remove the first measured value, as required by the bench top Satlantic FIRe Systems calibration procedure. The script used (courtesy of Dr Brian Hopkinson) was validated against the one used by Dr Mark Moore (Appendix C) during CROZEX (Moore et al., 2007b). A comparison of the two scripts confirmed that interpretations were independent of model choice. Any values above 10% error levels were removed from the data set.

Blanks were run (at the gain setting of that station) for each site to be included in the photosynthesis parameter calculations. These were done by filtering a fraction of each sample under positive pressure through a 25mm Whatman glass fibre filter (GF/F) to remove all biological material, and the filtrate was then run through the Satlantic FIRe System at the same settings as its corresponding sample. Blanks were run both on board (SANAE 50 only) as well as back at the University of Cape Town (UCT) (SANAE 49 and SANAE 50). Sample fluorescence values were corrected for the blank values in MATLAB (Appendix B).

*SANAE 49 Sampling*

For the SANAE 49 cruise, water samples from the top six CTD depths were collected to measure active phytoplankton community fluorescence through the water column. At each depth, nine replicate samples were run on the FIRe instrument.
**SANAE 50 sampling**

**CTD**

For the SANAE 50 cruise, water samples from the top six CTD depths were run as above. At each depth, three replicates samples were run on the FIRe instrument. The reason for the difference in number of replicates between SANAE 49 and SANAE 50 was time constraints. The large number of samples required to be run in the experimental stage of SANAE 50, the time it took, and available personnel, lead to the decrease in the number of replicates on SANAE 50 compared to SANAE 49.

**Underway**

*In situ* FRRf samples were taken as part of the biological sampling, every four hours. As for biological samples, water was collected from the uncontaminated surface Fe-fish supply. When, however, the Fe-fish could not be deployed, the FRRf sample was collected with bucket samples. Due to time and personnel constraints during the incubations, the FRRf 4am sample was not run. This corresponded with the biological station that did not include a Fe sample. At each station, three replicates samples were run on the FIRe instrument.

**Fe Incubations**

At the beginning of each Fe-incubation, three replicate samples were run on the FIRe instrument. Once the incubations were running, the sub sampling saw that one sample was run from each bottle, with a corresponding blank.
Background Statistical check

For the biological station data, a brief comparison between the $F_v/F_m$ and $\sigma_{PSII}$ parameters and the time of day was run to see if the phytoplankton where being negatively affected by radiation through photo-inhibition. It was found that the time of day did not influence the $F_v/F_m$ and $\sigma_{PSII}$ parameters.

2.4.3) Nutrients

Nutrients were analysed in two different ways on the two cruises. On SANAЕ 49 the nutrients were analysed manually, while on SANAЕ 50 they were analysed automatically.

SANAE 49: Manual analysis of nutrients

Samples for analysis of inorganic nutrients were drawn from Niskin bottles from all depths sampled and analysed immediately. Concentrations of SiO$_4$, PO$_4$, NO$_3$, NH$_4$ and urea were analysed according to manual methods described in Grasshoff et al., (1983) and Parsons et al., (1984), scaled to 5 ml sample sizes.

Problems encountered with silicate on SANAÉ 49

Frozen back-up SiO$_4$ samples were re-run back at the UCT, using the automatic method after the manually run samples were found to be far too high. This occurred only for SiO$_4$ samples in very low SiO$_4$ regions, such as north of the Subtropical Front (STF).
Problems encountered with phosphate on SANAE 49

Manual analysis of PO$_4$ is run with a concurrent analysis of a standard solution. On SANAE 49, the originally standard solution of PO$_4$ was double the concentration that it was required to be. Due to the high concentration of this standard, and the sensitivity range of the test, one could not simply half the results of the standard. The standard is used in the calculation of the sample. Thus, to gain the true PO$_4$ value from the samples, a correction needed to be applied. This was corrected by using the average corrected standard (0.92 µmol.L$^{-1}$) from SANAE 48, the previous annual cruise run at the same time, using the same methods, through the same waters.

Problems encountered with nitrate on SANAE 49

On the Northward Leg, after 65.8° S, the ammonium chloride (used as a buffer for the cadmium column after it is repacked [Mostert, 1983]) ran out, although samples were still run. A comparison between SANAE 48 and SANAE 49 NO$_3$ and PO$_4$ data as well as the NO$_3$:PO$_4$ ratio (Appendix D) revealed that the exclusion of ammonium chloride in the method made a substantial difference (the greatest difference between the two years, at the same latitudes, reaching 20 µmol l$^{-1}$). This difference was not initially picked up by the standard, due to the difference in pH between the fresh water standard and sea water samples. See Appendix D [a]) for the NO$_3$ readings. An attempt was made to rebuild the NO$_3$ data using the combined SANAE 48 and 49 NO$_3$:PO$_4$ ratio ($y = 0.0634x + 0.2041$, Appendix E). As the analysis of NO$_3$ is salt sensitive, this data had to be corrected using the above mentioned ratio. The combination of this correction and the PO$_4$ correction does undermine the quality of this data set.
SANAE 50: Automated nutrient analysis

The Lachat QuikChem 8500 series 2 Flow Injection Autoanalyzer was used to measure NO$_3$ and SiO$_4$ concentrations on SANAE 50. However PO$_4$ was still determined manually according to the method described in Grasshoff et al., (1983) and Parsons et al., (1984).

All nutrient samples were measured during the day. Hence samples taken during the day were analysed that same day, but nutrient samples collected at night were filtered and frozen to be analysed a few days later.

All incubation nutrient samples were filtered (through Whatman GF/F), frozen, and analysed at the UCT after the cruise. Unfortunately this process renders the SiO$_4$ analyses suspect at best, or simply invalid.

The differences in the time lines between the underway and incubation nutrient analysis was due to an unfortunate and frustrating set of personnel, political and funding events, both on the ship and on the land. It is understood that the delay in analysis of the incubation nutrient samples compromises this data sets quality (see appendix J).

2.4.4) Chlorophyll-a

Chl-a concentrations were determined for the top six CTD depths and for the in situ underway biological samples. Chl-a samples were collected by vacuum filtering 250 ml of seawater through 25 mm Whatman GF/F filters to trap phytoplankton cells. Chl-a pigment
was extracted from the filter with 7 ml (SANAE 49) or 8 ml (SANAE 50) of 90% acetone over a 12 - 24 hour period a dark fridge. Samples were then read on a Turner Designs Trilogy fluorometer that was calibrated with chl-a standards (Sigma, UK) before the voyage. The sample chl-a concentration (μg.l⁻¹) was derived using the calibration curve regression.

2.4.5) Community structure: (HPLC)

For both SANAE 49 and SANAE 50, community composition was estimated from diagnostic pigment composition and the pigment ratios measured using HPLC for both CTD and in situ underway samples. For HPLC analysis, water samples of 1 - 2 L (depending on particulate loads), were filtered under positive pressure through 25 mm Whatman GF/F filters to capture phytoplankton. These filters were then stored immediately in liquid nitrogen.

Post-cruise, HPLC samples were stored in a -80ºC freezer. HPLC analyses were conducted at the National Oceanography Centre (NOC), Southampton. Phytoplankton cells were raptured in 90% acetone in a Sonics & Materials Inc. Vibracell sonicator (run for 30 s). Pigments were extracted by centrifugation using a MSE Mistral 1000. After filtering through a 0.2 μm filter, sample pigments were analysed on a thermo separation product following the protocol of Gibbs as described by Barlow et al., (1997). This involves pigment separation through a 3 μm Hypersil MOS2 C8 column, followed by detection by absorbance and identification by retention time and online diode array spectroscopy.

Analysis of raw pigment signatures was done using ChromQuest 4.1 software. Pigment information was to characterise phytoplankton species, described by Wright and Jeffery (2006) and Jeffery et al., (1997) (see Appendix F). For each species, the average
percentage of occurrence in the population was determined. Only species that occurred on average in more than 5% of the population were included in this study.

2.4.7) Physics

UCTD and XBT’s

XBT’s and UCTD’s produce salinity and temperature profiles and were used on the cruise to determine frontal positions. UCTD data were processed using SeaBird Seasoft, and validated against CTD measurements.

CTD’s

CTD measurements included a SBE 43 Dissolved Oxygen Sensor, a fluorometer, an underwater PAR unit and a surface reference PAR sensor.

Underway PAR

The underway PAR sensor was mounted on the port side of the ships ‘monkey island’ and provided incoming irradiance measurements (every 30 seconds). Average PAR for each experiment was calculated using standard statistical methods (average = sum of irradiance for full experiment/number of observations). This was to calculate the full irradiance available per experiment.
2.4.8) Nitrogen uptake and f-ratio

To provide a measure of nitrogen and carbon export, the f-ratio (nitrate uptake / total nitrogen uptake \( \frac{\rho \text{NO}_3}{\text{Total } \rho N} \)) was calculated for the surface waters using stable isotope \(^{15}\text{N} \) uptake techniques (see Dugdale and Goering, 1967 and Lucas et al., 2007). Under appropriate time and space scales, the f-ratio represents the fraction of primary production (as PON [particulate organic carbon]), fuelled by NO\(_3\), that is potentially exported to the deep ocean through the sinking of phytoplankton particles. This can be translated into carbon (POC) export if the canonical Redfield Ratio of 6:1 (C:N) is adopted (Redfield, 1963). Uptake measurements were made at eight stations during the SANAE 49 cruise (Figure 4).

_Underwater irradiance and sample depths_

At each of the eight CTD stations, water samples were collected at the following light depths: 86%, 47%, 15%, 6%, 3.5% and 0.7% surface irradiance. These light depths were calculated using an underwater PAR sensor on the downward cast of the CTD. After converting PAR values into natural log values, the following equation was used to determine the depth for each sample:

\[
w = (-1/k) \times \ln(\text{LD}/100)
\]

where: \( k \) = the slope of the natural log of PAR vs. depth

\( \text{LD} \) = light depth (%)

\( W \) = depth which represents that light percentage (m)

The linear nature between PAR and depth plots within the Southern Ocean allowed \( k \) to be calculated as described above.
Sea-water samples from each depth were decanted into 3 x 1.0 L polycarbonate bottles. To measure simultaneous ρNO₃ and carbon fixation, one bottle from each depth was spiked with ¹⁵N-NO₃ (1 µmol K¹⁵NO₃ / 100 µl) and with ¹³C (49.4 µmol / 100µl). To separate bottles at each depth, NH₄ (0.1 µmol ¹⁵NH₄Cl / 100 µl) and urea spikes (0.1 µmol CO (¹⁵NH₂)₂ /100 µl) were also added at ~10% of ambient concentrations to avoid stimulating production.

After spiking, the bottles were placed in simulated in situ on-deck incubators for 24 hours, which were shaded to the appropriate light depth with neutral density filters, and cooled to ambient sea surface temperatures with running seawater.

At the end of the incubations, all samples were filtered onto pre-ashed Whatman GF/F filters (25 mm) and stored frozen prior to being run on UCT’s Thermo Finnegan Mass Spectrometer in the Archaeometry Department. Calculation of uptake rates followed the protocol of Dugdale and Goering (1967), as well as Lucas et al., (2007).

As calculation of update rates (and thus f-ratios) are strongly dependent on accurate nutrient measurements. A sensitivity study was undertaken, this sensitivity study looked at what changes would occur to the f-ratio if there was a 10% variation in the NO₃ data. Results of this study can be found under section 3.1.4.2.
2.5) **Statistical analyses**

2.5.1) **Statistical analyses of the controls on FRRf parameters**

To determine any factors that could be controlling the $F_v/F_m$ and $\sigma_{PSII}$ values (code in Appendix G), linear regressions between variables were run using the programme R (version 2.14.0, 2011). A comparison between the f-ratio, $F_v/F_m$ and $\sigma_{PSII}$ allowed one to search for a relationship between the photosynthetic “health” ($F_v/F_m$ and $\sigma_{PSII}$) and carbon export to the deep ocean (f-ratio), based on the premise that healthy Fe-replete diatoms are likely to assimilate significant amounts of NO$_3$, as reflected in a high f-ratio (Lucas *et al.*, 2007).

2.5.2) **Statistical analysis of results from Fe incubations experiments**

For the Fe enrichment experiments, a paired-sampled t-test was conducted to separately compare the $F_v/F_m$, $\sigma_{PSII}$ and chl-a, values under the following conditions:

1) The control bottles and the Fe addition bottles at 50% light levels
2) The control bottles and the Fe addition bottles at 25% light levels
3) The control bottles at 50% light levels and 25% light levels
4) The Fe addition bottles at 50% light levels and 25% light levels
5) All the control bottles and all the Fe addition bottles
6) The contamination control bottle and the control bottle at 50% light level (this was only done for experiments: B, C and D).

This was done in Microsoft Excel (2010) to calculate any significant difference between the two light levels, for the Fe alleviated bottles and the controls.
CHAPTER 3: RESULTS

SECTION 1: Observational Results

3.1.1) Antarctic Sea Ice Data

Table 1: A comparison of the total sea Antarctic sea ice area for the duration of the SANAE 49 and SANAE 50 cruises (NSIDC, 2012)

<table>
<thead>
<tr>
<th></th>
<th>Antarctic Sea Ice Area (million square km)</th>
</tr>
</thead>
<tbody>
<tr>
<td>December</td>
<td>6.9</td>
</tr>
<tr>
<td>January</td>
<td>3.2</td>
</tr>
<tr>
<td>February</td>
<td>2</td>
</tr>
</tbody>
</table>

Table 1 reveals that the area covered by sea ice was greater for the full three months duration of the SANAE 49 (2009/10 cruise) compared to SANAE 50 (2010/11).

3.1.2) Water masses

The temperature sections (figures 7, 8) from the north westward legs of both SANAE 49 and SANAE 50, between Antarctica and South Georgia (see figure 4), show evidence of a subsurface temperature minimum layer. This Winter Water (WW) is cold, fresh and oxygen rich (Park et al., 1998). There is a strong thermocline dividing this water from the Antarctic Surface Water (AASW) and Upper Circumpolar Deep Water (UCDW). The UCDW and Lower Circumpolar Deep Water (LCDW) is distinguished by salinity differences, with UCDW having low salinity water found between densities of 27.35 kg m$^{-3}$ – 27.75 kg m$^{-3}$, while the younger, LCDW has low salinity water with a density above 27.75 kg m$^{-3}$ (Veth et al., 1997).
On the north westward leg of SANA E 49 (figure 7), the southern boundary of the Antarctic Circumpolar Current (SBdy) is positioned at 60.45° S, with the Southern Antarctic Circumpolar Current Front (SACCF) at 57.97° S. On the south eastward leg, the SBdy and SACCF shifted to 58.8° S and 58.97° S respectively. On SANA E 50 the SBdy and SACCF, although less distinct, were placed at 56.7° S or 57.3° S and 50.17° S on the south eastward leg respectively.

MLD is deeper for the south eastward legs than the north westward legs of both SANA E 49 and 50. This is specifically noticeable south of the SBdy (figures 7 and 8).

**Figure 7:** Vertical Temperature (°C) salinity (PSU) and density (kg m⁻³) sections on the north westward (left) and south eastward (right) legs of SANA E 49. The positions of the SBdy and the SACCF are shown with a grey and black lines respectively. The dotted black line at 60°S on SANA E 49’s south eastward leg represents more than one CTD deployment at that latitude (see figure 4). CTD and UCTD stations are marked with a triangle and circle respectively.

Any discontinuity seen in the plots around 60° S on the south eastward leg of SANA E 49 (figure 7) were due to numerous CTD’s been conducted at one latitude (different
longitudes). This was due to an unfortunate set of weather conditions which forced the ship to follow this particular route (see figure 4).

The northward legs, from Antarctica to Cape Town, show little variability between the two years (figure 9). The presence of WW is still evident close to Antarctica, with waters warming as one approached South Africa. The presence of Agulhas Rings south of South Africa is evident during SANAE 50’s northward leg south of the Southern STF (figure 9). During SANAE 50, the Northern STF (39.33° S), Southern STF (40.56° S), Subantarctic Front (SAF) (44.01° S), Antarctic Polar Front (APF) (49.36° S), SACCF (53.22° S), SBdy (55.41° S) of the Antarctic Circumpolar Current (ACC) fronts were clearly present at the latitudes indicated. During SANAE 49, the fronts were positioned as follows: Southern STF (41.59° S), SAF (43.96° S), APF (50.08° S), SACCF (53.39° S) and SBdy (55.62° S).

Figure 8: Vertical Temperature (°C) salinity (PSU) and density (kg m\(^{-3}\)) sections on the north westward (left) and south eastward (right) legs of SANAE 50. The positions of the SBdy and the SACCF are shown with a grey and black line respectively. CTD and UCTD stations are marked with a triangle and circle respectively.
3.1.3) Nutrient concentrations

3.1.3.1) Nitrate

NO$_3$ concentrations in surface waters (AASW and upwelled UCDW) of the north westward leg of both cruises (Figures 10, 11) exceeded 25 µmol l$^{-1}$.

During SANAE 49, a NO$_3$ maximum (34.87 µmol l$^{-1}$) penetrated into AASW near South Georgia (figure 10), extending to just north of the SACCF (north westward leg). LCDW (lower circumpolar deep water) also contained high concentrations of NO$_3$, which mixed into the WW layer, reaching a maximum concentration (> 34.9 µmol l$^{-1}$) at about 60º S. NO$_3$ concentrations for AASW along the south eastward leg exceeded 16.7 µmol l$^{-1}$, with
minima concentrations occurring between 65.1° S and 62° S, as well as between 54° S - 56° S. On this leg the NO₃ concentrations increased substantially (>30 µmol l⁻¹) in deep waters (>500m). In increase in NO₃ concentrations is noted in the deeper waters between 62° S and 66° S on the south eastward leg.

Figure 10: Vertical section of a) NO₃ (µmol l⁻¹) from the north westward, b) south eastward and c) northward legs of SANAE 49. The SBdy and the SACCF are shown in (a) and (b) with a grey and black line respectively. The thin black line in (b) represents the region where more than one station occurred at 60° S on the south eastward leg.

During SANAE 50 along the south eastward leg, NO₃ concentrations decreased throughout the water column between 67° and 61.5° S, where surface concentrations were 15 µmol l⁻¹ (figure 11). Similar concentrations were evident in the surface waters just off Antarctica.

Along the northward legs of both SANAE 49 and SANAE 50 cruises, there were clear southward increases in NO₃ concentrations. Notable increases occurred before the fronts (figure 10 and 11), specifically at the SAF.
3.1.3.2) Silicate

SiO$_4$ concentrations observed during SANAE 49 varied little between the north westward and south eastward legs (figure 12). Concentrations generally exceeded 40 µmol l$^{-1}$, with maximum concentrations (>70 µmol l$^{-1}$) evident below 350m between the SBdy and SACCF, as well as below 450m north of 55º S. Minimum values (5.45 µmol l$^{-1}$) were observed in the AASW near South Georgia on the north westward leg, steadily increasing with depth to 150m.

In surface waters between Antarctica and South Georgia, SiO$_4$ concentrations exceeded 70 µmol l$^{-1}$ south of the SACCF (figure 13). Between this front, and South Georgia, concentrations decreased sharply to a minimum of 4.45 µmol l$^{-1}$. Along the south eastward leg of SANAE 49, maximum SiO$_4$ concentrations reached 150 µmol l$^{-1}$ below 150m between 67º and 61.5º S.
Figure 12: Vertical section showing SiO$_4$ (µmol l$^{-1}$) distributed on a) the north westward, b) the south eastward leg and c) the northward legs of SANAE 49. The SBdy and the SACCF are shown in (a) and (b) with a grey and black line respectively. The thin black line in (b) represents the region where more than one station occurred at 60º S on the south eastward leg.

Figure 13: Surface distribution of SiO$_4$ (µmol l$^{-1}$) along SANAE 50’s north westward, south eastward and northward legs. Vertical section of SiO$_4$ was obtained from the CTD line during the south eastward leg.

