



Microbial food web dynamics during spring phytoplankton blooms in the naturally iron-fertilized Kerguelen area (Southern Ocean)

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Abstract. Microbial food web dynamics were determined during the onset of several spring phytoplankton blooms induced by natural iron fertilization off Kerguelen Island in the Southern Ocean (KEOPS2). The abundances of heterotrophic bacteria and heterotrophic nanoflagellates, bacterial heterotrophic production, bacterial respiration, and bacterial growth efficiency, were consistently higher in surface waters of the iron-fertilized sites than at the reference site in HNLC (high nutrient low chlorophyll) waters. The abundance of virus-like particles remained unchanged, but viral production increased by a factor of 6 in iron-fertilized waters. Bacterial heterotrophic production was significantly related to heterotrophic nanoflagellate abundance and viral production across all sites, with bacterial production explaining about 70 and 85 %, respectively, of the variance of each in the mixed layer (ML). Estimated rates of grazing and viral lysis, however, indicated that heterotrophic nanoflagellates accounted for a substantially higher loss of bacterial production (50 %) than viruses (11 %). Combining these results with

rates of primary production and export determined for the study area, a budget for the flow of carbon through the microbial food web and higher trophic levels during the early (KEOPS2) and the late phase (KEOPS1) of the Kerguelen bloom is provided.

1 Introduction

The Southern Ocean has a unique geography with major implications for the global ocean circulation and climate system. It is also the largest HNLC (high nutrient low chlorophyll) ocean where iron limits phytoplankton primary production, resulting in a large stock of unused major inorganic nutrients (Martin and Fitzwater, 1990). A pronounced shift to larger phytoplankton cells, in particular diatoms, has been generally observed upon natural (Blain et al., 2007; Pollard et al., 2009) or artificial (Boyd et al., 2007; Smetacek et al., 2012) iron additions. Natural and artificial iron fertilization

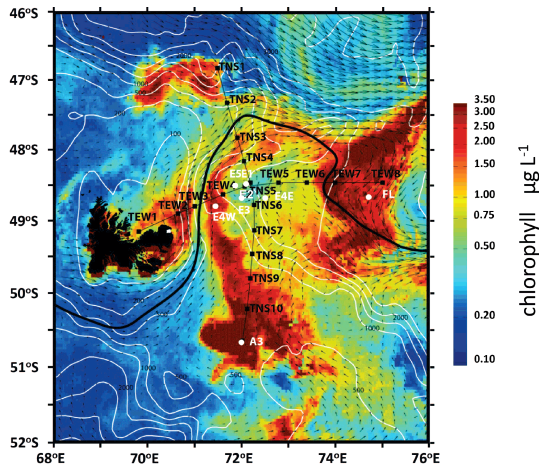


Figure 1. KEOPS2 study area from October to November 2011. Chl *a* (colour scale), surface velocity fields (arrows), the polar front (PF, black line), and the position of the different stations: stations 1–11 of the north–south transect (TNS); stations 1–9 of the east–west transect (TEW). The Chl *a* rich stations: A3, on the Kerguelen plateau visited twice; F-L and E-4W north and south of the polar front; and “E” stations sampled in a quasi-Lagrangian manner (E-1, E-2, E-3, E-4E, and E-5) within a complex meander south of the polar front. The reference HNLC station (R-2) is not shown as it is out of the area of the map (66.692743 °E longitude, 50.38954 °N latitude). Map is courtesy of Y. Park and colleagues. To note that, the chlorophyll content represented on the map corresponds to the last week of the KEOPS2 cruise.

studies have suggested regional variability in the ecosystem response to iron addition and in the carbon export (Blain et al., 2007; Pollard et al., 2009; Boyd et al., 2007). Although the underlying mechanisms are not fully understood, the variability in carbon export seems to be connected to the duration, the mode and magnitude of iron supply (De Baar et al., 2005; Boyd et al., 2007). The ratio of carbon exported from the surface layer to primary production is in part determined by microbial food web processes. Variable fractions of primary production are channelled through heterotrophic bacteria, and most of this phytoplankton-derived organic carbon is respired to carbon dioxide (CO₂). The question on how much carbon fixed by primary production is mineralized within the microbial food web, and what part of this particulate organic carbon is made available for higher trophic levels or export, is important in the present context.