Low values of SiO$_4$ (<40 µmol l$^{-1}$) were found north of the APF on the northward legs of both SANAE 50 and SANAE 49 cruises, as well as north of the SACCF on the other legs (figures 13, 12).
3.1.4) **Phytoplankton**

3.1.4.1) **Chlorophyll-a**

*Figure 14: Vertical sections of chlorophyll (µg.l⁻¹) distribution on a) the north westward, b) the south eastward and c) the northward leg of SANAE 49. The SBdy and the SACCF are shown in (a) and (b) with a grey and black line respectively. The thin black line in (b) represents the region where more than one station occurred at 60° S on the south eastward leg.*

Between Antarctica and South Georgia (SANAE 49) there were two main areas of moderate chl-a concentrations (>1 µg.l⁻¹) (figure 14). These blooms, although apparent on both legs, were deeper (>100m), with a higher chl-a maxima (2.5 – 3 µg.l⁻¹) on the south eastward leg. These blooms occurred in surface waters from 67° S to just South of 60° S and again near South Georgia. Chl-a concentrations (SANAE 50; figure 15) and (SANAE 49; figure 14) between Antarctica and South Georgia were not similar. Though, SANAE 50’s south eastward leg revealed a chl-a bloom between 67° S and 62° S (similar in position to one of the SANAE 49 blooms), the north westward leg of SANAE 50 does not reveal a bloom in this position (figure 15). Both the north westward and south eastward legs of SANAE 50 see an increase in chl-a near South Georgia.
Along the northward legs of SANAE 49 (figure 14) and SANAE 50 cruises (figure 15), chl-a concentration decreased (from 2.57 to 0.03 µg.l\(^{-1}\) and from 3.65 to 0.26 µg.l\(^{-1}\) respectively) with distance away from Antarctica, but increased at each of the fronts.

3.1.4.2) Nitrogen uptake and f-ratio

The f-ratio is only available on the SANAE 49 cruise. On the north westward leg; low f-ratios (0.022 to 0.06) characterised deeper waters towards South Georgia, while the highest f-ratio (f = 0.37) present in AASW (>20m depth) near Antarctica (figure 16). On the south eastward leg, low (0.016 to 0.06) f-ratios characterised the water column between the SBdy and 65° S, with increasing f-ratios (up to 0.22) on either side.
Table 2: f-ratio sensitivity test results, showing changes in f-ratio caused by a +/-10% variation in ambient NO$_3$.

<table>
<thead>
<tr>
<th>Station</th>
<th>latitude</th>
<th>Longitude</th>
<th>depth</th>
<th>Fratio -10% ambient NO$_3$</th>
<th>ambient NO$_3$</th>
<th>+10% ambient NO$_3$</th>
<th>% difference in f ratio</th>
</tr>
</thead>
<tbody>
<tr>
<td>BR-10</td>
<td>-65.52</td>
<td>-6.06</td>
<td>42</td>
<td>0.35</td>
<td>0.37</td>
<td>0.39</td>
<td>5%</td>
</tr>
<tr>
<td>BR-22</td>
<td>-61.15</td>
<td>-17.00</td>
<td>40</td>
<td>0.09</td>
<td>0.10</td>
<td>0.11</td>
<td>8%</td>
</tr>
<tr>
<td>BR-34</td>
<td>-57.68</td>
<td>-30.66</td>
<td>63</td>
<td>0.06</td>
<td>0.06</td>
<td>0.07</td>
<td>9%</td>
</tr>
<tr>
<td>BR-44</td>
<td>-54.14</td>
<td>-36.41</td>
<td>64</td>
<td>0.04</td>
<td>0.04</td>
<td>0.05</td>
<td>9%</td>
</tr>
<tr>
<td>BR-52</td>
<td>-54.34</td>
<td>-27.50</td>
<td>34</td>
<td>0.20</td>
<td>0.22</td>
<td>0.24</td>
<td>7%</td>
</tr>
<tr>
<td>BR-64</td>
<td>-60.00</td>
<td>-17.25</td>
<td>63</td>
<td>0.07</td>
<td>0.07</td>
<td>0.08</td>
<td>8%</td>
</tr>
<tr>
<td>BR-76</td>
<td>-61.83</td>
<td>-5.00</td>
<td>79</td>
<td>0.03</td>
<td>0.03</td>
<td>0.03</td>
<td>9%</td>
</tr>
<tr>
<td>BR-89</td>
<td>-70.04</td>
<td>-7.93</td>
<td>66</td>
<td>0.14</td>
<td>0.16</td>
<td>0.17</td>
<td>7%</td>
</tr>
</tbody>
</table>

Table 2 shows that a 10% variation in ambient NO$_3$ causes a less than 10% (5-9%) change in the f-ratio.

3.1.4.3) Phytoplankton Community Composition from HPLC

The community structure is reported here for dominant phytoplankton groups: i.e. those that represent more than 5% of the population on average. The data reveals only the north westward, south eastward (figure 17) legs of the SANAE 49 cruise. The data are unfortunately patchy due to loss of samples that were stored with the SANAE 50 samples, which were lost due to thawing.

Diatoms

Diatoms dominated much of the Southern Ocean between Antarctica and South Georgia, where they made up 80% of the populations.
Diatoms were abundant on both the north westward (45 - 88%) and south eastward (43 - 79%) legs from South Georgia to the SBdy (figure 17). Higher diatom abundance (73 - 60% and reaching 65% respectively) was evident closer to Antarctica. Relatively lower diatom abundances (>40%) characterised the open-ocean basin between the fronts, with a maximum (71%) just north of 64º S on the north westward leg.

**Figure 17:** Vertical section of a) diatoms, b) haptophytes, c) chromophytes and d) Fuc:Hex from the north westward (left) and south eastward (right) legs of SANAE 49. The SBdy and the SACCF are indicated using a grey and black line respectively. The dotted line represents the region where more than one station occurred at 60º S on the south eastward leg. Red squares are used to highlight areas of interest.

**Haptophytes**

Inversely to diatoms, haptophytes were least abundant near South Georgia and Antarctica (<30%), with their highest abundances (53 - 88%) found in the open ocean south of the SBdy (figure 17), specifically on the north westward leg.
Chromophytes

Chromophytes made up a much smaller proportion (0.1 - 25%) of the phytoplankton community structure, compared to diatoms and haptophytes. This group rose in proportion close to Antarctica (reaching 25% on the south eastward leg only), South Georgia (20% and 25% for the north westward and south eastward legs respectively), on the SACCF (~20%), south of the SBdy (~20%) and in the case of the north westward leg, just north of 66° S (21%) (figure 17).

Fuc:Hex

The diatom to haptophyte ratio (Fuc:Hex [Fucoxanthin:19’-hexanoyloxyfucoxanthin]) rose close to South Georgia (reaching 8.8 and 11) and Antarctica (6.5 and 4.8) on both the north westward and south eastward legs respectively (figure 17), This increase in Fuc:Hex ratio extended further south (from South Georgia) on the south eastward leg, compared to the north westward leg. On the north westward leg, this ratio also increases in the subsurface waters just north of 64° S, and on the SACCF.

3.1.4.4) Phytoplankton photo-physiology (FRRf)

$F_v/F_m$

The photo-physiological health of phytoplankton in this study is represented by characteristic $F_v/F_m$ and $\sigma_{PSII}$ values, as shown in the SANAE 49 (figures 18 and 20 respectively) and SANAE 50 sections (figures 19 and 21).
On SANAE 49 the highest $F_v/F_m$ values ($0.50 \pm 0.012$) were found near South Georgia, on the south eastward leg (figure 18). The distribution of the $F_v/F_m$ maxima (~0.45) around South Georgia differed between the south eastward and north westward legs. The north westward leg revealed that this maximum extended from South Georgia to just beyond 55º S, and throughout the water column to below 100m depth. On the south eastward leg, this maximum (~0.45) was present only below 50m, from South Georgia to 60º S. The $F_v/F_m$ values along the north westward leg, near Antarctica, revealed a similar pattern to the south eastward leg, though with more moderate values ($0.22 \pm 0.017 - 0.43 \pm 0.023$) then the south eastward leg. Low $F_v/F_m$ values (~0.12) were evident in the surface waters near Antarctica and in the middle of both legs. At about 60º S on the north westward leg, an increase in the $F_v/F_m$ values ($0.45 \pm 0.02$) in deeper waters was apparent.
The surface $F_v/F_m$ values on SANAE 50 (figure 19) are continuously moderate to low (<0.2), with an exception of near South Africa (0.38 ± 0.063 - 0.49 ± 0.023), on the northward leg, around South Georgia (reaching 0.42 ± 0.007) on both the north westward and south eastward legs, as well as South of 60º S on the south eastward leg (reaching 0.43 ± 0.048) above topographical features, all of which show high $F_v/F_m$ values. The deeper $F_v/F_m$ values seen on the south eastward leg shows pulses of increased $F_v/F_m$ values under 40m (>0.35) between 61º and 70º S.

$\sigma_{PSII}$

On the north westward leg of SANAE 49 (figure 20), minimum $\sigma_{PSII}$ values (152 ± 6.5 - 185 ± 7) were found near South Georgia where $\sigma_{PSII}$ remained below 200 over the full 100m depth until ~55º S. Such minimum values were also evident in the surface waters between 65º S and 67º S and below 30m between 62º S and 63º S. Maximum values of $\sigma_{PSII}$ (>260) were
found north of the SBdy until ~57° S; crossing both sides of the SACCF. These values were also present below 60m between 65° - 70° S, and in surface waters close to Antarctica and between 62° - 65° S.

**Figure 20**: Vertical section of $\sigma_{\text{PSII}}$ from the north westward (left) and south eastward (right) legs of SANAE 49. The SBdy and the SACCF are indicated using grey and black lines respectively. The dotted line represents the region where more than one station occurred at 60° S on the south eastward leg.

On SANAE 49, the south eastward leg (figure 20), minimum values of $\sigma_{\text{PSII}}$ (162 ± 38.2 - 200 ± 49) were evident in surface waters between 65° - 67° S, and at 60° S. Lowest values (156 ± 81 - 200 ± 49) were found at 50m depth, south of the SBdy. Maximum $\sigma_{\text{PSII}}$ values (253 ± 17.6 - 308 ± 73.6) occurred closest to South Georgia and below 30m between 65°-70° S, as well as in surface waters near Antarctica and between 62° - 65° S.

There is less evidence of a pattern of $\sigma_{\text{PSII}}$ from SANAE 50 (figure 21). $\sigma_{\text{PSII}}$ Values varied between 102 ± 51.1 and 336 ± 17.1. On the northward leg, these surface values are low (<200) north of 50° S. In the depth profile on the south eastward leg, there is an overall increase in $\sigma_{\text{PSII}}$ values as one moves south from ~63° S (~200 to ~300).
3.1.5) \( \text{SiO}_4^+ : \text{NO}_3^- \) Ratios

3.1.5.1) \( \text{SiO}_4^+ : \text{NO}_3^- \) Ratios: SANAE 49

For SANAE 49, the combined north westward and south eastward legs (figures 22) reveal a distinct drop in \( \text{SiO}_4^+ : \text{NO}_3^- \) ratio (dropping below 1) around South Georgia.
3.1.5.2) SiO$_4$:NO$_3$ Ratios: SANA 50

Table 3 shows the SiO$_4$:NO$_3$ ratio, SiO$_4$ difference and NO$_3$ difference in different ocean regions (divided by fronts) in the different legs of SANA 50

<table>
<thead>
<tr>
<th>Ocean Region</th>
<th>SiO$_4$:NO$_3$</th>
<th>difference in SiO$_4$</th>
<th>Difference in NO$_3$</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Southward leg</td>
<td>Northward leg</td>
<td>Northwestward leg</td>
</tr>
<tr>
<td>Between South Africa and northern STF</td>
<td>1.62</td>
<td>4.77</td>
<td>-</td>
</tr>
<tr>
<td>Between northern STF and southern STF</td>
<td>0.39</td>
<td>0.38</td>
<td>-</td>
</tr>
<tr>
<td>Between southern STF and SAF</td>
<td>0.22</td>
<td>0.16</td>
<td>-</td>
</tr>
<tr>
<td>Between SAF and APF</td>
<td>0.39</td>
<td>0.12</td>
<td>-</td>
</tr>
<tr>
<td>Between APF and SACCF</td>
<td>1.55</td>
<td>0.50</td>
<td>0.25</td>
</tr>
<tr>
<td>Between SACCF and Sbdy</td>
<td>2.87</td>
<td>1.45</td>
<td>2.48</td>
</tr>
<tr>
<td>Between Sbdy and 69° S</td>
<td>4.21</td>
<td>3.65</td>
<td>3.96</td>
</tr>
<tr>
<td>Between 69° S and the ice shelf</td>
<td>6.01</td>
<td>3.28</td>
<td>4.30</td>
</tr>
</tbody>
</table>

Combined north westward and south eastward legs of SANA 50

There is a high (>1.5) SiO$_4$:NO$_3$ ratio throughout this region except around South Georgia (between the APF and SACCF) where the average SiO$_4$:NO$_3$ ratio for this region is 0.25 and 0.32 for the north westward and south eastward legs respectively.

The difference between the early north westward leg and the later south eastward leg’s SiO$_4$ and NO$_3$ is positive in all regions expect around South Georgia (-1.45 µmol l$^{-1}$ and -0.76 µmol l$^{-1}$ for SiO$_4$ and NO$_3$ respectively).

Combined northward and southward legs of SANA 50

There is a high (>1.5) SiO$_4$:NO$_3$ ratio between South Africa and the northern STF as well as south of the SACCF. Regions between the northern STF and APF have low (<1.5)
SiO$_4$:NO$_3$ ratio. In the region between the APF and SACC, the ratio changed from a high (1.55) to a low (0.50) ratio between the southward and northward legs.

The difference between the early southward leg and the later northward leg’s SiO$_4$ and NO$_3$ is positive for all regions, except between South Africa and the northern STF for SiO$_4$ (-0.75 µmol l$^{-1}$), and south of 69º S for NO$_3$ (-25.01 µmol l$^{-1}$).

3.1.6) Statistical analyses

3.1.6.1) Linear regression

To determine whether various factors that might influence $F_v/F_m$ and $\delta_{\text{PSII}}$ values were significant, linear regressions were performed for a number of variables from the SANAE 49 cruise only, since this is the most complete data set. Two alternative linear regressions were run. The first (tests a and b) included the dominant phytoplankton groups that on average made up more than 5% of the population. The second (tests c and d) used the Fux:Hex ratio.

Test a)

\[
Y(F_v/F_m) = 3.385e+00 \pm (-1.175e-03)\text{SiO}_4 + (-9.648e-03)\text{PO}_4 + (6.800e-03)\text{NO}_3 + (4.049e-02)\text{chlorophyll} + (-6.515e-03)\text{temperature} + (-7.500e-02)\text{salinity} + (-6.959e-02)\text{oxygen} + (-2.280e-02)\text{fluorescence} + (-2.604e-04)\text{PAR} + (-8.572e-04)\text{diatoms} + (-2.174e-03)\text{haptophytes} + (-2.051e-03)\text{chromophytes}
\]

$P$ value $<0.001$ (chlorophyll, PAR)

$P$ value $<0.01$ (SiO$_4$, oxygen, fluorescence)

$P$ value $<0.05$ (haptophytes)
Test b)

\[ Y(\Delta_{PSII}) = -2.001e+03 + (1.061e-01)SiO_4 + (2.089e+00)PO_4 + (-4.732e+00)N_3 + \\
(-1.858e+01)\text{chlorophyll} + (-2.241e+00)\text{temperature} + (7.177e+01)\text{salinity} + \\
(-2.120e+01)\text{oxygen} + (2.013e+01)\text{fluorescence} + (1.549e-02)\text{PAR} + \\
(1.176e+00)\text{diatoms} + (-8.329e-01)\text{haptophytes} + (1.567e+00)\text{Chromophytes} \]

\( P \text{ value <0.001 (fluorescence)} \)

\( P \text{ value <0.01 (chlorophyll, salinity)} \)

\( P \text{ value <0.05(NO}_3, \text{ diatoms, chromophyte)} \)

Test c)

\[ Y(F_v/F_m) = 5.020e+00 + (-1.041e-03)SiO_4 + (-6.409e-03)PO_4 + (8.141e-03)N_3 + \\
(3.330e-02)\text{chlorophyll} + (1.524e-03)\text{temperature} + (-1.230e-01)\text{salinity} + \\
(-9.384e-02)\text{oxygen} + (-2.005e-02)\text{fluorescence} + (2.615e-04)\text{PAR} + \\
(7.083e-03)\text{Fuc:Hex} \]

\( P \text{ value <0.0001 (Fuc:Hex)} \)

\( P \text{ value <0.001 (oxygen, PAR)} \)

\( P \text{ value <0.01 (chlorophyll, salinity)} \)

\( P \text{ value <0.05(SiO}_4, \text{ NO}_3, \text{ fluorescence)} \)

Test d)

\[ Y(\Delta_{PSII}) = -1.809e+03 + (-2.255e-01)SiO_4 + (9.016e+00)PO_4 + (-1.999e-01)N_3 + \\
(-1.804e+01)\text{chlorophyll} + (5.458e+00)\text{temperature} + (5.935e+01)\text{salinity} + \\
(6.992e-02)\text{oxygen} + (1.740e+01)\text{fluorescence} + (6.170e-02)\text{PAR} + \\
(3.266e+00)\text{Fuc:Hex} \]

\( P \text{ value <0.01 (fluorescence, Fuc:Hex)} \)

\( P \text{ value <0.05(chlorophyll, salinity)} \)

3.1.6.2) Specific parameter comparison

Specific parameter comparisons with F_v/F_m show a positive relationship with F_v/F_m and NO_3, PO_4, % diatoms, SiO_4:NO_3, while showing a negative relationship between F_v/F_m
and SiO$_4$, f-ratio (Figure 23). However, in this analysis, due to the complexity of the system, these trends have a very low gradient ($R^2$ ranges between 0.081 and 0.2055).

**Figure 23:** Graphs showing SANAЕ 49 F$_V$/F$_{m}$ in relation to individual parameters namely: NO$_3$ (a), PO$_4$ (b), SiO$_4$ (c), % Diatoms (d), Fuc:hex ratio (e), f-ratio (f) and SiO$_4$:NO$_3$ (g).
SECTION 2: Bio-Assay Experiment Results

Due to logistical reasons each experiment was started on a different day, thus experiencing a range of conditions, reflecting both spatial and temporal variability within the Southern Ocean (table 4).

3.2.1) Initial conditions of bio-assay experiments

Table 4: The position and associated water mass for each on board Fe-light incubation experiment on SANAE 50. The positions for the 6 experiments are marked on figure 5.

<table>
<thead>
<tr>
<th>Exp. Sampling Date</th>
<th>Latitude</th>
<th>Longitude</th>
<th>SST (°C)</th>
<th>Nitrate (μM)</th>
<th>Silicate (μM)</th>
<th>Phosphate (μM)</th>
<th>SiO₄:NO₃ for region</th>
<th>Chl-a (μg.l⁻¹)</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>A 04-Jan</td>
<td>-64.021</td>
<td>-15.3282</td>
<td>-0.82</td>
<td>24.49</td>
<td>80.87</td>
<td>1.44</td>
<td>3.96 ± 0.58 ± 0.014</td>
<td>North-Eastern leg, South of the southern boundary, still icebergs and growlers around</td>
<td></td>
</tr>
<tr>
<td>B 15-Jan</td>
<td>-62.276</td>
<td>-21.8786</td>
<td>0.55</td>
<td>15.77</td>
<td>103.00</td>
<td>1.35</td>
<td>3.96 ± 0.56 ± 0.099</td>
<td>South-Western leg, South of the southern boundary</td>
<td></td>
</tr>
<tr>
<td>C 19-Jan</td>
<td>-68.911</td>
<td>-11.2692</td>
<td>0.30</td>
<td>22.02</td>
<td>92.66</td>
<td>1.57</td>
<td>4.40 ± 1 ± 0.000</td>
<td>South-Western leg, icebergs, South of the Southern Boundary</td>
<td></td>
</tr>
<tr>
<td>D 26-Jan</td>
<td>-70.345</td>
<td>-7.8767</td>
<td>0.28</td>
<td>17.17</td>
<td>81.69</td>
<td>1.35</td>
<td>4.52 ± 1.93 ± 0.077</td>
<td>South of the pole circle, marginal ice zone. High productive area with a lot of top predators e.g. killer whales, minke whales, humpback whales and leopard seals</td>
<td></td>
</tr>
<tr>
<td>E 09-Feb</td>
<td>-55.962</td>
<td>0.0203</td>
<td>0.88</td>
<td>24.67</td>
<td>90.36</td>
<td>1.81</td>
<td>1.45 ± 0.77 ± 0.026</td>
<td>Northward leg, between SBdy and SACCF, water blue, low productivity, miserable, cloudy weather. 1-3 m swell. Weather cleared a bit during the day. Whales and albatrosses sighted</td>
<td></td>
</tr>
<tr>
<td>F 11-Feb</td>
<td>-50.335</td>
<td>1.0675</td>
<td>6.17</td>
<td>19.00</td>
<td>0.29</td>
<td>1.02</td>
<td>0.12 ± 1.28 ± 0.042</td>
<td>Northward leg, in the APZ, after the Polar Front. Highly productive warm waters</td>
<td></td>
</tr>
</tbody>
</table>

3.2.2) Physical changes experienced by Fe incubations

Table 5: Average (graphically depicted) sea temperature and PAR (ambient, high and low light bottles) for each Fe-light experiment on SANAE 50. Columns from left to right show: average, standard deviation and range. The logging of the data is explained in the methods section (page 41).

<table>
<thead>
<tr>
<th>Experiment</th>
<th>average sea temperature (°C)</th>
<th>average par (W/m²)</th>
<th>average par for high light (W/m²)</th>
<th>average par for low light (W/m²)</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>3 (+/-1.6)</td>
<td>319 (+/-534)</td>
<td>224 (+/-267)</td>
<td>112 (+/-133)</td>
</tr>
<tr>
<td>B</td>
<td>0.5 (+/-0.1)</td>
<td>264 (+/-241)</td>
<td>132 (+/-120)</td>
<td>66 (+/-60)</td>
</tr>
<tr>
<td>C</td>
<td>0.1 (+/-0.5)</td>
<td>1001 (+/-885)</td>
<td>501 (+/-442)</td>
<td>250 (+/-221)</td>
</tr>
<tr>
<td>D</td>
<td>0.1 (+/-0.4)</td>
<td>971 (+/-783)</td>
<td>486 (+/-391)</td>
<td>243 (+/-196)</td>
</tr>
<tr>
<td>E</td>
<td>3.5 (+/-3.3)</td>
<td>173 (+/-578)</td>
<td>239 (+/-339)</td>
<td>119 (+/-169)</td>
</tr>
<tr>
<td>F</td>
<td>30.1 (+/-1.8)</td>
<td>350 (+/-772)</td>
<td>280 (+/-386)</td>
<td>140 (+/-193)</td>
</tr>
</tbody>
</table>

3.2.2.1) Sea temperature changes

Experiments E (7.5 ± 3.3 °C) and F (10.1 ± 1.8 °C) were characterised by the greatest average sea temperatures as well as the greatest ranges in temperature (1.3 - 13 °C and 5.9 -
13.2 °C respectively) the other four experiments had very low sea temperatures and ranges of: -1.4 °C to 4.5 °C for experiment A, -1 °C to 0 °C for experiment B, -1.5 °C to 1.6 °C for experiment C and -1.2 °C to 0.8 °C for experiment D (table 5).

3.2.2.2) PAR readings

Average PAR for ‘high’ and ‘low’ light incubations represented 50% and 25% of ambient irradiance respectively.

Experiment C experienced the highest average PAR of 1001 ± 885 µE.m\(^{-2}\).s\(^{-1}\) (table 5). Experiment D experienced 971 ± 783 µE.m\(^{-2}\).s\(^{-1}\); experiment A 449 ± 534 µE.m\(^{-2}\).s\(^{-1}\); experiment E 478 ± 678 µE.m\(^{-2}\).s\(^{-1}\) and F (560 ± 772 µE.m\(^{-2}\).s\(^{-1}\). Experiment B experienced notably lower irradiances 264 ± 772 µE.m\(^{-2}\).s\(^{-1}\) than any other experiment.

3.2.2.3) Summary of weather observations

As is clear from the PAR values, experiment A was overcast throughout the experiment except for the evening of the second day until 6am on the third day.

Experiment B experienced mostly overcast weather, except for 2 hours at 20:00 on the second day, and full sunshine on the last day.

Experiment C experienced mostly full sunshine except for day 1, which was overcast, with heavy clouds, rain, and snow.
Experiment D started off sunny, becoming overcast by 8am on day 1, but the dull weather cleared by the end of day 2, only to return in the latter half of day 3 with snow.

Experiment E and F remained overcast except for an hour or two of patchy sunshine every day.

3.2.3) **Experiment results**

3.2.3.1) *Variation in physiological response time*

Experiment A showed a change in the photo-physiological response from day 2 and 3 for $\varphi_{\text{PSII}}$ and $F_v/F_m$ respectively (figure 24). Experiment B responded quickly from day 1 and also showed the greatest change. Photo-physiological responses for Experiments C, E and F also started on day 1, with experiment D starting at days 0 and 1 for $F_v/F_m$ and $\varphi_{\text{PSII}}$ respectively.

3.2.3.2) *Variation in ranges*

Experiment B shows the greatest response to Fe alleviation (figure 24). This is evident especially through the $F_v/F_m$ response. Experiment B began at a $F_v/F_m$ value of $0.27 \pm 0.000$. By the end of day 5, the Fe addition bottles reached $0.47 \pm 0.014$ and $0.47 \pm 0.016$ for 50% and 25% light respectively, while the control bottles for 50% and 25% light were at $0.29 \pm 0.049$ and $0.28 \pm 0.031$ respectively.
Figure 24: Averaged $F_v/F_m$ (left), $\sigma_{PSII}$ (middle) and chl-a (right) results for Fe incubation experiments on SANAE 50. Red represents the Fe addition bottles. Blue represents the control bottles. A triangle and square represents 50% and 25% light respectively. Circles symbolise the overall control.
Despite huge variation in starting photosynthetic parameters, by day 5 (excluding experiments E and F, which didn’t reach day 5) each experimental Fe addition bottles reached values ~0.4 and ~200 for $F_v/F_m$ and $\phi_{PSII}$ respectively, while the control bottles were below these values for $F_v/F_m$ and above for $\phi_{PSII}$.

3.2.3.3) FIRE

A paired-samples t-test was conducted to separately compare the $F_v/F_m$ and $\phi_{PSII}$ values under the following conditions:
1) All the control bottles + the Fe addition bottles
2) The control bottles + the Fe addition bottles at 50% irradiance
3) The control bottles + the Fe addition bottles at 25% irradiance
4) The control bottles at 50% irradiance and 25% light irradiance
5) The Fe addition bottles at 50% irradiance and 25% irradiance
6) The contamination control bottle and the control bottle at 50% irradiance (Experiments B, C and D only)

Summary: FRRf responses to Fe additions

All experiments showed an increase in $F_v/F_m$ and a decrease in $\phi_{PSII}$ for bottles with Fe additions (figure 24). The degree to which this occurred was highly variable.

Experiment B showed the greatest statistically significant response (table 6) to Fe alleviation with both $F_v/F_m$ and $\phi_{PSII}$ responding significantly under both high light (50%) and low light (25%) conditions ($P < 0.001$ for all). Experiment C showed a full response at high irradiance ($P = 0.0031$, and $P = 0.0005$ for $F_v/F_m$ and $\phi_{PSII}$ respectively), while experiments D
(P <0.001 for $F_v/F_m$ and $\delta_{PSII}$) and F (P =0.0288; P =0.0148 for $F_v/F_m$ and $\delta_{PSII}$ respectively) showed statistically significant responses at low irradiances.

Table 6: The $F_v/F_m$, $\delta_{PSII}$ and chl-a results for all the Fe-light incubation experiments. X is used to show no significant difference

<table>
<thead>
<tr>
<th>experiments</th>
<th>response measured</th>
<th>All the control bottles and all the Fe addition bottles</th>
<th>The control bottles and the Fe addition bottles at 50 % light levels</th>
<th>The control bottles and the Fe addition bottles at 25 % light levels</th>
<th>The control bottles at 50 % light levels and 25% light levels</th>
<th>The Fe addition bottles at 50 % light levels and 25% light levels</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>$F_v/F_m$</td>
<td>Significant difference $(p=0.0092)$</td>
<td>x</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>A</td>
<td>$\delta_{PSII}$</td>
<td>Significant difference $(p=0.0064)$</td>
<td>x</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>A</td>
<td>Chlorophyll-a</td>
<td>Significant difference $(p=0.0003)$</td>
<td></td>
<td>x</td>
<td>x</td>
<td></td>
</tr>
<tr>
<td>B</td>
<td>$F_v/F_m$</td>
<td>Significant difference $(p=0.0000001)$</td>
<td>x</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>B</td>
<td>$\delta_{PSII}$</td>
<td>Significant difference $(p=0.0000000017)$</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>B</td>
<td>Chlorophyll-a</td>
<td>Significant difference $(p=0.0003)$</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>C</td>
<td>$F_v/F_m$</td>
<td>Significant difference $(p=0.00001)$</td>
<td>x</td>
<td>x</td>
<td></td>
<td></td>
</tr>
<tr>
<td>C</td>
<td>$\delta_{PSII}$</td>
<td>Significant difference $(p=0.000001)$</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>C</td>
<td>Chlorophyll-a</td>
<td>Significant difference $(p=0.0003)$</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>D</td>
<td>$F_v/F_m$</td>
<td>Significant difference $(p=0.000000003)$</td>
<td>x</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>D</td>
<td>$\delta_{PSII}$</td>
<td>Significant difference $(p=0.000000005)$</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>D</td>
<td>Chlorophyll-a</td>
<td>Significant difference $(p=0.000000019)$</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>E</td>
<td>$F_v/F_m$</td>
<td>Significant difference $(p=0.0000000019)$</td>
<td>x</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>E</td>
<td>$\delta_{PSII}$</td>
<td>Significant difference $(p=0.00000000017)$</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>E</td>
<td>Chlorophyll-a</td>
<td>Significant difference $(p=0.0003)$</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>F</td>
<td>$F_v/F_m$</td>
<td>Significant difference $(p=0.00000000017)$</td>
<td>x</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>F</td>
<td>$\delta_{PSII}$</td>
<td>Significant difference $(p=0.000000000017)$</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>F</td>
<td>Chlorophyll-a</td>
<td>Significant difference $(p=0.0003)$</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Experiment C shows no significant statistical photo-physical response to Fe addition in low light levels; however a significant statistical photo-physical response was seen in high light conditions. Experiment F shows no significant statistical photo-physical response to Fe addition in high light levels; however a significant statistical photo-physical response was seen in low light conditions. In experiments A, D, E (high light) a significant difference between Fe supplemented bottles and controls without Fe was only shown by $\delta_{PSII}$ (P =0.0003, P =0.0025, P =0.0022 for A, D and E respectively). Under low light in experiment
A, \( F_v/F_m \) responded significantly (\( P =0.0105 \)) to Fe addition, while in experiment E, \( \phi_{PSII} \) responded significantly (\( P =0.0074 \)) under low light to Fe alleviation.

Experiment D was the only experiment where the controls between light levels differed.

3.2.3.4) Chlorophyll-a

Similarly to the Fire data (section 3.1.1), a paired-samples t-test was conducted to separately compare chl-a under the same conditions.

Although there was an observable increase seen in chl-a concentrations in response to Fe additions (figure 24) in all experiments this was only significantly different from the controls in experiments D (\( P =0.041 \); 25% light), E (\( P =0.0175 \); 50% light) and F (\( P =0.0092 \); 50% light) (see table 6).

3.2.3.5) Nutrient concentrations

Unfortunately, the experimental nutrients values were incomprehensible and unrealistic (see appendix I for example), almost certainly due to poor on-board analyses combined with poor storage options (See appendix J for critique on these methods).
CHAPTER 4: DISCUSSION

Although the Southern Ocean is a HNLC ecosystem (Levitus et al., 1993; Sullivan et al., 1993; Moore and Abbott, 2000; Boyd et al., 2002; Seeyave et al., 2007), intense and sporadic chl-a blooms do occur (Arrigo et al., 2008) and their presence is controlled by the ocean's physical and chemical environment. The open ocean north of the ice zone, away from fronts and shallow topography tends to have low primary production that has been attributed to Fe limitation (Sunda and Huntsman, 1997; Boyd et al., 2000; Sokolov and Rintoul, 2007). However blooms are known to form around frontal zones (Laubscher et al., 1993; Joubert et al., 2011), sub-Antarctic islands (Blain et al., 2001; Seeyave et al., 2007) and in the marginal ice zone (Boyd, 2002; Dierssen et al., 2002; Lannuzel et al., 2006).

Given the important contribution of these phytoplankton blooms to the Southern Ocean biological pump (Longhurst, 1991; Schlitzer, 2002), it is important for us to observe and characterise variability in both the physical mechanisms controlling primary production and the biological response to these forcing mechanisms.

The following section describes the physical and chemical environment encountered in the South Atlantic sector of the Southern Ocean in the austral summers of 2009 (SANAE 49) and 2010 (SANAE 50) together with the biological response in chlorophyll. In addition it makes inter-annual comparisons between the two years describing notable inter-annual variability in the physical and chemical environment as well as the biological response.
I: Physical, chemical and biological environment

4.1.1) Hydrography

Phytoplankton distribution, diversity, biomass and production depend on the physiological responses of phytoplankton to the often extreme conditions under which they live. The hydrography of the Southern Ocean is a key determinate of phytoplankton distribution via its effects on the light environment (through changes in the MLD) and the nutrient supply (through adjustments in stratification and mixing).

4.1.1.1) Hydrography between Antarctica and South Georgia

WW, the relatively homogeneous, subsurface temperature minimum layer (Park et al., 1998), is evident on both cruises. This water originates from cooling in the previous winters mixed layer and is capped by seasonal warming and ice melt in the summer. UCDW and Lower Circumpolar Deep Water (LCDW) were present (below ~100m north of SACC) in both years (figure 7 and 8). UCDW originates from the Indian and Pacific basin and is characterised by relatively warm, highly saline waters that are nutrient rich but low in oxygen (Callahan, 1972; Orsi et al., 1995; Veth et al., 1997; Park et al., 1998). Conversely, LCDW (below ~200m south of the SBdy) is nutrient poor and denser (figure 7 and 8); hence its position below UCDW.

4.1.1.2) Hydrography from Antarctica to Cape Town

The very distinctive temperature increase seen just south of South Africa is caused by the Agulhas Current, originating in the Indian Ocean. The warm (≥18ºC), salty (≥35.5 PSU), subtropical surface water of the Agulhas Current interacts with and warms the South Atlantic
through both anticyclonic ring (averaged 50 km wide, 50 m deep) shedding from the Agulhas Retroflection Region and warm filaments of Agulhas Current Water (Gordon et al., 1987; Lutjeharms and Cooper, 1996).

Temperatures decrease significantly as one moves south towards Antarctica. The deeper, colder, sinking WW can be seen in both years south of 50º S (figure 9).

4.1.1.3) Inter-annual variability in hydrography

Although frontal positions remained consistent within the property ranges given by Orsi et al., (1995), some variability in position was observed between years. The SACCF moved from ~58º S to ~50º S from SANAE 49 to SANAE 50. The SBdy showed a slight northward movement between years moving from between ~59º S and ~60º S to ~57º S. On the northward leg, between the summers of 2009 (SANAE 49) and 2010 (SANAE 50) the STF and APF moved ~1º north, the SAF moved ~1º south, while the SACCF and SBdy moved <1º south and north respectively. As frontal regions are areas associated with increased chl-a (due to the upwelling of deep nutrient rich waters caused by diverging surface waters), slight movement of these fronts influences the distribution of chl-a blooms.

Changes in the MLD play a particularly important role in the Southern Ocean’s biology due to its influence on the light environment that phytoplankton are exposed to. A deep mixed layer relative to the euphotic zone means that phytoplankton will be mixed outside of a favourable light environment, therefore negatively impacting photosynthesis. MLD’s were deeper on the south eastward legs compared to the north westward legs of both SANAE 49 and 50. This is specifically noticeable south of the SBdy (figures 7 and 8).
Deeper MLD’s later in the summer season are counter-intuitive given that MLD’s are expected to shoal with seasonal warming. As such the deeper MLD’s south of the SBdy on the south eastward legs are likely the result of localised wind events deepening the mixed layer.

4.1.1.4) Inter-annual variability in sea ice

As Southern Ocean blooms are tied to the timing between seasonal ice melt and available sunlight (Moore and Abbott, 2000), an overall comparison between the two years is useful. Melting ice is known to release Fe into surface waters (Dierssen et al., 2002), additionally the increase in buoyancy from the input of fresh melt water leads to increased stratification in the water column, allowing the phytoplankton to experience favourable light conditions. These combined features allow both sufficient Fe and light needed for the formation of a phytoplankton bloom event. Sea surface conditions between the two consecutive summers were remarkably different, notably in terms of seasonal ice coverage. In 2009, extensive and thick ice was evident, while in 2010, sea-ice cover was minimal (table 1; NSIDC, 2012). This highlights inter-annual-variability, as frequently noted for example by Murphy et al., (1998) and Reid and Croxall (2001).

A consequence of variable sea ice cover is that the warm surface waters that were observed stretching southwards from South Georgia extended further south in 2010 (SANAE 50, figure 8) than they did in 2009 (SANAE 49, figure 7).
4.1.2) **Nutrients**

Nutrient concentrations are known to limit the growth of phytoplankton. This affects both the size (Sunda and Huntsman, 1995) and community structure (de Baar and Boyd, 2000; Smetacek *et al.*, 2004; Seeyave *et al.*, 2007) of the phytoplankton community. The most important limiting nutrients are Fe, NO$_3$ and SiO$_4$ (Boyd, 2002; Arrigo *et al.*, 2008).

Fe is known to be limiting within the Southern Ocean, specifically in open ocean conditions north of the sea ice zone, away from fronts and continental land masses (Boyd *et al.*, 2000; Sokolov and Rintoul, 2007). As the Fe data (responsibility of the CSIR and the University of Stellenbosch) remain unavailable at the time of writing, secondary indicators of Fe availability have been used.

Although all Southern Ocean waters are known to have high concentrations of NO$_3$ and PO$_4$, concentrations of SiO$_4$ are known to differ markedly from north to south. Subantarctic waters north of the Antarctic Polar Frontal (APF) have low SiO$_4$ concentrations (1 to 5 µmol l$^{-1}$), whereas high SiO$_4$ concentrations (> 60 µmol l$^{-1}$) are found south of the APF (Coale *et al.*, 2004).

4.1.2.1) **Nutrients between Antarctica and South Georgia**

Neither NO$_3$ (>10 µmol l$^{-1}$) nor SiO$_4$ (>15 µmol l$^{-1}$) concentrations were limiting between Antarctica and South Georgia on either SANAE 49 or SANAE 50.

Both NO$_3$ and SiO$_4$ show an increase in concentrations with depth, and a decrease in concentration in the surface waters around South Georgia. This indicates the use of these
nutrients (by phytoplankton) in the surface waters, specifically around South Georgia (figures 10, 11, 12 and 13).

In increase in NO$_3$ concentrations is noted in the deeper waters between 62° S and 66° S on the south eastward leg of SANAE 49 (figure 10).

4.1.2.2) Nutrients from Antarctica to Cape Town

On both Antarctica to Cape Town transects neither NO$_3$ (>1 µmol l$^{-1}$) nor SiO$_4$ (>2 µmol l$^{-1}$) were limiting south of ~50° S (figure 10, 11, 12 and 13), however north of the APF (found at 50.08° S and 49.36° S on SANAE 49 and SANAE 50 respectively), SiO$_4$ concentrations dropped to below <2 µmol l$^{-1}$ a concentration that is often considered limiting for diatom growth (Ragueneau et al., 2000; Peterson et al., 2005; Whitney et al., 2005). North of the northern STF at Northern STF ~39° S NO$_3$ concentrations fell to <1 µmol l$^{-1}$, limiting phytoplankton growth in these subtropical waters (e.g. Joubert et al., 2011).

4.1.2.3) Inter-annual variability in nutrients

Although, the relative geographic distribution of nutrients was similar for both SANAE 49 and 50, there were inter-annual differences between minima and maxima. On SANAE 49, the SiO$_4$ minimum was 0.09 µmol l$^{-1}$ lower than on SANAE 50, whereas the NO$_3$ minimum was 0.36 µmol l$^{-1}$ higher on SANAE 49. On SANAE 49, the SiO$_4$ maximum was 52.15 µmol l$^{-1}$ lower than on SANAE 50, whereas the NO$_3$ maximum was 2.44 µmol l$^{-1}$ higher on SANAE 49 (figures 10, 11, 12 and 13). Differences in nutrient concentrations can
be accounted for by natural variability and by differences in biological uptake and physical replenishment.