For a better understanding of the role of the microbial food web for the fate of organic carbon, a comprehensive picture of the biomass, productivity, and mortality of heterotrophic bacteria is required. While the measurement of bacterial heterotrophic production (BP) in natural and artificial iron-fertilization studies has been regularly undertaken, measurements of bacterial respiration (BR) and growth efficiency have been rare (Obernosterer et al., 2008; Bonilla-Findji et al., 2008). The role and relative importance of bacteria in the carbon flow also needs to be completed by the

fate of BP through grazing by protists and viral lysis. Grazing channels part of the bacterial carbon to higher trophic levels, while viral induced mortality of bacteria is reported to lead to an increase in BR and finally to a decrease in carbon export (Middelboe and Lyck, 2002; Bonilla-Findji et al., 2008).

The oceanographic cruise KEOPS1 (Kerguelen Ocean and Plateau Study, January–February 2005) demonstrated that the 3-month spring phytoplankton bloom above the Kerguelen plateau is sustained from below by low level supplies of iron and other nutrients (Blain et al., 2007). KEOPS1 took place during the third month of the bloom, from its peak to its decline. Heterotrophic bacteria were more abundant in the bloom, and revealed substantially higher rates of production and respiration than in surrounding HNLC waters (Christaki et al., 2008). Heterotrophic bacteria processed a significant portion of primary production, with most of it being rapidly respired (Obernosterer et al., 2008). Furthermore, while heterotrophic nanoflagellates did not seem to efficiently control BP in the bloom (Christaki et al., 2008), viruses enhanced the role of bacteria as oxidizers of organic matter, hence as producers of CO₂ (Bonilla-Findji et al., 2008; Malits et al., 2014).

The KEOPS2 cruise aimed to complement the findings of KEOPS1 above the Kerguelen plateau, and gain new insights on the biogeochemistry and ecosystem response to iron fertilization by extending the study area to the offshore waters east of Kerguelen Island. The sampling strategy covered spatially diverse fertilized regions at early bloom stages (October–November 2011).

The main objective of the present study was to provide for the first time insight into seasonal dynamics of the microbial food web functioning in the context of natural Fe fertilization of the Southern Ocean. The major biogeochemical and biological parameters reported in this study are full depth profiles of microbial stocks (viruses, heterotrophic bacteria, and heterotrophic nanoflagellates), bacterial production, potential bacterial grazing, and viral lysis, while respiration was measured at selected depths in surface waters.

2 Methods

2.1 Sample collection

The present study was carried out during the KEOPS2 cruise from 15 October to 20 November 2011. Water samples for this study were collected at 10 stations along a north–south transect (TNS; stations 1, 2, 3, 4, 5, 6, 7, 8, 9, 10) and at seven stations along an east–west transect (TEW; Stations 1, 2, 3, 4, 5, 7, 8) (Fig. 1, Table 1), both covering the bloom area above and off the Kerguelen plateau, (Park et al., 2014) and providing an overview of the region. In addition to the “historical” A3 station situated on the Kerguelen plateau (Blain et al., 2008), a further seven main stations were sampled in

Table 1. Overview of the parameters determined in the present study. VLPs: virus-like particles; HB: heterotrophic bacteria; HNF: heterotrophic nanoflagellates; BP: bacterial heterotrophic production; BR: bacterial respiration; TNS: north–south transect; TEW: east–west transect; main stations: R-2, A3, E-4W, F-L, E-1, E-2, E-3, E-5, E-4E; ML: mixed layer; Z_{ML} : depth of the mixed layer.