4.1.3) Chlorophyll-a

Chl-a is used as a proxy for phytoplankton biomass. As a complex interplay between nutrient availability and light controls the formation of chl-a blooms in the Southern Ocean (Raven, 1990; Sunda and Huntsman, 1997; Boyd et al., 1999; Boyd, 2002; Hassler et al., 2012), interpreting phytoplankton distribution in relation to the physical and chemical environment is key to understanding the mechanisms leading to phytoplankton adaptations and distributions in the oceans. This understanding becomes an imperative, as climate projections predict changes in the MLD and stratification, both of which influence nutrient and light availability (Greenblatt and Sarmiento, 2004).

4.1.3.1) Chlorophyll-a between Antarctica and South Georgia

There were two main areas containing chl-a blooms (here described as chl-a>1 µg.l$^{-1}$) between Antarctica and South Georgia (figure 14 and 15). These blooms occurred around South Georgia as well as between ~67° S and ~61° S. A third bloom was present in SANAE 50 at ~70° S (figure 15).

Both years saw an increase (reaching a maximum of 2.5 - 3 µg.l$^{-1}$ and 3.8 µg.l$^{-1}$ for 2009 and 2010 respectively) in chl-a associated with South Georgia (figure 14 and 15) which is likely due to natural Fe fertilization from shallow sediments associated with sub Antarctic islands (De Baar et al., 1995; Blain et al., 2001; Holeton et al., 2005; Seeyave et al., 2007) as
well as increased stratification due to increase buoyancy from fresh-water runoff (Joubert et al., 2011).

The blooms occurring between ~67° S and ~61° S are in the open ocean, largely away from divergent fronts. However in this region of marginal ice zone melting, which extends to ~55° S (at 10° E) in winter (NSIDC, 2012) is known to increase Fe in surface waters and improve light conditions through increased buoyancy (Joubert et al., 2011). This likely explains the presence of these blooms. In addition, during SANAE 49, it is noticeable that on the south eastward leg, the chl-a bloom at ~61° S (figure 14) corresponded with an increase in deeper NO₃ concentrations (figure 10). The deeper mixed layer on this return leg could account for mixing of this increased nutrient concentration in to the surface water, thus aiding this chl-a bloom.

4.1.3.2) Inter-annual variability in Chlorophyll-a

There were some substantial differences in the distribution and abundance of chl-a between 2009 (figure 14) and 2010 (figure 15). Although both years showed a general northward decrease in chl-a biomass between Antarctica and South Africa, this decrease differed slightly in both concentration (SANAE 49’s maximum was higher than SANAE 50 by 0.68 µg.l⁻¹) and distribution of chl-a maxima. The distribution of the high chl-a region varied from 65.8° S to 57.5° S and from 70.6° S to 59.3° S for 2009 and 2010 respectively (figures 14 and 15). On SANAE 50, the maximum was higher (3.65 µg.l⁻¹) and further south (70.6° S) compared to SANAE 49 (2.57 µg.l⁻¹ at 65.8° S). A likely explanation of the more southward peak in chl-a on SANAE 50 was the ice conditions. SANAE 50 saw a smaller area
of sea ice cover (table 1) than during SANAE 49, thus allowing the formation of a more southerly bloom.

The chl-a blooms associated with South Georgia on the two years were different in that chl-a maxima were higher in 2010 (3.8 ug l\(^{-1}\)) than in 2009 (2.5-3.0 ug l\(^{-1}\)) (figure 15 and 14 respectively). Other differences include the notable lack of a bloom event on SANAE 50’s north westward leg at 67º S and 62º S (figure 15). Chl-a blooms were evident at these positions on all three of the other transects (figures 14 and 15) namely SANAE 50’s south eastward leg, as well as both of SANAE 49’s legs to and from South Georgia. These differences could be due to a lack of a localised wind event on SANAE 50’s north westward leg. In the Southern Ocean one expects the phytoplankton to bloom in spring with the release from light limitation (Venables and Moore, 2010), with these blooms decreasing in intensity as nutrients become limiting over summer. The deeper MLD’s on the south eastward legs (of SANAE 49 and 50) compared to the north westward legs suggest increased wind events on the later south eastward legs that potentially mixed limiting nutrients into the surface waters accounting for the increase in Chl-a concentrations found later in the season.

### II: Phytoplankton photo-physiology

Variability in FRRf values as a measure of physiological competency has been attributed to differences in phytoplankton community structure (Suggett et al., 2004, 2009; Moore et al., 2005), as well as light and Fe co-limitation (Greene et al., 1991; Boyd and Abraham, 2001). Meanwhile macronutrient availability, temperature and salinity play lesser roles in regulating the FRRf response (Kolber et al., 1988; Boyd and Abraham, 2001).
A low \( F_v/F_m \) and high \( \sigma_{PSII} \) reveals physiological stress brought on by environmental parameters, while the inverse reveals a healthy, efficient phytoplankton population (Holeton et al., 2005; Suggett et al., 2006; Moore et al., 2007b). This section focuses on discussing the photo-physiological “health” of phytoplankton based on FRRf measurements during SANAE 49 and 50 and assesses the environmental controls potentially accounting for differences in the measured photo-physiological responses.

4.2.1.1) Observed phytoplankton photo-physiology (FRRf)

Corresponding with past literature (Holeton et al., 2005; Moore et al., 2007b), \( F_v/F_m \) and \( \sigma_{PSII} \) values were the converse of each other (figures 18 and 20) throughout the summer of 2009. Over the summer of 2010, \( F_v/F_m \) and \( \sigma_{PSII} \) values were largely the converse of each other (figures 19 and 21), with corresponding inverse values (of \( F_v/F_m \) and \( \sigma_{PSII} \)) north of 63° S (figure 19 b and 21 b), north of 40° S and around South Georgia (on the north westward leg) (figure 19 a and 21 a). During SANAE 49, there was however, one notable difference near South Georgia (on the south eastward leg), where the \( F_v/F_m \) values were high (reaching a maximum of \( \sim 0.45 \)), indicating physiologically healthy cells, but \( \sigma_{PSII} \) values were also unexpectedly high (253 ± 17.6 - 308 ± 73.6), rather than low as anticipated. This suggests that although the high \( F_v/F_m \) shows an increase in photosynthetic efficiency, part of the photosynthetic process is undergoing photosynthetic stress (high \( \sigma_{PSII} \)). Hence it is possible that the population in this region is undergoing change, either in photosynthetic health or in community structure. Considering the photosynthetic health, enough Fe could be allowing \( F_v/F_m \) to function efficiently, but with insufficient Fe for this to apply to \( \sigma_{PSII} \) as well. However, considering that both parameters are an indication of the functioning of PS II, this explanation is unlikely. Alternatively, the low light, storm conditions that occurred during
this section of the transect supports the notion that though some Fe alleviation occurred, the phytoplankton were still struggling to function efficiently under low light conditions. Considering community structure, the data (figure 17) shows an increase in diatom dominance (from ± 45% to >60%, figure 17 a) as well as an increase in the diatom to haptophyte ratio (showing a move towards diatom dominance) on the south eastward leg around South Georgia (figure 17 d). This dominance corresponds with that found by Gibberd et al., (2013), who studied the same area, the previous year. It is possible that the changing phytoplankton community was receiving enough Fe to increase its photosynthetic efficiency through $F_v/F_m$ and become a diatom dominated community, despite the still prevalent inefficiency in $\delta_{PSII}$.

The phytoplankton photo-physiology FRRf parameters of $F_v/F_m$ were remarkably similar between the austral summers of 2009 and 2010. High $F_v/F_m$ (reaching a maximum of >0.4 for both years) characterised the region around South Georgia and South of 60° S, although highest $F_v/F_m$ values were at depth in 2009, but on the surface in 2010. The moderate to low values (<0.2) in the remaining areas between Antarctica and South Georgia were also consistent for both years.

From these data, the areas of high photosynthetic efficiency are near South Georgia, in the deeper waters near Antarctica and to a lesser extent around the SBdy and SACCF. These are areas known for elevated Fe concentrations (Laubscher et al., 1993; Blain et al., 2001; Seeyave et al., 2007; Joubert et al., 2011) and as such, Fe alleviation is thought to be the main driver of the high, efficient FRRf values found here. Conversely, the areas between these high photosynthetic efficiency regions are typically low in Fe as they are distant from fronts and shallow topography and terrestrial sources (Sunda and Huntsman, 1997; Boyd et
such that Fe limitation is the likely cause of low photosynthetic efficiency values found here.

4.2.2) **Statistical evidence for controls on photo-physiology**

To identify the major controlling factors behind the FRRf values, two main statistical tests were carried out to relate the photosynthetic parameters ($F_v/F_m$ and $\sigma_{PSII}$) to environmental factors. These tests focused on the SANAE 49 (2009) cruise only due to the more complete nature of this data set, which included HPLC. The first test was a linear analysis that compared $F_v/F_m$ and $\sigma_{PSII}$ to a number of environmental factors (temperature, salinity, PAR, SiO$_4$, NO$_3$, oxygen) and phytoplankton indices (chl-a, fluorescence, community structure). The community structure was classified according to three main phytoplankton groups (diatoms, 49%; haptophytes, 29%; and chromophytes, 8%).

The following variables were all significantly related to values of $F_v/F_m$: chl-a and PAR ($P < 0.001$), SiO$_4$, oxygen and fluorescence ($P < 0.01$) and haptophytes ($P < 0.05$).

Whereas the following where all significantly related to $\sigma_{PSII}$: Fluorescence ($P < 0.001$), chl-a and salinity ($P < 0.01$), NO$_3$, diatoms and chromophytes ($P < 0.05$).

In the second test, the community structure data was replaced with the Fuc:Hex ratio (as used by: Barlow *et al.*, 1998; Smith and Asper, 2001; Hirata *et al.*, 2008; Feng *et al.*, 2010; Alderkamp *et al.*, 2012). Fuc:Hex ratio is used to determine the relative abundance of diatoms to *P. antarctica* within Antarctic phytoplankton populations (Alderkamp *et al.*, 2012). If the Fuc:Hex ratio increases, there are more diatoms relative to *P. antarctica* and
vice versa. The main known cause of a shift towards a diatom-dominated community is Fe alleviation (Boyd et al., 2000). Within Fe fertilised areas diatom growth can be further increased by increased light and SiO$_4$ concentrations (Boyd, 2002), as well as increased CO$_2$ concentrations (Tortell et al., 2008). Increased stratification of surface waters is also known to favour an increase in the Fuc:Hex ratio (Arrigo et al., 1999). Furthermore, an increase in diatoms relative to *P. antarctica* implies major changes in the heterotrophic community. This likely results in an increase in downward carbon fluxes due to an increase in the efficiency of the herbivorous food web, thus enhancing the export of CO$_2$ into the deeper ocean (Fonda Umani et al., 2002, 2005).

The community structure in this instance now gains the most significant influence; with $F_v/F_m$ now significantly related to Fuc:Hex ($P <0.0001$), oxygen and PAR ($P <0.001$), chl-a and salinity ($P <0.01$), SiO$_4$, NO$_3$ and fluorescence ($P <0.05$).

The statistical relationship with $\delta_{\text{PSII}}$ and community structure also improved as follows: Fuc:Hex and fluorescence ($P <0.01$), chl-a and salinity ($P <0.05$).

Though strongly significant relationships can be drawn between $F_v/F_m$ and $\delta_{\text{PSII}}$ in the linear regressions (section 3.1.6.2), the complexity of the South Ocean system, in relation to photosynthetic efficiency, is highlighted by Figure 23 where individual parameter comparison is inconclusive.

The results of the linear regression statistical tests are elaborated on in the following sections, explaining the role of the various environmental and biological factors influencing the statistical relationships between phytoplankton photo-physiology and the environment.
4.2.2.1) Fe availability as a driver of photo-physiology

Past literature shows that the most influential factors behind phytoplankton’s photosynthetic efficiency are known to be Fe and taxonomic community structure (Suggett et al., 2009). Unfortunately, the Fe data for this investigation is not available at this time, hence the interpretations presented here between photo-physiology and Fe are speculative, but based on published observations.

Indirect indicators for the presence of Fe in the HNLC waters of the Southern Ocean include: phytoplankton community structure and NO\textsubscript{3} uptake. Haptophyte dominance is found in areas of limiting Fe, while diatom dominated communities are found in regions of sufficient Fe (Sakshaug et al., 1991; Laubscher et al., 1993; de Baar et al., 1995; Smetacek et al., 1997; Boyd, 2002; Gibberd et al., 2013). In addition, f-ratios typically increase with increasing Fe availability (Lucas et al., 2007) due to the high Fe demand of NO\textsubscript{3} reduction (Lucas, 2009). Conversely, where Fe is limited, NO\textsubscript{3} uptake and its intracellular reduction to NH\textsubscript{4} via the Fe dependent enzymes nitrate and nitrite reductase is compromised (Lehninger, 1975), resulting in low f-ratios (<0.2) and low specific uptake rates (VNO\textsubscript{3}) despite high ambient NO\textsubscript{3} concentrations (Lucas et al., 2007).

NO\textsubscript{3} uptake in the Southern Ocean may also be inferred by changes in the SiO\textsubscript{4}:NO\textsubscript{3} ratios. If the SiO\textsubscript{4}:NO\textsubscript{3} ratio becomes high (>1.5), there is little SiO\textsubscript{4} removed relative to NO\textsubscript{3}. This occurs in Fe limiting regions. On the other hand, increased uptake rates of SiO\textsubscript{4} to an equal uptake rate of SiO\textsubscript{4} and NO\textsubscript{3} in the presence of sufficient Fe can shift the ambient SiO\textsubscript{4}:NO\textsubscript{3} ratios towards a low ratio of 1.1. Although not conclusive, increased concentrations of NO\textsubscript{3} and SiO\textsubscript{4} in surface waters can imply upwelling of deeper nutrient rich
waters with similarly increased Fe concentrations. Such combined upwelling of Fe and other macronutrients has been found off Peru (Bruland et al., 2005).

Following this argument, on the south eastward leg from 58° S to 54° S (figure 16) high f-ratios (up to 0.22) suggests the presence of Fe associated with South Georgia. Similarly, Fe limitation is thought to account for the low f-ratios (<0.06) found in the middle of the ocean away from known Fe sources (Boyd et al., 2000; Sokolov and Rintoul, 2007). On the north westward leg from 66° S to 62° S high f-ratios (reaching 0.37) similarly reveal a likely Fe source in surface waters off Antarctica. Fe enrichment in these waters is known to occur through upwelling of Fe rich water along the continental margin (Lannuzel et al., 2006) that is enriched by glacial scouring of underlying rocks (Dierssen et al., 2002) and by accumulation from atmospheric deposition of dust that is released into the surface ocean during ice melt (Dierssen et al., 2002; Lannuzel et al., 2006). F-ratios decreased with depth, most notably below 30m on the north westward leg, nearing South Georgia (figure 16). This finding of low f-ratios at depth can be explained by light limitation rather than Fe limitation, as NO$_3$ uptake is energy expensive and has a high light demand (as seen by the Michaelis-Menten equation [MacIsaac and Dugdale, 1972; Kudela et al., 1997]). Rather than low Fe concentrations, low f-ratios found near South Georgia likely reveal phytoplankton’s preference for reduced nitrogen in the form of NH$_4$ (Eppley and Peterso, 1979; Lucas et al., 2007; Joubert et al., 2011), which is often abundant in surface waters of subantarctic archipelago’s due to high annual rainfall transporting terrestrial sources offshore (Ismail, 1990).

As f-ratios are dependent on ambient NO$_3$ data, it should be remembered here that the SANAЕ 49 NO$_3$ data underwent a correction. However, table 2 reveal that a 10% change in
ambient NO\textsubscript{3} leads to a 5-9% change in f-ratio. This shows a low sensitivity to variation in NO\textsubscript{3}.

Increases in NO\textsubscript{3} concentrations (figure 10) corresponded with increases in F\textsubscript{v}/F\textsubscript{m} (figure 18), specifically near South Georgia, implying that upwelled waters are enriched by both NO\textsubscript{3} and Fe (Hiscock, 2004), where elevated F\textsubscript{v}/F\textsubscript{m} values are likely due to enhanced Fe availability (Suggett et al., 2009). Similar results were found in a study by Fung et al. (2000) and Pollard et al., (2009) where Fe was made available to phytoplankton around Crozet, from a shallow sedimentary source and from island runoff, resulting in high values of F\textsubscript{v}/F\textsubscript{m}, characteristic of Fe-replete cells (Suggett et al., 2009). This conclusion is statistically supported by the significant positive correlation between NO\textsubscript{3} and F\textsubscript{v}/F\textsubscript{m} in test c (methods section 3.1.5) (P <0.05) and the significant negative correlation between NO\textsubscript{3} and \(\delta_{\text{PSII}}\) in tests b (P <0.05). These observations suggest a benthic or sedimentary source of Fe into surface waters around South Georgia, as with the Crozet Islands (Pollard et al., 2009; Venables and Moore, 2010). Increased NO\textsubscript{3} concentrations (figure 10) corresponding with increases in F\textsubscript{v}/F\textsubscript{m} (figure 18) were also noted below 40m around the SBdy on both legs where frontal upwelling is expected to increase Fe supply to surface waters (Laubscher et al., 1993; Joubert et al., 2011). Similarly, near Antarctica photo-physiological responses were improved at depth where F\textsubscript{v}/F\textsubscript{m} increased suggest upwelling of Fe along the continental shelf.

The presence of Fe downstream of South Georgia and adjacent to Antarctica is also indicated through relative SiO\textsubscript{4} depletion (which implies active diatom growth) and community structure adjustments to a diatom dominated community (figure 17). Diatoms thrive in Fe alleviated waters (Coale et al., 1996; de Baar and Boyd, 2000) and correspond with the drawdown of SiO\textsubscript{4} (Coale et al., 1996). In the Southern Ocean, between Antarctica
and South Georgia, SiO$_4$ is never limiting (Timmermans, 2004 and figure 12), and so not expected to negatively control diatom growth. However, low SiO$_4$ concentrations (5.45 µmol l$^{-1}$) in surface waters downstream of South Georgia (figure 12) statistically correlate negatively (P <0.01 and <0.05 for tests a and c respectively) with high F$_v$/F$_m$, (figure 18) and positively with low δ$_{PSII}$ (figure 20). This indicates that where F$_v$/F$_m$ values were high SiO$_4$ concentrations were lowered, such that an Fe-replete diatom community was responsible for the reduction in SiO$_4$ concentrations. Community structure data and in situ SiO$_4$:NO$_3$ ratios further support this argument with diatoms dominating the community structure close to Antarctica and near South Georgia (figure 17), while lower SiO$_4$:NO$_3$ ratios (figure 22) found in surface waters around South Georgia support either a terrestrial runoff source of Fe or upwelling very close to the island that supports diatom growth. Similarly, the lower SiO$_4$:NO$_3$ ratios in deeper waters off Antarctica support Fe upwelling in this region. Though unusual to have enhanced diatom growth at depth, SANAE 49 HPLC data (figure 17) shows it can occur off South Georgia. Higher SiO$_4$:NO$_3$ removal ratios that depart from 1:1 towards 3:1 or more, as one moves away from South Georgia and Antarctica into open waters, implies increased Fe-limitation of diatom growth as well as a shift towards NO$_3$ uptake by phytoplankton other than diatoms. A similar scenario was observed in studies around the Crozet Islands (Moore *et al.*, 2007a, 2007b).

4.2.2.2) Community structure as a driver of photo-physiology

The development of different phytoplankton under various environmental conditions ensures adaptation, through evolution, of taxa. As different taxa are exposed to different yearly light and nutrient conditions, their photo-physiology will vary. This section looks at the community structure (though HPLC data), and the influence it has on the photosynthetic parameters over the summer of 2009.
It has already been shown that the ratio of diatoms to haptophytes significantly (P <0.0001) controls $F_v/F_m$, where the presence of actively growing diatoms elevate $F_v/F_m$, and less significantly (P <0.01) $\delta_{PSII}$ values.

Individually the three predominant phytoplankton groups (diatoms, haptophytes and chromophytes) have a less significant effect on the photosynthetic parameters compared to the Fuc:Hex ratio. An increase in haptophytes sees a significant decrease (P <0.05) in $F_v/F_m$, while an increase in diatoms and chromophyte both see a slight significant (both P <0.05) increase in $\delta_{PSII}$.

These statistical results show that when looking at photo-physiology and community structure; it is the population as a whole (seen by Fuc:Hex ratio) which is important in influencing the photosynthetic efficiency, rather than the individual phytoplankton groups.

The best-known haptophytes are coccolithophores and *Phaeocystis* species. Coccolithophores, with their calcareous ‘liths’, are key particulate inorganic carbon (PIC) exporters in the biological pump, despite CO$_2$ being released to the atmosphere during the process of CaCO$_3$ formation, (Poulton *et al*., 2007). Coccolithophores tend to occupy the open ocean basins north of the APF, in warmer temperatures and lower calcite saturation states (Poulton *et al*., 2007, 2010). *Phaeocystis* species occurs in the open ocean basins south of the APF where low SiO$_4$ concentrations limit diatom growth, with the exception of the Ross Sea, where the *Phaeocystis*’s efficient photo-physiology allows an early spring bloom (Arrigo *et al*., 1999; Boyd, 2002). *Phaeocystis* species are also dominant in the Atlantic section of the Southern Ocean near the ice shelf (Gibberd *et al*., 2013). Diatoms are found in
Fe alleviated area such as around fronts and islands (Sakshaug et al., 1991; Laubscher et al., 1993; de Baar et al., 1995; Smetacek et al., 1997; Boyd, 2002; Gibberd et al., 2013). Diatoms, with their silica ballast, are also important exporters of carbon through the biological pump. Diatoms and *Phaeocystis* species are often the dominant phytoplankton taxa in Southern Ocean waters (Arrigo et al., 1999; Boyd, 2002; Poulton et al., 2007; Hassler et al., 2012). Chromophytes on the other hand are scarce in the Southern Ocean relative to diatoms and haptophytes and are more usually dominated by brown seaweeds (Jeffrey et al., 1997).

The simplified statement that diatoms dominate around South Georgia and adjacent to Antarctica, while haptophytes occupy in the open ocean basins, can be explained through the Fe hypothesis. This hypothesis states that the physiological nature of phytoplankton and the environment in which they live will determine their distribution. The Fe hypothesis, first proposed by Martin et al., in 1990, explains the dominance of smaller pico- and nano phytoplankton (Gervais et al., 2002) such as haptophytes in Fe limited seas. Unlike larger phytoplankton (e.g. large diatoms), smaller species have high surface to volume ratios facilitating the uptake of Fe at low concentrations (Sunda and Huntsman, 1995). Smaller species, with lower overall demands for Fe, are likely to have higher photosynthetic efficiency under low Fe conditions, compared to larger phytoplankton, which cannot function optimally under the same low Fe conditions. When an area is freed from Fe limitation the community structure changes, becoming dominated by larger phytoplankton species such as diatoms (de Baar and Boyd, 2000; Smetacek et al., 2004; Seeyave et al., 2007). Apart from increased growth rates in Fe-replete waters, larger phytoplankton are less prone to grazing pressure (Hoffmann et al., 2006), so emerge as a dominant group. Herbivorous mesozooplankton that feed on large cells including large diatoms have a relatively long
generation time and therefore cannot control a fast growing phytoplankton biomass where doubling times are measured in hours rather than days. On the other hand, microzooplankton have a short generation time, and so are able to control the biomass of smaller phytoplankton species (Coale et al., 1996).

Unlike the other diatom patches near South Georgia and the Antarctic continent mentioned earlier, which all corresponded with high $F_v/F_m$ values, a more unusual diatom patch was encountered just north of 64º S (figure 17) where $F_v/F_m$ values were relatively low (~0.3) (figure 18). A possible explanation for the low photo-physiological efficiency of this particular patch of diatoms is that it developed previously due to an input of Fe into the system, which has now been exhausted. At low Fe concentrations the diatoms start becoming less photosynthetically efficient (Greene et al., 1991), but have not yet decreased in dominance. The weakening of this populations photosynthetic efficiency will likely result in a decrease in their dominance, and a shift towards a smaller phytoplankton population that is more efficient at taking up Fe at lower concentrations such that the $F_v/F_m$ values will once again increase, though this is not yet apparent in this patch. Hence, phytoplankton efficiency increases in response to both Fe alleviation and an associated shift in community structure; however the photosynthetic parameters indicate the health of the population rather than its population dynamics.
4.2.2.3) Other factors influencing $F_v/F_m$ and $\phi_{PSII}$

**Chlorophyll-a**

Chl-a concentration exhibits a significant positive correlation with $F_v/F_m$ ($P < 0.001$, $< 0.01$ for tests a and c respectively) and a significant negative correlation with $\phi_{PSII}$ ($P < 0.01$, $< 0.05$ for linear regressions tests b, and d respectively). This is hardly surprising for two reasons. Firstly, high chl-a concentrations imply fast growth rates of phytoplankton, such that biomass is accumulating despite predation pressure from zooplankton (Fielding *et al.*, 2007). This would not occur in an environment that was limited by either Fe or light. Secondly, although this is chl-a specific, where there is more chlorophyll present per phytoplankton cell, the more effective it will be at photosynthesis, thus reflecting higher $F_v/F_m$ values and lower $\phi_{PSII}$ values. Similar results have been recorded in a number of Fe addition bio-assay experiments where both chl-a concentrations and $F_v/F_m$ values are shown to increase in response to Fe addition while $\phi_{PSII}$ decreases (Moore *et al.*, 2007a, 2007b).

**PAR and fluorescence**

The Southern Ocean is often considered to be light limited (Hiscock, 2004), either due to low ambient PAR or to deep MLD’s. Light is a requirement for photosynthesis, such that $F_v/F_m$ is driven by a changing light field which is particularly significant in the often light limited Southern Ocean (Sunda and Huntsman, 1997; Lindley and Barber, 1998; Timmermans *et al.*, 2001; Moore *et al.*, 2007a, 2007b). Evolutionary trends indicate that different phytoplankton groups can be found at different light depths. Between these groups, the arrangement of pigments housed within the light-harvesting antennae will vary greatly.
(MacIntyre et al., 2002; Johnsen and Sakshaug, 2007), leading to differences in both PS II light-harvesting potential and efficiency (Lutz et al., 2001). This, along with the fact that the number of photochemically competent reaction centres can vary as a function of irradiance (Neale, 1987; Long et al., 1994; Vassiliev et al., 1994; Babin et al., 1996) ensures that $F_v/F_m$ and $\phi_{PSII}$ values vary with PAR and fluorescence (Suggett et al., 2009), as seen by the statistical analyses.

### III: Nutrient ratios and phytoplankton community structure

Nutrient inputs through ocean circulation, and winter mixing resets the Southern Oceans nutrient concentrations annually. Therefore seasonal changes in the nutrient ratios reflect nutrient uptake by the phytoplankton community structure that forms. However, the relationship between the source, use and therefore concentration of nutrients and the community structure of phytoplankton is complex. A diatom dominated community takes up $\text{SiO}_3$ and $\text{NO}_3$ and leaves behind a characteristically low $\text{SiO}_4:\text{NO}_3$ ratio (<1.5). However when Fe is limiting, haptophytes dominate the community such that the $\text{SiO}_4:\text{NO}_3$ uptake ratio increases, leaving a trace in the ratio $\text{SiO}_4:\text{NO}_3$ left behind (Takeda 1998). Fe availability also drives $\text{SiO}_3$ and $\text{NO}_3$ removal by individual species, with a limitation of Fe limiting $\text{NO}_3$ removal and enhancing the uptake of $\text{SiO}_4$ per unit of $\text{NO}_3$.

Concerns over the quality of the SANAE 49 nutrient data (Section 2.4.3), undermine the confidence of the following discussion concerning the SANAE 29 nutrient ratio’s and community structure. However, this lack of confidence does not apply regarding the SANAE 50 data sets.
4.3.1) SANAE 49: Nutrient ratio’s and Community Structure

The NO$_3$:PO$_4$ ratio bore no relationship to the community structure using HPLC, in contrast with Arrigo et al., (1999) who showed that the NO$_3$:PO$_4$ draw-down ratio was twice as high for *Phaeocystis antarctica* (19.2 ± 0.61) than for diatoms (9.69 ± 0.33). The SiO$_4$:NO$_3$ removal ratio does however reflect community composition, in that low SiO$_4$:NO$_3$ ratios (figure 22) coincide with diatom dominated communities (figure 17). This confirms Holeton’s et al., (2005) finding that variation in SiO$_4$ concentrations can indicate the presence of diatoms. In waters between Antarctica and South Georgia, a high SiO$_4$:NO$_3$ ratio was linked with high haptophyte and low diatom concentrations, as well as low Fuc:Hex values.

4.3.2) SANAE 50: Nutrient ratio’s and Community Structure

The low SiO$_4$:NO$_3$ drawdown ratio’s (table 3) found north of the SACC, around South Georgia (0.25 and 0.32 for the north westward and south eastward legs respectively) reveal a diatom dominated community in this region. This finding is supported by Gibbert et al., (2013). The dominance of diatoms can be explained by the presence of Fe (diatom dominated communities are found in areas of sufficient Fe [Sakshaug et al., 1991; Laubscher et al., 1993; de Baar et al., 1995; Smetacek et al., 1997; Boyd, 2002]). A 1:1 removal ratio indicates that both nutrients have been taken up by the phytoplankton in a 1:1 molar ratio (Brzezinski, 1985), implying Fe-availability.

Between South Africa and Antarctica, the regions between the northern STF and APF all reveal a low (<1.5) SiO$_4$:NO$_3$ removal ratio (table 3), suggesting a diatom dominated
community. These regions are all narrow frontal regions, which accounts for the diatom dominance which this ratio reveals (Laubscher et al., 1993; Joubert et al., 2011).

Between Antarctica and South Georgia, the moderately high (<1.5) SiO$_4$:NO$_3$ removal ratio’s found south of the SACCF (table 3) reveal a high removal of NO$_3$ relative to SiO$_4$. This suggests haptophyte-dominated community and associated iron limitation in this region. The high (>1.5) SiO$_4$:NO$_3$ ratio between South Africa and the northern STF reveals a haptophyte dominated community within this region, this finding is supported by Gibbert et al., 2013.

The changing removal ratios from a low (1.6) to a high (4.8) between South Africa and the northern STF reveal a shift to a more haptophyte dominated community here. While the inverse is seen between the APF and the iceshelf revealing a shift in community structure (between the southern and northward legs), with increasing diatom numbers, resulting in an outright shift to diatom dominance on the northward leg between the APF and the SBdy.

The positive differences in SiO$_4$ and NO$_3$ between the earlier north westward and south eastward legs reveal a continued drawdown of both these nutrients in these regions. The negative differences in SiO$_4$ (-1.45 µmol l$^{-1}$) and NO$_3$ (-0.76 µmol l$^{-1}$) between the earlier north westward and south eastward legs (around South Georgia) reveals an introduction of SiO$_4$ into the region (probably through upwelling, which may indicate Fe upwelling in the region as well), together with a tendency for phytoplankton to use nitrogen in the form of NH$_4$ rather than NO$_3$. Similarly, the negative difference (-0.75 µmol l$^{-1}$) in SiO$_4$ between the southward and northward legs, north of the northern STF also reveals an input of SiO$_4$ into the system around South Africa. While the negative difference in NO$_3$ (-25.01 µmol l$^{-1}$)
between the southward and northward legs south of 69º S show a preference for the phytoplankton to use NH₄ or nitrogen assimilation at the ice shelf.

IV: Experimental Fe incubation results during SANAE 50

4.4.1) Overall control and contamination of experiments

The form of Fe added in these experiments was FeSO₄; chosen for two reasons: Firstly, through the atmospheric deposition of dust, it is believed that Fe is transported between systems in the form of FeSO₄ (Zhuang and Duce, 1993). Secondly, IronEx2 clearly shows that any changes brought about by the addition of acidic FeSO₄, is due to the Fe, not the sulphate or miniscule changes in pH (Coale et al., 1996). Hence, changes brought about in phytoplankton production can be attributed to the Fe only.

Three of the experiments (B, C and D) contained overall controls (figure 24). These were bottles at 50% light that remained closed for the duration of the experiment, any difference between this bottle, and the 5th day 50% control bottles highlights any subsampling contamination in the experimental bottles. Experiment D reveals no differences in Fᵥ/Fₘ or ðPSII between the overall control and 50% control bottles on the 5th day, thus showing no contamination during subsampling in this experiment. Experiment C shows that the overall control (0.18) has a lower Fᵥ/Fₘ value than the 5th day 50% control (0.26 ± 0.048), revealing that there was probably slight contamination during the removal of sub-samples, and that the Fe fertilization has a greater effect than shown in this study (this contamination signal is not evident in ðPSII, suggesting that the contamination is only slight). Experiment B shows the Fᵥ/Fₘ for the overall control (0.38) to be higher than the 5th day 50% control (0.29
± 0.049) (no difference in $\delta_{\text{PSII}}$), suggesting that sub sampling saw the contamination with a substance that had a negative effect on the phytoplankton. Overall, there was little (experiment B and C) or no (experiment D) contamination during subsampling during this study.

4.4.2) Introduction to experiments:

The SOIREE (Boyd and Abraham, 2001) and CROZEX (Moore et al., 2007a, 2007b) Fe fertilization experiments clearly revealed an increase in photosynthetic efficiency of the phytoplankton (through an increase in $F_v/F_m$ values and a decrease in $\delta_{\text{PSII}}$ values), an increase in chl-a concentrations, a decrease in NO$_3$ concentration, and an increase in CO$_2$ drawdown. However the degree to which the efficiency is increased as well as the magnitude of the resulting Chl-a increase is still a matter of discussion. In this investigation, the Fe alleviation incubations were conducted in varying areas of the Southern Ocean south of Africa, under varying initial conditions. The experiments clearly show an increase in the photo-physiological health of phytoplankton, as previously shown (Boyd and Abraham, 2001; Moore et al., 2007b). The variation seen in the photophysical efficiencies response to Fe addition between experiments, and the factors behind it, are discussed below.

4.4.3) Relationship of photo-physiology to Fe alleviation

All the experiments show a statistically significant increase in photosynthetic ability for both, or, either $F_v/F_m$ and $\delta_{\text{PSII}}$ when Fe is added (figure 24 and table 6). Thus, when Fe is added in the Southern Ocean, the photosynthetic efficiency of phytoplankton increases, however, the degree of this changes, and the response time varies.
The variation observed between experiments, despite an overall positive response to Fe alleviation, is consistent with the results of Moore et al., (2007b). The differences they observed were accounted for by variability in initial community structure, stage of the bloom and availability of Fe and possibly silicic acid prior to manipulation (Moore et al., 2007a).

In this study, Experiment B, performed south of the SBdy in ice-free conditions (table 4), shows the most statistically significant changes in photosynthetic efficiency. The addition of Fe caused an increase in efficiency in both $F_v/F_m$ and $\Delta_{PSII}$ at low light ($F_v/F_m$ increased from 0.27 ± 0.000 to 0.47 ± 0.016 and $\Delta_{PSII}$ decreased from 214 ± 0.0 to 189 ± 15.9 [while the controls ended after five days at 251 ± 16.9 and 0.28 ± 0.031 for $F_v/F_m$ and $\Delta_{PSII}$ respectively]). This improvement in efficiency was also seen at high light levels (where $F_v/F_m$ increased from 0.27 ± 0.000 to 0.47 ± 0.014 and $\Delta_{PSII}$ decreased from 214 ± 0.0 to 198 ± 6.1 [while the controls ended after five days at 0.29 ± 0.049 and 219 ± 13.5 for $F_v/F_m$ and $\Delta_{PSII}$ respectively]). Although there is no statistical difference between the Fe addition bottles at 50% and 25% light, the clear statistical differences (figure 24 and table 6) between the Fe supplemented bottles and control bottles without Fe shows that Fe additions improved photosynthetic efficiency in all regions sampled in the Southern Ocean and consequently that primary production in these regions is Fe limited.

Experiment D, conducted at the ice shelf (table 5), was the only site that showed a statistical difference (final difference at day 5: 0.14 and 34 for $F_v/F_m$ and $\Delta_{PSII}$ respectively) in photophysical responses in the control bottles without Fe alleviation at the two different light levels (table 6). This difference in the importance of light levels diminishes under Fe alleviation. As at 50% light (Fe addition), only $\Delta_{PSII}$ remains significant ($P = 0.0025$), while at 25% light under Fe alleviation both the photophysical ($P$ value for $F_v/F_m$ and $\Delta_{PSII}$ is 0.0001
and 0.0000005 respectively) and chlorophyll (P =0.041) response improved significantly. Thus, at the iceshelf, Fe significantly changes the ability of phytoplankton to photosynthesis efficiently at depth where light is limited.

For experiments C (conducted south of the SBdy, in the presence of icebergs, table 4) and F (conducted in the Antarctic Polar Zone [APZ], table 4), both physiological indices (Fv/Fm and ðPSII) responded to Fe alleviation, however this positive response occurred at only one light depth but not the other. For experiment C, Fe alleviation elevated the physiological response at 50% light, whereas in Experiment F, Fe alleviation elevated the physiological response at 25%. These results suggest that the response to light and Fe co-limitation is not entirely predictable and almost certainly complicated by the taxonomic composition of the community, which is adapted to the in situ light environments (see also Moore et al., 2007a, 2007b). Complications introduced by the taxonomic composition of the community is discussed under the next section (Fe alleviation and chlorophyll response).

Again as Moore et al., (2007a, 2007b) noted, responses in the incubation experiments showed variability from day zero to 1-2 days that were independent of initial weather conditions or the previous light history (PAR) experienced by the phytoplankton. Since sampling of the experiments was done at night, potential ‘light-shock’ effects can be disregarded. Temperature does not significantly alter photo-physiological responses either, as shown by the statistical analyses in Chapter three (Section 3.1.5). Variability in Fv/Fm and ðPSII between experiments was almost certainly due to different taxa, as previously noted. Changes in Fv/Fm and ðPSII responded relatively quickly, but changes in biomass and nutrient take longer.
4.4.4) Fe alleviation and chlorophyll response

Past literature (Boyd and Abraham, 2001; Moore et al., 2007b) clearly indicates that Fe alleviation, whether natural or artificial causes a chl-a bloom that occurs after the PS II and PS I responses. Chl-a concentrations increased during all treatments in all experiments, in the Fe-supplemented bottles, but also in the controls. However, a statistically significant difference between the Fe addition and controls was only evident in three of the experiments in this investigation (experiment D, E and F).

In experiment D (conducted at the ice shelf, table 4) the chl-a in the 25% Fe alleviated bottles was higher (7.1 µg.l$^{-1}$ ± 0.32) than in the controls (4.3 µg.l$^{-1}$ ± 0.54) (there was an overall significant difference [p =0.02] between all the Fe bottles and all the controls). Similarly for experiment E (conducted between the SBdy and SACCF, table 4) chl-a in the 50% Fe alleviated bottles was (0.49 µg.l$^{-1}$ ± 0.04) versus the controls (0.42 µg.l$^{-1}$ ± 0.04) and in experiment F (conducted in the APZ, table 4) chl-a in the 50% Fe alleviated bottles was (2.1 µg.l$^{-1}$ ± 0.05) versus the controls (1.3 µg.l$^{-1}$ ± 0.24). Experiment F also showed an overall significant difference [p =0.02] between all the Fe bottles and all the controls. These results indicate that in three of the experiments Fe addition led to a significant increase in chl-a (relative to the non Fe controls) which is likely a direct result of enhanced productivity via increased photosynthetic efficiency in the Fe addition bottles.

FRRf can be interpreted in relation to community structure (Suggett et al., 2004, 2009; Moore et al., 2005) rather than photosynthetic efficiency affected by nutrient stress (Kolber et al., 1988; Greene et al., 1991; Boyd and Abraham, 2001). This could be used to further explain the lack of a significant chl-a increase in Fe alleviated incubations. Rather than an increase of the photosynthetic efficiency of the phytoplankton species present, the
FRRf values could show a changing community structure in response to Fe alleviation. However, a study of the controlling factors of Fv/Fm and \( \sigma_{PSII} \) (Section 2 of this discussion) reveal that the changes in FRRf parameters are more likely to be related to Fe related physiological changes then a community structure shift.

The lack of any statistically significant chl-a response in the remaining three experiments (A, B and C) may be explained by two different possibilities. Firstly the original community structure of the phytoplankton and secondly by bottle effects.

4.4.4.1) Community structure and Fe alleviation

The influence of community structure on Fe fertilization and resulting chl-a blooms is based on evolutionary theory. Different oceanic regions are subject to different conditions in which phytoplankton need to adapt and thrive, resulting in differing community structures throughout the world’s oceans. As phytoplankton species have adapted to different conditions, they will respond differently to changes in their environment. The community structure for these incubations was gained through the nutrient data (Section 3 of this discussion) and backed up by past published research.