Parameter	Station	Depth Layer
VLP, HB, HNF	TNS, TEW and main stations	Surface-bottom
BP	Main stations	Surface-bottom
BR	Main stations except E-2	ML
HNF grazing on bacteria	Main stations	ML, Z_{ML} –200 m, > 200 m
Virus induced bacterial mortality	Main stations	Surface-bottom

Table 2. Brief description of the main stations. The depth of the mixed layer (Z_{ML}) is based on a difference in sigma of 0.02 to the surface value. The mean Z_{ML} (\pm SD) of all CTD casts performed during the occupation of the stations is given. For chlorophyll *a* and major inorganic nutrients mean values \pm SD for the mixed layer are given. A3-1 and A3-2 refer to two consecutive visits of station A3. *: a single cast was performed at E-2

Station	Date (2011)	Longitude ($^{\circ}$ E)	Latitude ($^{\circ}$ S)	Depth (m)	Z_{ML} (m)	Chl <i>a</i> ($\mu\text{g L}^{-1}$) ^a	$\text{NO}_3^- + \text{NO}_2^-$ (μM) ^b	PO_4^{3-} (μM) ^b	Si(OH)_4 (μM) ^c
R-2	26/10	66.692743	50.38954	2450	105 \pm 15	0.25 \pm 0.08	26.0 \pm 0.2	1.83 \pm 0.03	12.3 \pm 0.3
A3-1	20/10	72.080388	50.62953	475	168 \pm 11	0.62 \pm 0.17	29.7 \pm 0.5	2.00 \pm 0.03	23.7 \pm 0.8
F-L	07/11	74.807422	48.61992	2690	38 \pm 7	4.00 \pm 1.58	20.5 \pm 1.9	1.06 \pm 0.21	7.7 \pm 0.8
E-4W	10/11	71.429833	48.76628	1398	61 \pm 11	2.38 \pm 0.31	25.4 \pm 1.0	1.79 \pm 0.10	18.5 \pm 1.2
A3-2	16/11	72.055283	50.62417	528	153 \pm 15	2.03 \pm 0.34	26.2 \pm 0.4	1.78 \pm 0.03	18.9 \pm 0.5
E-1	30/10	72.186833	48.45783	2050	72 \pm 38	0.94 \pm 0.08	25.7 \pm 0.5	1.75 \pm 0.05	15.1 \pm 0.4
E-2	01/11	72.077083	48.52343	2003	43*	0.78 \pm 0.50	27.3 \pm 0.6	1.84 \pm 0.13	15.9 \pm 2.0
E-3	04/11	71.966828	48.70215	1923	38 \pm 9	0.63 \pm 0.08	26.2 \pm 0.7	1.79 \pm 0.01	15.2 \pm 0.2
E-4E	13/11	72.566683	48.71500	2210	74 \pm 8	1.11 \pm 0.41	24.6 \pm 1.9	1.62 \pm 0.18	12.3 \pm 3.0
E-5	19/11	71.899917	48.4115	1920	46 \pm 13	1.15 \pm 0.07	25.4 \pm 0.2	1.74 \pm 0.04	11.7 \pm 0.2

Note: Z_{ML} at station E-2 is derived from a single CTD profile. ^a Lasbleiz et al. (2014). ^b Blain et al. (2014). ^c Closset et al. (2014)

the fertilized region (E-4W, F-L, E-1, E-2, E-3, E-4E, E-5, Fig. 1, Tables 1 and 2). Two of these main stations were chosen due to their high concentrations of surface chlorophyll, one located north (station F-L), and one located south of the southern branch of the polar front (E-4W). Based on the trajectories of two surface drifters, the other five main stations were sampled in a quasi-Lagrangian manner within a complex meander south of the polar front (E-1, E-2, E-3, E-4E, and E-5). Station A3 was visited twice (A3-1 and A3-2) during the onset of the bloom (KEOPS2), which complemented previous investigations following the decline of the phytoplankton bloom (KEOPS1). The reference site (station R-2) in high nutrient low chlorophyll waters (HNLC) was situated west of the plateau (Table 2). All water samples were collected with 12 L Niskin bottles mounted on a rosette equipped with a CTDO Seabird SBE911-plus. Sampling for microbial parameters presented here was performed at 11–14 depths at each station.