Experiment’s A, B and C all began with high SiO\(_4\):NO\(_3\) ratio’s (>1.5) found in the waters between Antarctica and South Georgia (before the SAGCF). These experiments were thus most likely initiated in haptophyte-dominated communities of the Fe-limited open ocean.

Conversely, experiments D, E and F are all considered to be initiated in a community where diatoms were prevalent. Low SiO\(_4\):NO\(_3\) ratios of experiment E and F were 1.45 and
0.12 respectively revealing a diatom dominated community. Diatoms are known to be prevalent in the high Fe waters of the marginal ice zone of Experiment D (Laubscher et al., 1993; de Baar et al., 1995; de Baar and Boyd, 2000; Boyd, 2002; Dierssen et al., 2002; Lannuzel et al., 2006). Despite the high SiO$_4$:NO$_3$ ratios found at this station, Ceinwen Smith, working on microscopy data, confirmed a haptophyte dominated community, but with the presence of the chain forming Chaetoceros diatoms (personal communication), the finding of a haptophyte dominated community in this marginal ice zone is further confirmed for the late summer of 2009 by Gibberd et al., (2013).

It is significant that the three experiments which developed statistically significant increases in chl-a all began in either diatom dominated (experiment E and F) communities or in communities containing large chain forming diatoms (experiment D). Diatoms have a faster growth rate than the herbivorous mesozooplankton that prey on them. As such they are able to escape grazing pressure allowing an increase in population abundance and the statistically significant increase in measured chl-a. Conversely, experiments A, B and C began in smaller haptophyte dominated communities, where an increase in Fe probably led to an increase in production (as witnessed by the increase in photosynthetic parameters). However, the microzooplankton which graze on small haptophytes and other small taxa have a short generation time that is more evenly matched by their haptophyte prey, such that they were able to control net community growth, so preventing an increase in observed chl-a biomass (Coale et al., 1996).
4.4.4.6) Bottle effects

A second theory states that bottle effects could be the reason for a lack of distinct chl-a increase when there is a clear and positive physiological response. Closed systems can create bottle effects that bias the results, through potentially unrealistic ecosystem dynamics due to the loss of grazers and advective processes. Grazing rates increase with increasing biomass accumulation, thus it is expected that grazing would lower the chl-a accumulation in the Fe addition bottles. This, along with the slight contamination seen at the beginning of the experiments (which increases the response in the control bottles), would remove or lower the statistically significant difference between Fe addition bottles and their controls. However, one would expect such an effect to be experienced by all the experiments (Cullen et al., 1992; Geider and Laroche, 1994; Moore et al., 2007a).

V: Conclusions

Between the two years of study, the chemical and physical environment of the South Atlantic sector of the Southern Ocean varied little. Variation in the intensity and distribution of the chl-a blooms is explained through variation in fronts, localised wind events and ice coverage between the two summers. Compared with the open ocean, the photosynthetic health of phytoplankton increases remarkably throughout the water column near South Georgia, and to a lesser degree, near Antarctica. Secondary indicators of the presence of Fe, all individually suggest Fe alleviation in these regions. Phytoplankton efficiency increases in response to both Fe alleviation and an associated shift in community structure; however the photosynthetic parameters indicate the health of the population rather than its population dynamics.
Increased Fe leads to an increase in the physiological efficiency of phytoplankton in the Southern Ocean and where diatoms were present, lead to an increase in chl-a biomass. This proves that the majority of the Southern Ocean is Fe limited, and that an increase in Fe would lead to an increase in photosynthetic efficiency, but not necessarily biomass or carbon export.

Given the anticipated changes in light and Fe availability to the Southern Ocean (Sarmiento et al., 1998; Boyd, 2002; Hillel and Rosenzweig, 2002; Greenblatt and Sarmiento, 2004), due to climate change, and the important role the Southern Ocean plays in alleviating atmospheric increases in CO$_2$ through the biological carbon pump (Volk and Hoffert, 1985; Longhurst, 1991; Schlitzer, 2002; Falkowski and Raven, 2007), assessing phytoplankton responses to Fe and light co-limitation is important. This research provides us with a better understanding of the nuances of the response of the Southern Oceans phytoplankton physiology and community structure to Fe and light variability and the impact this has on the potential for carbon export.
Appendixes

List of Appendix’s

Appendix A: A T-S plot of all the CTD stations on a) the north westward leg and b) the south eastward leg of SANAE 49 used to determine the position of the water masses.

Appendix B: MATLAB code by Dr Brian Hopkinson, a model determining $F_v/F_m$ and $\sigma_{PSII}$ values.

Appendix C: MATLAB code by Dr Mark Moore, a model determining $F_v/F_m$ and $\sigma_{PSII}$ values.

Appendix D: Graphs depicting a) the northward leg of NO$_3$ for SANAE 48, 49 and the adjusted NO$_3$ values for SANAE 49, b) The northward leg of PO$_4$ for SANAE 48 and 49, and c) the NO$_3$:PO$_4$ ratio for the northward leg of SANAE 49.

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Appendix F: Pigments to species table.

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Appendix H: Weather observations during experiments.

Appendix I: The nutrient results of experiment B.

Appendix J: Critique and recommendations on the bio-assay nutrient methods.
Appendix A: A T-S plot of all the CTD stations on a) the north westward leg and b) the south eastward leg of SANAE 49 used to determine the position of the water masses.
Appendix B: MATLAB code by Dr Brian Hopkinson, a model determining $F_v/F_m$ and $\sigma_{PSII}$ values.

```matlab
files = dir ('BR*.*')
for i = 1 : numel(files)
    infile = files(i).name;
    fid = fopen (infile);
    for j = 1:3
        waste = fgetl(fid);
    end
    line = fgetl(fid);
    gain = sscanf(line(6:9), '%i');
end
blanks = dir('ABL*.*)';
for x = 1 : numel(blanks)
    test = blanks(x).name;
    fidblank = fopen (test);
    for j = 1:3
        waste = fgetl(fidblank);
    end
    line = fgetl(fidblank);
    blankgain = sscanf(line(6:9), '%i');
    if gain == blankgain
        blank = blanks(x).name;
    end
    fclose(fidblank);
end
fclose(fid);
outfile = 'output.txt';
iter = 1;
test_firefit(infile, blank, outfile, iter);
```

```matlab
function paramsfit = test_firefit(infile, blank, outfile, iter)
% first need to open file and read header.
% now get data.
% infile = 'V1P27_1.000';
% blank = 'LP2400.blk';
% outfile = 'output.out';
% iter = 1;
disp(infile);
doplots = 0;
% infile = 'INC1_T0.000';
% blank = 'INC1_T0B.000';
% outfile = 'INC1_T0.pro';

fid = fopen(infile,'r');
rewind(fid);

% read in relevant information from header
for i = 1:6
    waste = fgetl(fid);
end;

line = fgetl(fid);
STS_num = sscanf(line(14:16), '%i'); % number of samples in single turnover saturation
line = fgetl(fid);
```

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STR_num = sscanf(line(19:21), '%i'); % number of samples in ST relaxation
line = fgetl(fid); % discard STRI
line = fgetl(fid); % discard MTF duration
line = fgetl(fid);
MTS_num = sscanf(line(19:21), '%i'); % number of samples in MT saturation
line = fgetl(fid);
MTR_num = sscanf(line(19:21), '%i'); % number of samples in MT relaxation
for i = 1:3
    waste = fgetl(fid); % discard additional lines
end;
line = fgetl(fid);
PARint = sscanf(line(18:23), '%f'); % input PAR intensity for P vs E curves
line = fgetl(fid);
if line(1) == 'C'
    lskip=4;
end
if line(1) == 'N'
    lskip = 7;
end
for i = 1:lskip % need to discard different number of lines depending on whether this is a standard sample or part of PvE curve
    waste = fgetl(fid);
end;
% read in sample file
% read in single turnover saturation
data = fscanf(fid,'%f %f %f %f',[4 inf]);
data = data';
STSsamp = round(data(1:STS_num,1))';
STS_t = data(1:STS_num,2)';
STS_ex = data(1:STS_num,3)';
STS_fl = data(1:STS_num,4)';

% read in ST relaxation
STRsamp = round(data(1:STR_num+1:STR_num+STR_num,1))';
STR_t = data(1:STR_num+1:STR_num+STR_num,2)';
STR_ex = data(1:STR_num+1:STR_num+STR_num,3)';
STR_fl = data(1:STR_num+1:STR_num+STR_num,4)';

N = STS_num+STR_num;
% read in multi-turnover saturation
MTSsamp = round(data(N+1:N+MTS_num,1))';
MTS_t = data(N+1:N+MTS_num,2)';
MTS_ex = data(N+1:N+MTS_num,3)';
MTS_fl = data(N+1:N+MTS_num,4)';
N = N+MTS_num;
% read in multi-turnover relaxation
MTRsamp = round(data(N+1:N+MTR_num,1))';
MTR_t = data(N+1:N+MTR_num,2)';
MTR_ex = data(N+1:N+MTR_num,3)';
MTR_fl = data(N+1:N+MTR_num,4)';
fclose(fid);
clear data;
%read in blank file
fib = fopen('blank','r');
rewind(fib);
for i = 1:16
    line = fgetl(fib);
end;
line = fgetl(fib);
if line(1) == 'C'
    lskip=4;
end
if line(1) == 'N'
    lskip = 7;
end
for i = 1:lskip  %need to discard different number of lines depending on whether this is a standard sample or part of PvE curve
    waste = fgetl(fib);
end;
data = fscanf(fib,'%f %f %f %f',[4 inf]);
data = data';
STSsampb = round(data(1:STS_num,1))';
STS_tb = data(1:STS_num,2)';
STS_exb = data(1:STS_num,3)';
STS_flb = data(1:STS_num,4)';
%read in ST relaxation
STRsampb = round(data(STS_num+1:STS_num+STR_num,1))';
STR_tb = data(STS_num+1:STS_num+STR_num,2)';
STR_exb = data(STS_num+1:STS_num+STR_num,3)';
STR_flb = data(STS_num+1:STS_num+STR_num,4)';
N = STS_num+STR_num;
%read in multi-turnover saturation
MTSsampb = round(data(N+1:N+MTS_num,1))';
MTS_tb = data(N+1:N+MTS_num,2)';
MTS_exb = data(N+1:N+MTS_num,3)';
MTS_flb = data(N+1:N+MTS_num,4)';
N = N+MTS_num;
%read in multi-turnover relaxation
MTRsampb = round(data(N+1:N+MTR_num,1))';
MTR_tb = data(N+1:N+MTR_num,2)';
MTR_exb = data(N+1:N+MTR_num,3)';
MTR_flb = data(N+1:N+MTR_num,4)';
%subtract blank from sample
STS_fl = STS_fl - STS_flb;
STR_fl = STR_fl - STR_flb;
MTS_fl = MTS_fl - MTS_flb;
MTR_fl = MTR_fl - MTR_flb;
%fits single turnover saturation
%define irradiance applied in saturation phase
skip = 2 ; %samples to skip at start of STS phase
irrad = 47248; %irradiance in uEi/m2/sec, from calibration sheet

irrad = irrad * 1E-6 * 6.02E23; %irradiance in quanta/m2/sec
STStint = 1E-6; %'flashlet' duration in sec, or time of light application between sampling (light is on constantly)
sigscale = 1e-20; % sigma in m2/quanta
pulse = irrad * STStint * sigscale;

for i = 1:STS_num,
pfd(i) = pulse;
end;

STS_fl2 = STS_fl(skip+1:STS_num);
STSparams = [STS_num skip];

%make initial guesses
fog = mean(STS_fl(3:5));
fmg = mean(STS_fl(STS_num - 4:STS_num));
sigg = 1000; % 1e-20 m2/quanta, see scaling of pfd above
pg = 0.6;
x0 = [fog, fmg, sigg, pg];

%set search bounds
ming = [fog*0.5, fmg*0.5, 10, 0]; %set lower bounds for fo, fm, sig, p
maxg = [fog*1.5, fmg*1.5, 2500, 1]; %set upper bounds for fo, fm, sig, p

%STSfitted = STSfit(x0, pfd, STSparams);
%plot(STS_t(skip+1:STS_num),STSfitted,STS_t,STS_fl,'r');

%fog = mean(STS_fl(3:5));
%fmg = mean(STS_fl(STS_num - 4:STS_num));
sigg = 1000; % 1e-20 m2/quanta, see scaling of pfd above
pg = 0.6;
x0 = [fog, fmg, sigg, pg];

%x = [fog, fmg, sigg, pg];
%ming = [fog*0.5, fmg*0.5, 10, 0]; %set lower bounds for fo, fm, sig, p
%maxg = [fog*1.5, fmg*1.5, 2500, 1]; %set upper bounds for fo, fm, sig, p

%STSFitted = STSfit(x0, pfd, STSparams);
%plot(STS_t(skip+1:STS_num),STSFitted,STS_t,STS_fl,'r');

%set options for lsqcurvefit
opts = optimset('lsqcurvefit');
%opts = optimset(opts,'Display','off');
%opts = optimset(opts,'MaxIter',4000);
%opts = optimset(opts,'Diagnostics','off');
%opts = optimset(opts,'MaxFunEvals',24000);
%opts = optimset(opts,'TolFun',1e-9);
%opts = optimset(opts,'TolX',[0.001 0.001 0.1 0.01]);

[xfit, resnorm, residual, exitflag, output] = lsqcurvefit('STSfit', x0, pfd, STS_fl2, ming, maxg, opts, STSparams);
%lsqcurvefit setup
%[x,resnorm,residual,exitflag,output] = lsqcurvefit(fun,x0,xdata,ydata,lb,ub,options)
fvfm = (xfit(2)-xfit(1))/xfit(2);

STSbestfit = STSfit(xfit, pfd, STSparams);

if doplots == 1,
    figure(iter); %open new figure window for each iteration of the firefit program
    subplot(2,2,1);
    plot(STS_t(skip+1:STS_num),STSbestfit,STS_t,STS_fl2,'r'),
    title(strcat(infile,' STS'));
end;
%fit single turnover relaxation data, oxidation of Qa

fo = mean(STR_fl(STR_num - 3: STR_num));
fm = mean(STR_fl(1:2));
params = [fo fm];
tau0 = 4000;
%oxidation timescale of Qa in us
taumin = 10;
%minimum tau 10 us
taumax = 100000;
%maximum tau 100 ms

[taufit, resnorm, residual, exitflag, output] = lsqcurvefit('STRfit',
tau0,STR_t, STR_fl,taumin, taumax,opts, params);
STRbestfit = STRfit(taufit, STR_t, params);

if doplots == 1,
 subplot(2,2,2);
 plot(log(STR_t), STRbestfit, log(STR_t), STR_fl, 'r'), title('STR');
end;

%fit multi turnover saturation data
%divide up MT phase into chunks, average, and chose max of these

nbblocks = fix(MTS_num / 20); %break into blocks of 20 samples, leaving off trailing samples after final block of 20;

for i = 1:nblocks,
   Blockavg(i) = mean(MTS_fl(1 + 20*(i-1):20*i));
end;
fmMT = max(Blockavg);

if doplots == 1,
 subplot(2,2,3);
 plot(MTS_t,MTS_fl), title('MTS');
end;

%fit relaxation phase after multi turnover saturation, oxidation of PQ pool.

foMT = mean(MTR_fl(MTR_num - 3: MTR_num));
fmMT = mean(MTR_fl(1:3));
paramsMTR = [foMT fmMT];
tauPQ_0 = 20000; % oxidation time of PQ in us, guess 20 ms
tauPQ_min = 1000; % minimum oxidation time 1 ms
tauPQ_max = 200000; % maximum oxidation time 200 ms

[tauPQ_fit, resnorm, residual, exitflag, output] = lsqcurvefit('MTRfit',
tauPQ_0, MTR_t, MTR_fl,tauPQ_min, tauPQ_max, opts, paramsMTR);
MTRbestfit = MTRfit(tauPQ_fit, MTR_t, paramsMTR);

if doplots == 1,
 subplot(2,2,4);
 plot(log(MTR_t), MTRbestfit, log(MTR_t), MTR_fl, 'r'), title('MTR');
end;

%export to an outfile
first = exist(outfile); %test if the outfile has
already been created and written to
if first == 0, %for the first time write
 header and data to file
 fid = fopen(outfile, [w]);
 fprintf(fid,'sample \t fo \t fm \t fv/fm \t sig \t p \t tauQa \t fmMT \t tauPQ \t tau \n');
end;
fprintf(fid,'%s 	 %5.2f 	 %5.2f 	 %1.3f 	 %6.2f 	 %2.2f 	 %6.2f 	 %5.2f 	 %6.2f 	 %6.2f',infile, xfit(1), xfit(2), fvfm, xfit(3), xfit(4),...
    taufit, fmMT, tauPQ_fit, PARint);
else                                        %after first access just write data
    fid = fopen(outfile,'a');
    fprintf(fid,'%s 	 %5.2f 	 %5.2f 	 %1.3f 	 %6.2f 	 %2.2f 	 %6.2f 	 %5.2f 	 %6.2f 	 %6.2f',infile, xfit(1), xfit(2),fvfm, xfit(3), xfit(4),...
    taufit, fmMT, tauPQ_fit, PARint);
    fprintf(fid,'
');
end
fclose(fid);

paramsfit = [xfit(1) xfit(2) xfit(3) xfit(4) taufit fmMT tauPQ_fit PARint];
return;

Appendix C: MATLAB code by Dr Mark Moore, a model determining Fv/Fm and σPSII values.

% This code fits variable fluorescence data from the Satlantic FIRe instrument to the model of Kolber et al. 1998
% Version V3 disregards the first 3 micro seconds of data from the ST saturation phase
% IMPORTANT NOTE! The Connectivity value recovered when disregarding the first 3 micro seconds is likely to be inaccurate

clear all;
close all;

LED_intensity = 31000; % muMol m-2 s-1 % Must be set for absolute values of sigPSII

% Request input files
in_file = input('Enter raw data file name (excluding extensions): ','s');
d = dir([in_file '.0*']); % Look for all files of correct format. This should be changed if there are >100 files!

for filecount = 1:length(d)
    fid = fopen(d(filecount).name,'r'); % Open this file
    frewind(fid);
    f=find(d(filecount).name=='.');
    outfile = ([d(filecount).name(1:f-1),'proc']); % Define name of output file
    textoutfile = ([d(filecount).name(1:f-1),'proc.txt']); % Define name of output file
    record_no = 0;

    fprintf(fid,'%s 	 %5.2f 	 %5.2f 	 %1.3f 	 %6.2f 	 %2.2f 	 %6.2f 	 %5.2f 	 %6.2f 	 %6.2f', infile, xfit(1), xfit(2), fvfm, xfit(3), xfit(4),...
        taufit, fmMT, tauPQ_fit, PARint);
    fprintf(fid,'
');
    else                                        %after first access just write data
        fid = fopen(outfile,'a');
        fprintf(fid,'%s 	 %5.2f 	 %5.2f 	 %1.3f 	 %6.2f 	 %2.2f 	 %6.2f 	 %5.2f 	 %6.2f 	 %6.2f', infile, xfit(1), xfit(2),fvfm, xfit(3), xfit(4),...
            taufit, fmMT, tauPQ_fit, PARint);
        fprintf(fid,'
');
    end
    fclose(fid);

    paramsfit = [xfit(1) xfit(2) xfit(3) xfit(4) taufit fmMT tauPQ_fit PARint];
    return;
% ================ Read in header information, PAR etc. ===============

line = fgets(fid); % Read in headers
line = fgets(fid); % This line is the time
time_secs(filecount) = str2num(line(end-9:end));
line = fgets(fid); % Read in headers: LED
line = fgets(fid); % Read in headers: Gain
Gain = str2num(line(6:9));
line = fgets(fid); % Read in headers: Sample delay
line = fgets(fid); % Read in headers: Number of samples

line = fgets(fid);, STF = str2num(line(end-4:end)); % Duration of ST saturation sequence
line = fgets(fid);, STRP = str2num(line(end-4:end)); % Number of points in ST relaxation sequence
line = fgets(fid);, STRI = str2num(line(end-4:end)); % Interval between first ST relaxation measurement
line = fgets(fid); % MTF length (ms)

line = fgets(fid);, MTF = str2num(line(end-4:end)); % Number of points in MT saturation sequence
line = fgets(fid);, MTRP = str2num(line(end-4:end)); % Number of points in MT relaxation sequence
line = fgets(fid);, MTRI = str2num(line(end-4:end)); % Interval between first MT relaxation measurement
line = fgets(fid);
line = fgets(fid);

if line(end-4:end-2) == 'Off' % Read in the PAR data iff Actinic source was used
   PAR(filecount) = 0;
   for i = 1:5
     line = fgets(fid);
   end
else if line(end-4:end-2) == 'On'
   line = fgets(fid);
   f=find(line==':');
   PAR(filecount) = str2num(line(f+1:end));
   for i = 1:8
     line = fgets(fid);
   end
end

%==================================================================================================

% ========= Read in Data ========

while (feof(fid) == 0) % while not end of input file.....

   line = fgets(fid); % start loading data
   line = str2num(line);
   record_no = record_no + 1;
   Time(record_no) = line(2); % in microseconds
   Ex(record_no) = line(3);
   Yield(record_no) = line(4);
end % loading data

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% Correct yields to absolute values

if Gain == 2200
    Yield = Yield./283;
end
if Gain == 2000
    Yield = Yield./195;
end
if Gain == 1800
    Yield = Yield./123;
end
if Gain == 1600
    Yield = Yield./78.3;
end
if Gain == 1400
    Yield = Yield./50;
end

Yield_STS = Yield(1:STF); % Break records into separate phases
Yield_STR = Yield(STF+1:STF+STRP);
Yield_MTS = Yield(STF+STRP+1:STF+STRP+MTF);
Yield_MTR = Yield(STF+STRP+MTF+1:STF+STRP+MTF+MTRP);

Time_STS = Time(1:STF);
Time_STR = Time(STF+1:STF+STRP)-Time(STF);
Time_MTS = Time(STF+STRP+1:STF+STRP+MTF);
Time_MTR = Time(STF+STRP+MTF+1:STF+STRP+MTF+MTRP);

Yield_STS_no3 = Yield_STS(4:end); % Reduced ST section ignoring first 3 micro secs

% =============== Perform fitting

PFD_STS = ones(size(Time_STS)).*LED_intensity.*6.023e23/1e32;
PFD_STS_no3 = [sum(PFD_STS(1:4)) PFD_STS(5:end)]; % Reduced PFD for ST section ignoring first 3 micro secs

sparams_saturate = [Yield_STS(1) Yield_STS(end) 300 0.5]; % Set initial guesses for relaxation parameter fits
sparams_relax_ST = [200 2000 20000 0.3 0.3 0.3 Yield_STR(end)];
Yield_STR(1)]; % Set initial guesses for relaxation parameter fits
sparams_relax_MT = [2000 20000 200000 0.3 0.3 0.3 Yield_MTR(end)];
Yield_MTR(1)]; % Set initial guesses for relaxation parameter fits

[f,sfitparams_STS,kvg,iter,corp,covp_exp,covr,stdresid,Z,r2]=...
nleasqr(PFD_STS_no3',Yield_STS_no3',sparams_saturate,'Saturation_function' ); % Perform fitting for STS kinetic
[f,sfitparams_STS_nop,kvg,iter,corp,covp_nop,covr,stdresid,Z,r2]=...
nlleasqr(PFD_STS_no3',Yield_STS_no3',sparams_saturate(1:3),'Saturation_function_nop'); % Perform fitting for STS kinetic without p

[f,sfitparams_STR,kvg,iter,corp,covp_exp,covr,stdresid,Z,r2]=...
nlleasqr(Time_STR',Yield_STR',sparams_relax_ST,'Relaxation_function'); % Perform fitting for STR kinetic

[f,sfitparams_MTR,kvg,iter,corp,covp_exp,covr,stdresid,Z,r2]=...
nlleasqr(Time_MTR',Yield_MTR',sparams_relax_MT,'Relaxation_function'); % Perform fitting for MTR kinetic

Fm_MT(filecount) = max(Yield_MTS);

% pause

% ======= Copy outputs from fitting routine into vectors
===================

Fo_nop(filecount)  = sfitparams_STS_nop(1);
Fm_nop(filecount)  = sfitparams_STS_nop(2);
Sig_nop(filecount) = sfitparams_STS_nop(3);
Fo(filecount)      = sfitparams_STS(1);
Fm(filecount)      = sfitparams_STS(2);
Sig(filecount)     = sfitparams_STS(3);
p(filecount)       = sfitparams_STS(4);

erFo_nop(filecount) = covp_nop(1,1)^0.5;
erFm_nop(filecount) = covp_nop(2,2)^0.5;

per_erFo_nop(filecount) = 100.*erFo_nop(filecount)/Fo_nop(filecount);
per_erFm_nop(filecount) = 100.*erFm_nop(filecount)/Fm_nop(filecount);

tau1_ST(filecount) = sfitparams_STR(1);
tau2_ST(filecount) = sfitparams_STR(2);
tau3_ST(filecount) = sfitparams_STR(3);
alp1_ST(filecount) = sfitparams_STR(4);
alp2_ST(filecount) = sfitparams_STR(5);
alp3_ST(filecount) = sfitparams_STR(6);
Fo_relax_ST(filecount) = sfitparams_STR(7);
Fm_relax_ST(filecount) = sfitparams_STR(8);

tau1_MT(filecount) = sfitparams_MTR(1);
tau2_MT(filecount) = sfitparams_MTR(2);
tau3_MT(filecount) = sfitparams_MTR(3);
alp1_MT(filecount) = sfitparams_MTR(4);
alp2_MT(filecount) = sfitparams_MTR(5);
alp3_MT(filecount) = sfitparams_MTR(6);
Fo_relax_MT(filecount) = sfitparams_MTR(7);
Fm_relax_MT(filecount) = sfitparams_MTR(8);
Average_F(filecount) = mean(Yield(2:STF));
Fo_rough(filecount) = mean(Yield(3:6));
Fm_rough(filecount) = mean(Yield(STF-2:STF));
GAIN(filecount) = Gain;

% ==== This is an attempt to calculate rate of PQ pool reduction,
% ==== doesn't work very well yet!
%  Delta_Fm_ST_MT(filecount) = Fm_MT(filecount) - Fm_nop(filecount);
%  T_half_ST_MT(filecount) = min(Time_MTS(find(Yield_MTS >=
(Fm_nop(filecount) + Fm_MT(filecount))./2)));

% ====== Plot fits ======
clf;
subplot(311), plot(Yield,'r','LineWidth',2)
xlabel('Measurement no.')
ylabel('F (a.u.)')

out = Saturation_function_nop(PFD_STS_no3, sfitparams_STS_nop); %
Predicted fit for single turnover relaxation
outp = Saturation_function(PFD_STS_no3, sfitparams_STS); % Predicted
fit for single turnover relaxation with p
subplot(323), plot(out,'b','LineWidth',3)
hold on,
plot(outp,'g','LineWidth',3)
% plot(Yield_STS,'ok','MarkerFaceColor',[0.7 0.7 0.7],'MarkerSize',4)
plot([1:length(Yield_STS)-3],Yield_STS_no3,'ok','MarkerFaceColor','w','MarkerSize',4)
xlabel('Measurement no.')
ylabel('F (a.u.)')

out = Relaxation_function(Time_STR, sfitparams_STR); % Predicted fit
for single turnover relaxation
subplot(324), semilogx(Time_STR,out,'r','LineWidth',3)
hold on,
plot(Time_STR,Yield_STR,'ok','MarkerFaceColor','w','MarkerSize',4)
xlim([10 5e6])
xlabel('Time (\mus)')
ylabel('F (a.u.)')

subplot(325), plot(Time_MTS,Yield_MTS,'b')
hold on
plot(Time_MTS,ones(size(Time_MTS)).*Fm_nop(filecount),'k:')
plot(Time_MTS,ones(size(Time_MTS)).*Fo_nop(filecount),'k:')
plot(Time_MTS,ones(size(Time_MTS)).*Fm_MT(filecount),'k:')
% plot(Time_MTS,ones(size(Time_MTS)).*(Fm_nop(filecount)+Delta_Fm_ST_MT(filecount))./2),'r:')
plot(T_half_ST_MT(filecount).*ones(1,2),[Fo_nop(filecount)
Fm_nop(filecount)+Delta_Fm_ST_MT(filecount))./2],'r:')
out = Relaxation_function(Time_MTR, sfitparams_MTR); % Predicted fit
for single turnover relaxation
xlabel('Time (\mus)')
ylabel('F (a.u.)')

subplot(326), semilogx(Time_MTR,out,'r','LineWidth',3)
hold on,
plot(Time_MTR,Yield_MTR,'ok','MarkerFaceColor','w','MarkerSize',4)
xlim([10 5e6])
xlabel('Time (\mus)')
ylabel('F (a.u.)')
pause%(0.01);

% ===============
fclose(fid); % close this input file
% ====== Save the evaluated parameters to the output file

eval(['save ', outfile,' PAR Fo_nop Fm_nop per_erFo_nop per_erFm_nop Sig_nop Fo Fm Sig p tau1_ST tau2_ST tau3_ST alpha1_ST alpha2_ST alpha3_ST tau1_MT tau2_MT tau3_MT alpha1_MT alpha2_MT alpha3_MT Fm_MT Fo_relax_ST Fm_relax_ST Fo_relax_MT Fm_relax_MT Average_F'])

end % of all files for this filename

alpha_sum_ST = alpha1_ST + alpha2_ST + alpha3_ST; % Check whether sum of alpha terms equals 1
alpha_sum_MT = alpha1_MT + alpha2_MT + alpha3_MT; % Check whether sum of alpha terms equals 1

% out = [Fo_nop(2:2:end)' Fm_nop(2:2:end)' Fm_MT(2:2:end)' Fm_nop(1:2:end)' Fm_nop(1:2:end)' Fm_MT(1:2:end)'] % For copying to text/excel
out = [Average_F' Fo_nop' Fm_nop' per_erFo_nop' per_erFm_nop' Fo_rough' Fm_rough' Fm_MT' Sig_nop' tau1_ST' tau2_ST' tau3_ST' alpha1_ST' alpha2_ST' alpha3_ST' tau1_MT' tau2_MT' tau3_MT' alpha1_MT' alpha2_MT' alpha3_MT'] % For writing to text out file

fid = fopen(textoutfile,'w');
fprintf(fid,' AvgF Fo Fm ErrorFo ErrorFm Fo_rough Fm_rough FmMT SigmaPSII tau1_ST tau2_ST tau3_ST alpha1_ST alpha2_ST alpha3_ST tau1_MT tau2_MT tau3_MT alpha1_MT alpha2_MT alpha3_MT
');
fprintf(fid,'%6.3f %6.3f %6.3f %6.3f %6.3f %6.3f %6.3f %6.3f %6.2f %6.3f %6.3f %6.3f %6.3f %6.3f %6.3f %6.3f %6.3f %6.3f %6.3f %6.3f %6.3f
',out');
close(fid);

nlleasqr

function

[f,p,kvg,iter,corp,covp,covr,stdresid,Z,r2]=nlleasqr(x,y,pin,func,stol,niter,wt,dp,dfdp,options)

%function[f,p,kvg,iter,corp,covp,covr,stdresid,Z,r2]=
  % nlleasqr(x,y,pin,func,stol,niter,wt,dp,dfdp,options))

% Version 3. beta
% Levenberg-Marquardt nonlinear regression of f(x,p) to y(x), where:
% x=vec or mat of indep variables, 1 row/observation: x=[x0 x1....xm]
% y=vec of obs values, same no. of rows as x.
% wt=vec(dim=1 or length(x)) of statistical weights. These should be set
% to be proportional to (sqrts of var(y))^-1; (That is, the covaraince
% matrix of the data is assumed to be proportional to diagonal with
% equal to (wt.^2)^-1. The constant of proportionality will be
% default=1.
% pin=vector of initial parameters to be adjusted by leasqr.
% dp=fractional incr of p for numerical partials,default=.001*ones(size(pin))
% Note: dp(j)=0 holds p(j) fixed i.e. leasqr wont change initial guess:
% pin(j)
% func=name of function in quotes,of the form y=f(x,p)
% dfdp=name of partials M-file in quotes default is prt=dfdp(x,f,p,dp,func)
% stol=scalar tolerances on fractional improvement in ss, default stol=.0001
% niter=scalar max no. of iterations, default = 20
% options=matrix of n rows (same number of rows as pin) containing
% column 1: desired fractional precision in parameter estimates.
%   Iterations are terminated if change in parameter vector (chg) on two
%   consecutive iterations is less than their corresponding elements
%   in options(:,1).  [ie. all(abs(chg*current parm est) < options(:,1))
%   on two consecutive iterations.], default = zeros().
% column 2: maximum fractional step change in parameter vector.
%   Fractional change in elements of parameter vector is constrained to
% be
%   at most options(:,2) between successive iterations.
%   [ie. abs(chg(i))=abs(min([chg(i) options(i,2)*current param
% estimate]]),
%   default = Inf*ones().)
% OUTPUT VARIABLES
% f=vec function values computed in function func.
% p=vec trial or final parameters. i.e, the solution.
% kvg=scalar: =1 if convergence, =0 otherwise.
% iter=scalar no. of interations used.
% corp= correlation matrix for parameters
% covp= covariance matrix of the parameters
% covr = diag(covariance matrix of the residuals)
% stdresid= standardized residuals
% Z= matrix that defines confidence region
% r2= coefficient of multiple determination
% ():= optional parameters
% ss=scalar sum of squares=sum-over-i(wt(i)*(y(i)-f(i)))^2.

% All Zero guesses not acceptable
% Richard I. Shrager (301)-496-1122
% Modified by A.Jutan (519)-679-2111
% Modified by Ray Muzic 14-Jul-1992
%      1) add maxstep feature for limiting changes in parameter estimates
%      at each step.
%      2) remove forced columnization of x (x=x(:)) at beginning. x could
% be
%      a matrix with the ith row of containing values of the
% independent variables at the ith observation.
%      3) add verbose option
%      4) add optional return arguments covp, stdresid, chi2
%      5) revise estimates of corp, stdev
% Modified by Ray Muzic 11-Oct-1992
%      1) revise estimate of Vy.  remove chi2, add Z as return values
% Modified by Ray Muzic 7-Jan-1994
%      1) Replace ones(x) with a construct that is compatible with
% versions
%      newer and older than v 4.1.
%      2) Added global declaration of verbose (needed for newer than v4.x)
%      3) Replace return value var, the variance of the residuals with
covr,
%      the covariance matrix of the residuals.
%      4) Introduce options as 10th input argument. Include
%      convergence criteria and maxstep in it.
%      5) Correct calculation of xtx which affects covariance estimate.
%      6) Eliminate stdev (estimate of standard deviation of parameter
% estimates) from the return values. The covp is a much more
meaningful expression of precision because it specifies a confidence region in contrast to a confidence interval. If needed, however, stdev may be calculated as stdev=sqrt(diag(covp)).

7) Change the order of the return values to a more logical order.
8) Change to more efficient algorithm of Bard for selecting epsL.

References:

% set default args

plotcmd='plot(x(:,1),y,''o'',x(:,1),f,''+''); shg';
if (sscanf(version,'%f') >= 4),
    global verbose
    plotcmd='plot(x(:,1),y,''o'',x(:,1),f,''+''); figure(gcf)';
end;

if(exist('verbose')~=1), verbose=0; end;
if (nargin <= 8), dfdp='dfdp'; end;
if (nargin <= 7), dp=.001*(pin*0+1); end; %DT
if (nargin <= 6), wt=1.0; end;
if (nargin <= 5), niter=20; end;
if (nargin <= 4), stol=.0001; end;

y=y(:); wt=wt(:); pin=pin(:); dp=dp(:); %change all vectors to columns
% check data vectors- same length?
m=length(y); n=length(pin); p=pin;
if m1~m ,error('input(x)/output(y) data must have same number of rows ') ,end;

if (nargin <= 9),
    options=[zeros(n,1) Inf*ones(n,1)];
    nor = n; noc = 2;
else
    [nor noc]=size(options);
    if (nor ~= n),
        error('options and parameter matrices must have same number of rows'),
        end;
    if (noc ~= 2),
        options={options(noc,1) Inf*ones(noc,1)};
    end;
end;
pprec=options(:,1);
maxstep=options(:,2);

% set up for iterations
f=feval(func,x,p); fbest=f; pbest=p;
r=wt.*(y-f);
sbest=r'*r;
nrm=zeros(n,1);
chgprev=Inf*ones(n,1);
kvg=0;
epsLlast=1;
epstab=[.1 1 1e2 1e4 1e6];

% do iterations

for iter=1:niter,
    pprev=pbest;
    prt=feval(dfdp,x,fbest,pprev,dp,func);
    r=wt.*(y-fbest);
    sprev=sbest;
    sgoal=(1-stol)*sprev;
    for j=1:n,
        if dp(j)==0,
            nrm(j)=0;
        else
            prt(:,j)=wt.*prt(:,j);
            nrm(j)=prt(:,j)'*prt(:,j);
            if nrm(j)>0,
                nrm(j)=1/sqrt(nrm(j));
            end;
        end;
        pr(:,j)=nrm(j)*prt(:,j);
    end;
    [prt,s,v]=svd(prt,0);
    s=diag(s);
    g=prt'*r;
    for j=1:length(epstab),
        epsL = max(epsLlast*epstab(j),1e-7);
        se=sqrt((s.*s)+epsL);
        gse=g./se;
        chg=((v*gse).*nrm);
        %   check the change constraints and apply as necessary
        ochg=chg;
        for iii=1:n,
            if (maxstep(iii)==Inf), break; end;
            chg(iii)=max(chg(iii),-abs(maxstep(iii)*pprev(iii)));
            chg(iii)=min(chg(iii),abs(maxstep(iii)*pprev(iii)));
        end;
        if (verbose & any(ochg ~= chg)),
            disp([\'Change in parameter(s): ' ...
            sprintf('%d ',find(ochg ~= chg)) \'were constrained\']);
        end;
        aprec=abs(pprec*pbest); %---
        if (any(abs(chg) > 0.1*aprec)),%--- % only worth evaluating function
            p=chg+pprev;
            % there is some non-miniscule change
            f=feval(func,x,p);
            r=wt.*(y-f);
            ss=r'*r;
            if ss<sbest,
                pbest=p;
                fbest=f;
                sbest=ss;
            end;
            if ss<sgoal,
                break;
            end;
        end;
    end;
end;
epsLlast = epsL;
if (verbose),
    eval(plotcmd);
end;
if ss<eps,
    break;
end
aprec=abs(pprec.*pbest);
%
[aprec chg chgprev]
if (all(abs(chg) < aprec) & all(abs(chgprev) < aprec)),
    kvg=1;
    if (verbose),
        fprintf('Parameter changes converged to specified precision\n');
    end;
    break;
else
    chgprev=chg;
end;
if ss>sgoal,
    break;
end;
end;

% set return values
%
p=pbest;
f=fbest;
ss=sbest;
kvg=((ss>sgoal)|(ss<=eps)|kvg);
if kvg ~= 1 , disp(' CONVERGENCE NOT ACHIEVED! '), end;

% CALC VARIANCE COV MATRIX AND CORRELATION MATRIX OF PARAMETERS
% re-evaluate the Jacobian at optimal values
jac=feval(dfdp,x,f,p,dp,func);
msk = dp ~= 0;
n = sum(msk);       % reduce n to equal number of estimated parameters
jac = jac(:, msk);   % use only fitted parameters

% following section is Ray Muzic's estimate for covariance and correlation
% assing covariance of data is a diagonal matrix proportional to
% diag(1/wt.^2).
% cov matrix of data est. from Bard Eq. 7-5-13, and Row 1 Table 5.1
Qinv=diag(wt.*wt);
Q=diag((0*wt+1)./(wt.^2));
%[nrw ncw]=size(wt);
%Q=ones(nrw,ncw)./wt; Q=diag(Q.*Q);
resid=y-f;
covr=resid'*Qinv*resid/(m-n);  %covariance of residuals
Vy=1/(1-n/m)*covr;  % Eq. 7-13-22, Bard  %covariance of the data
covr=diag(covr);
Z=((m-n)*jac'*Qinv*jac)/(n*resid'*Qinv*resid);  %for compact storage
stdresid=resid./sqrt(diag(Vy));
jtgjinv=inv(jac'*Qinv*jac);
covp=jtgjinv*jac'*Qinv*Vy*Qinv*jac*jtgjinv;  % Eq. 7-5-13, Bard  %cov of parm est
for k=1:n,
    for j=k:n,
        corp(k,j)=covp(k,j)/sqrt(abs(covp(k,k)*covp(j,j)));
    end;
end;
corp(j,k)=corp(k,j);
end;
end;

%%% alt. est. of cov. mat. of parm.: (Delforge, Circulation, 82:1494-1504, 1990
%%% disp('Alternate estimate of cov. of param. est."
% acovp=resid'*Qinv*resid/(m-n)*jtgginv

% Calculate R^2 (Ref Draper & Smith p.46)
% r=corrcoef(y,f);
r2=r(1,2).^2;

% if someone has asked for it, let them have it
% if (verbose),
eval(plotcmd);
disp(' Least Squares Estimates of Parameters"
disp(p')
disp(' Correlation matrix of parameters estimated"
disp(corp)
disp(' Covariance matrix of Residuals ')
disp(covr)
disp( 'Correlation Coefficient R^2'

disp(r2)
sprintf('95\% conf region: F(0.05)(%.0f,%.0f)\geq '
delta_pvec''*Z*delta_pvec',n,m-n)
Z
end;

% A modified version of Levenberg-Marquardt
% Non-Linear Regression program previously submitted by R.Schrager.
% This version corrects an error in that version and also provides
% an easier to use version with automatic numerical calculation of
% the Jacobian Matrix. In addition, this version calculates statistics
% such as correlation, etc....
% %
% Version 3 Notes
% Errors in the original version submitted by Shragter (now called version 1)
% and the improved version of Jutan (now called version 2) have been
% corrected.
% Additional features, statistical tests, and documentation have also been
% included along with an example of usage. BEWARE: Some the the input and
% output arguments were changed from the previous version.
% $ Ray Muzic  rfm2@ds2.uh.cwru.edu
% $ Arthur Jutan  jutan@charon.engga.uwo.ca

Saturation_function

% FRRF Saturation curve for FIRE_processing (see Kolber et al. 1998)

function out = Saturation_function(pfd, params);

fo = params(1);
fm = params(2);
sig = params(3);
p = params(4);

out = zeros(1,length(pfd));
c = zeros(1,length(pfd));

c(1) = pfd(1) * sig;
for i = 2:length(out),
c(i) = c(i-1) + pfd(i) * sig * (1 - c(i-1))/(1 - p * c(i-1));
end;

out = fo + (fm - fo)*c*(1-p)./(1-c*p) ;

out = out';

dfdp

function prt=dfdp(x,f,p,dp,func)
% numerical partial derivatives (Jacobian) df/dp for use with leasqr
% --------INPUT VARIABLES---------
% x=vec or matrix of indep var(used as arg to func) x=[x0 x1 ....]
% f=func(x,p) vector initialised by user before each call to dfdp
% p= vec of current parameter values
% dp= fractional increment of p for numerical derivatives
%   dp(j)>0 central differences calculated
%   dp(j)<0 one sided differences calculated
%   dp(j)=0 sets corresponding partials to zero; i.e. holds p(j) fixed
% func=string naming the function (.m) file
%   e.g. to calc Jacobian for funcion expsum
prt=dfdp(x,f,p,dp,'expsum')
%--------OUTPUT VARIABLES-------
% prt= Jacobian Matrix prt(i,j)=df(i)/dp(j)
%================================

m=length(x);n=length(p);      %dimensions
ps=p; prt=zeros(m,n);del=zeros(n,1);          % initialise Jacobian to Zero
for j=1:n
  del(j)=dp(j) .*p(j);    %cal delx=fract(dp)*param value(p)
  if p(j)==0
    del(j)=dp(j);     %if param=0 delx=fraction
  end
  p(j)=ps(j) + del(j);    
  if del(j)<0, f1=feval(func,x,p);
    if dp(j) < 0, prt(:,j)=(f1-f)./del(j);
  else
    p(j)=ps(j)- del(j);
    prt(:,j)=(f1-feval(func,x,p))./(2 .*del(j));
  end
  end
p(j)=ps(j);       %restore p(j)
end
return

Saturation -function_nop

% FRRF Saturation curve for FIRe_processing (see Kolber et al. 1998)

function out = Saturation_function_nop(pfd,params);

fo = params(1);
\[ \text{out} = \text{fo} + (\text{fm} - \text{fo}) \times (1 - \exp(-\text{sig} \times \text{cumsum(pfd)})) ; \]

**Relaxation_function**

\[
\text{function} \quad \text{out} = \text{Relaxation_function}(\text{Time\_Relax}, \text{params})
\]

\[
\text{tau1} = \text{params}(1);
\text{tau2} = \text{params}(2);
\text{tau3} = \text{params}(3);
\alpha_1 = \text{params}(4);
\alpha_2 = \text{params}(5);
\alpha_3 = \text{params}(6);
\text{fo\_relax} = \text{params}(7);
\text{fm\_relax} = \text{params}(8);
\]

\[
\text{out} = \text{fo\_relax} + (\text{fm\_relax} - \text{fo\_relax}) \times (\alpha_1 \times \exp(-\text{Time\_Relax}/\text{tau1}) + \alpha_2 \times \exp(-\text{Time\_Relax}/\text{tau2}) + \alpha_3 \times \exp(-\text{Time\_Relax}/\text{tau3})) ;
\]
Appendix D: Graphs depicting a) the northward leg of NO$_3$ for SANAE 48, 49 and the adjusted NO$_3$ values for SANAE 49, b) The northward leg of PO$_4$ for SANAE 48 and 49, and c) the NO$_3$:PO$_4$ ratio for the northward leg of SANAE 49.
Appendix E: Graph showing the NO$_3$:PO$_4$ ratios from SANAЕ 48 and SANAЕ 49 used to correct the NO$_4$ data on the northward leg.

\[ y = 12.621x + 1.9597 \]
\[ R^2 = 0.8004 \]

Appendix F: Pigments to species table.
Appendix G: Code used for statistical analysis in R regarding the influences on Fv/Fm and σPSII.
####  first try

data=read.csv("E://Rplaywithsilicate//ALLDATAwithsilicate6.csv",header=TRUE)
attach(data)
data=data[1:175,]
data

Fv.Fm.average

model3=lm(Fv.Fm.average~SiO4+PO4+N03+Chlorophyll+Temperature+Salinity+Oxygen+Fluorescence+PAR)
model3
summary(model3)

model4=lm(Sigma.average~SiO4+PO4+N03+Chlorophyll+Temperature+Salinity+Oxygen+Fluorescence+PAR)
model4
summary(model4)

#### depth data devision

depth1=data[1:30,]
end

---

**Appendix II:** Weather observations during experiments.
<table>
<thead>
<tr>
<th>Day</th>
<th>Time</th>
<th>Experiment A</th>
<th>Experiment B</th>
<th>Experiment C</th>
<th>Experiment D</th>
<th>Experiment E</th>
<th>Experiment F</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>20:00</td>
<td>Just starting to become sunny</td>
<td>sunny, beautiful - lots of wildlife - killer whales, humpbacks and leopard seals</td>
<td>overcast</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>22:00</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>00:00</td>
<td>cloudy</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
</tr>
<tr>
<td></td>
<td>04:00</td>
<td>Full sunshine, beautiful!</td>
<td>dark, cloudy &amp; misty</td>
<td>high productivity 0.8, started raining just after station</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>06:00</td>
<td>sunshine</td>
<td>getting light on horizon, cloudy, mist cleared</td>
<td>light, 6 SCI (Toaotiy), high cloud, light rain</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>08:00</td>
<td>sunshine</td>
<td>miserable, overcast day</td>
<td>cloudy, clearer</td>
<td>cloudy cover, drizzle</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>10:00</td>
<td>overcast</td>
<td>miserable, overcast day</td>
<td>cloudy</td>
<td>warm</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>12:00</td>
<td>overcast</td>
<td>little windy, dark, cold. Chl-a up to 1.5 on fluorometer! Huge bloom.</td>
<td>sunny</td>
<td>beautiful - lots of wildlife - killer whales, humpbacks and leopard seals</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>14:00</td>
<td>overcast, cold</td>
<td>cloudy</td>
<td>miserable, overcast day</td>
<td>misty, calm</td>
<td>cloudy</td>
<td></td>
</tr>
<tr>
<td></td>
<td>16:00</td>
<td>overcast, cold</td>
<td>light, a little windy, overcast</td>
<td>miserable day with heavy cloud, wind, snow</td>
<td>overcast, cold</td>
<td>sunny</td>
<td></td>
</tr>
<tr>
<td></td>
<td>18:00</td>
<td>overcast, cold</td>
<td>overcast, cold</td>
<td>overcast</td>
<td>sunny</td>
<td></td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>20:00</td>
<td>Cloudy, windy, light swell</td>
<td>sunny</td>
<td>sunny</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>22:00</td>
<td>No ice. Overcast</td>
<td>overcast</td>
<td>near dark, overcast</td>
<td>cloudy</td>
<td>sunny</td>
<td></td>
</tr>
<tr>
<td>00:00</td>
<td>Start of weddell ACC front Rapid increase in Temp and decrease in CO2</td>
<td>Overcast, windy</td>
<td>overcast</td>
<td>overcast</td>
<td>dark, increasing resolution</td>
<td>it was dark</td>
<td></td>
</tr>
<tr>
<td>02:00</td>
<td>overcast</td>
<td>overcast</td>
<td>overcast</td>
<td>overcast</td>
<td>Warm &amp; dark</td>
<td>sunny</td>
<td></td>
</tr>
<tr>
<td></td>
<td>04:00</td>
<td>overcast</td>
<td>sunny</td>
<td>sunny</td>
<td>sunshine</td>
<td>overcast, temperature dropped</td>
<td></td>
</tr>
<tr>
<td></td>
<td>06:00</td>
<td>Stormy, snowing, almost white out</td>
<td>sunny</td>
<td>sunny</td>
<td>high productivity 0.8, started raining just after station</td>
<td>overcast, temperature dropped</td>
<td></td>
</tr>
<tr>
<td></td>
<td>08:00</td>
<td>Stormy, sleet</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
<td></td>
</tr>
<tr>
<td></td>
<td>10:00</td>
<td>Stormy</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
<td></td>
</tr>
<tr>
<td></td>
<td>12:00</td>
<td>Cold, stormy</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
<td></td>
</tr>
<tr>
<td></td>
<td>14:00</td>
<td>Cold</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
<td></td>
</tr>
<tr>
<td></td>
<td>16:00</td>
<td>Cold</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
<td></td>
</tr>
<tr>
<td></td>
<td>18:00</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>20:00</td>
<td>Sun shining, wind, swell, white horses</td>
<td>sunny</td>
<td>sunny</td>
<td>dark, windy, productivity increasing, on the STF?</td>
<td>dark, windy, productivity increasing, on the STF?</td>
<td></td>
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<tr>
<td></td>
<td>22:00</td>
<td>Overcast</td>
<td>sunny</td>
<td>sunny</td>
<td>overcast, temperature dropped</td>
<td>sunny</td>
<td></td>
</tr>
<tr>
<td></td>
<td>00:00</td>
<td>dark</td>
<td>sunny</td>
<td>sunny</td>
<td>dark, overcast</td>
<td>sunny</td>
<td></td>
</tr>
<tr>
<td></td>
<td>02:00</td>
<td>cold, windy, dark</td>
<td>sunny</td>
<td>sunny</td>
<td>it was dark</td>
<td>6m swell, dark, windy, sparklingSusan, floe clouds</td>
<td></td>
</tr>
<tr>
<td></td>
<td>04:00</td>
<td>Overcast, light sleet and wind</td>
<td>sunny</td>
<td>sunny</td>
<td>clear with very bright stars</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>06:00</td>
<td>Sunny</td>
<td>clear with very bright stars</td>
<td>getting light, windy, partly cloudy</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>08:00</td>
<td>Wind perfect for upwelling</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
<td></td>
</tr>
<tr>
<td></td>
<td>10:00</td>
<td>Sunny</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
<td></td>
</tr>
<tr>
<td></td>
<td>12:00</td>
<td>Calm, clear</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
<td></td>
</tr>
<tr>
<td></td>
<td>14:00</td>
<td>Calm</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
<td></td>
</tr>
<tr>
<td></td>
<td>16:00</td>
<td>Calm, clear</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
<td>sunny</td>
<td></td>
</tr>
<tr>
<td></td>
<td>20:00</td>
<td>Sun shining, wind, swell, white horses</td>
<td>sunny</td>
<td>sunny, clear, beautiful, cold</td>
<td>sunny</td>
<td>sunny, beautiful - lots of wildlife - killer whales, humpbacks and leopard seals</td>
<td></td>
</tr>
</tbody>
</table>

Legend:
- **ACC Front**: Antarctic Circumpolar Current Front
- **STF**: South Tasman Front
- **SCI**: Surface Current Index
- **Upwelling**: Process where nutrients and oxygen are brought to the surface of the ocean
- **Productivity**: Ability of an ecosystem to produce biomass
- **Toaotiy**: Pacific Ocean Current
- **Floe**: A large piece of ice
- **Susan**: A term used in the context of the Southern Ocean
- **Flux Clouds**: Clouds that indicate the presence of winds
- **Chl-a**: Chlorophyll-a, a pigment produced by phytoplankton
- **Jupiter**: A large gas planet in the solar system
- **Scorpios**: The constellation of Scorpions, known for its tail-like appearance
- **Sunrise, Sunset**: Refers to the time of day when the sun is at its lowest point in the sky.
<table>
<thead>
<tr>
<th>Time</th>
<th>Weather Conditions</th>
<th>Visibility</th>
<th>Remarks</th>
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<tbody>
<tr>
<td>00.00</td>
<td>Clearing</td>
<td>sunny</td>
<td>sunny</td>
</tr>
<tr>
<td>02.00</td>
<td>Cloudy, calmish, a little wind</td>
<td>sunny</td>
<td>windy, a sky full of sparkles</td>
</tr>
<tr>
<td>04.00</td>
<td>Cloudy, calm, Icebergs against</td>
<td>sunny</td>
<td>6m swell, dark, windy, sparkling ocean, flaw clouds</td>
</tr>
<tr>
<td>06.00</td>
<td>Overcast, cold, misty</td>
<td>sunny</td>
<td>getting light, windy, partly cloudy</td>
</tr>
<tr>
<td>08.00</td>
<td>Overcast, cold</td>
<td>sunny</td>
<td>weather starting to turn, ice on deck</td>
</tr>
<tr>
<td>12.00</td>
<td>Overcast, cold, windy</td>
<td>sunny</td>
<td>horses cantering</td>
</tr>
<tr>
<td>16.00</td>
<td>Overcast, cold, windy</td>
<td>sunny</td>
<td>sunny</td>
</tr>
<tr>
<td>18.00</td>
<td>Overcast, cold, windy</td>
<td>sunny</td>
<td>sunny</td>
</tr>
<tr>
<td>20.00</td>
<td>Overcast, cold, still light</td>
<td>sunny</td>
<td>sunny</td>
</tr>
<tr>
<td>02.00</td>
<td>Cloud cover, still light</td>
<td>sunny</td>
<td>overcast, snowing</td>
</tr>
<tr>
<td>04.00</td>
<td>Cloud cover, dark, clouds dropping</td>
<td>sunny</td>
<td>overcast, snowing</td>
</tr>
<tr>
<td>06.00</td>
<td>Cloudy, calm, light wind</td>
<td>sunny</td>
<td>overcast, snowing</td>
</tr>
<tr>
<td>08.00</td>
<td>Cloudy, calm, light wind</td>
<td>sunny</td>
<td>overcast, snowing</td>
</tr>
<tr>
<td>10.00</td>
<td>Overcast</td>
<td>sunny</td>
<td>overcast, snowing</td>
</tr>
<tr>
<td>14.00</td>
<td>Overcast</td>
<td>sunny</td>
<td>overcast, snowing</td>
</tr>
<tr>
<td>16.00</td>
<td>Taken as leaving South Georgia, cloudy &amp; Hailing</td>
<td>sunny</td>
<td>overcast, snowing</td>
</tr>
<tr>
<td>18.00</td>
<td>Overcast</td>
<td>sunny</td>
<td>overcast, snowing</td>
</tr>
<tr>
<td>20.00</td>
<td>Overcast</td>
<td>sunny</td>
<td>just starting to become sunny</td>
</tr>
<tr>
<td>22.00</td>
<td>Overcast, dark, cloudy</td>
<td>sunny</td>
<td>overcast, snowing</td>
</tr>
<tr>
<td>02.00</td>
<td>Overcast, cold, rainy</td>
<td>sunny</td>
<td>Full sunshine, beautiful!</td>
</tr>
<tr>
<td>04.00</td>
<td>Overcast</td>
<td>sunny</td>
<td>sunny</td>
</tr>
<tr>
<td>06.00</td>
<td>Overcast</td>
<td>sunny</td>
<td>sunny</td>
</tr>
<tr>
<td>08.00</td>
<td>Overcast</td>
<td>sunny</td>
<td>sunny</td>
</tr>
</tbody>
</table>
Appendix I: The nutrient results of experiment B.
**Appendix J: Critique and recommendations on the bio-assay nutrient methods**

**Critique**

*Phosphate*

Variation with PO$_4$ can be explained by the lack of chloroform treatment before freezing. Without this treatment marked changes in the level of inorganic PO$_4$ can occur (Gilmartin, 1967). These changes may be seen through increases due to ‘bacterial or enzymatic decomposition of organic phosphorus’ or decreases from either the ‘utilization by growing bacteria and plankton or by adsorption on detritus or sample bottle walls, or both’ (Gilmartin, 1967). Furthermore, analysis of frozen PO$_4$ data should occur within 4 months of collection to prevent the steady decrease in concentration (Clementson and Wayte, 1992) that occurs after this time. The experimental data from SANAE 50 began to be analysed after 6 months due to available laboratory and personal time of the person charged with analysing these nutrients.

*Silicate*

Variation with SiO$_4$ can be explained through two experimental errors that occurred in the process. One being a straight experimental error of on board filtering of all samples through a Whatman GF/F (made intrinsically of SiO$_4$). This could unexpectedly raise sample values depending on the amount of SiO$_4$ transferred from the filter. Secondly, Zhange and Ortner (1998) discovered that the thawing process of water samples is important in the recovery of the intrinsic value of SiO$_4$. Their recommendations, four days thawing in a 4°C freezer, for the 100% recovery of SiO$_4$ was not followed. This could lead to lower values of SiO$_4$. 


Nitrate

With regard to the NO$_3$ data, consultation with various international experts both at NOC and in South Africa, confirm that sometimes this just happens with frozen samples. The length of time for sample storage (6 months – year) was probably a largely contributing factor (Kremling and Wenck, 1986).

Recommendations

Though straight freezing and thawing is fine in oligotrophic regions (Dore et al., 1996), the results of this experiment along with experiences of a similar nature by other scientists suggest that it is not sufficient in the Southern Ocean. Hence I recommend that the processes recommended by Zhange and Ortner (1998) and Gilmartin (1967) and for SiO$_4$ and PO$_4$ respectively should be investigated and implemented in future cruises of this nature. If possible nutrients should be run on board the ship to prevent these complications. If this is not possible then they should be run immediately upon return to land.

ARCHER, D. 2009. The long Thaw: How humans are changing the next 100,000 years of Earth’s climate. Princeton University Press, United States of America


http://www.ieagreen.org.uk/prghgt42.htm (1 of 11) 13 July 2003


Assessment Report of the Intergovernmental Panel on Climate Change (S. Solomon, D. Qin, M. Manning, et al., [editors]). Cambridge University Press, Cambridge, United Kingdom and New York, NY, USA


http://www.R-project.org


RINTOUL et al., (in press) [http://www.sciencedaily.com/releases/2012/05/120521104635.htm](http://www.sciencedaily.com/releases/2012/05/120521104635.htm)


VOLK, T. and HOFFERT, M.I. 1985. Ocean carbon pumps: Analysis of relative strengths and efficiencies in ocean-driven atmospheric CO2 changes. In E.T. Sundquist and W.S. Broecker


Acts

Air Pollution Prevention Act in 1965 (South Africa)

Clean Air Act of 1956, updated in 1993 (Britain)

Clean Air Act of 1999 (Philippine)

Clean Air Act of 1963, Public Law (United States) 88-206, 77 United States Statutes at Large 392, 1963-12-17

Clean Air Act Extension of 1970, 84 United States Statutes at Large 1676, Public Law (United States) 91-604, 1970-12-31

Clean Air Act Amendments of 1977, Public Law (United States) 95-95, 91 United States Statutes at Large 685, 1977-08-07

Clean Air Act Amendments of 1990, Public Law (United States) 101-549, 104 United States Statutes at Large 2399, 1990-11-15

National Environmental Management: Air Quality Act 2004 (South Africa)