2.2 Abundances of microbial components and heterotrophic nanoflagellate grazing

The abundance of virus-like particles (VLPs) of heterotrophic bacteria [(HB) – sensu stricto heterotrophic Bacteria + Archaea] and HNF (heterotrophic nanoflagellates) were determined by flow cytometry. Subsamples (2.5 mL for VLP and HB, and 4.5 mL for HNF) were fixed with a transmission electron microscope (TEM) grade glutaraldehyde (final concentration 1 %) for VLP and HNF; and with formaldehyde (2 % final concentration) for HB. VLP and HB, and HNF were refrigerated for 10 to 20 min and 2 h, respectively, then frozen in liquid nitrogen and stored at -80°C until analysis. Counts of VLP and HB were made using a FACSCalibur flow cytometer (BD-Biosciences) equipped with an air-cooled laser, providing 15 mW at 488 nm with the standard filter set-up. VLP and HB were stained with SYBRGreen I, as described in detail in Marie et al. (1999) and Brussaard (2004). Populations of VLP and HB differing in fluorescence intensity were distinguished on plots of side scatter versus green fluorescence (530 nm wavelength, fluorescence channel 1 of the

instrument). Flow cytometry list modes were analysed using CellQuest Pro software (BD Biosciences, version 4.0). HNF were stained with SYBRGreen I and analysed according to Christaki et al. (2011) with a FACSCanto (BD-Biosciences).

To establish the size of cells of the three cytometric populations identified during this study (HNF1, HNF2, and HNF3, Fig. S1 in the Supplement), cells from each population were sorted with a FACS Aria cell sorter (BD-Biosciences). The sorted cells (1000–3000 cells per sample) were collected on Nuclepore filters (0.2 µm pore size, 25 mm diameter), and examined using a Zeiss AX10 microscope at 1000×. The mean biovolume of each cytometric population was calculated based on the linear dimensions of the cells, applying a prolate spheroid equation. Clearance rates (nl HNF⁻¹ h⁻¹) were estimated based on 10⁵ biovolume⁻¹ h⁻¹ (Fenchel, 1982; Christaki et al., 2001). The potential of HNF grazing accounting for the relative loss of bacterial heterotrophic production (BP) was then calculated as

$$\% \text{BP loss} = \text{bacterial cells "cleared"} \text{ L}^1 \text{ h}^{-1} \times 100 / \text{bacterial cells produced L}^{-1} \text{ h}^{-1}. \quad (1)$$

The number of bacterial cells produced was calculated from BP, as determined by leucine incorporation (see below), using a conversion factor of 12.4 fg C cell⁻¹ (Fukuda et al., 1998).

2.3 Phage-infected bacteria and burst size

For observations on the Transmission Electron Microscope (TEM), the 4.5 mL subsamples collected at different depths were pooled before ultracentrifugation for the following layers: the mixed layer (ML); from the bottom of the mixed layer to 200 m and the layer from below 200 m to the bottom. This operation resulted in one TEM observation per layer per station (Fig. S2 in the Supplement). Bacterial cells were harvested by ultracentrifugation onto 400 mesh copper electron microscope grids with carbon-coated Formvar film using a Beckman Coulter SW40Ti swing-out-rotor at 70 000× g for 20 min at 4 °C. Each grid was then stained for 30 s with uranyl acetate (2 % w/v) and examined in a JEOL 1200EX TEM operated at 80 kV at a magnification of 20 000–40 000×. For each grid, 600 bacterial cells per sample were examined to determine the frequency of visibly infected bacterial cells (FVICs), and the number of mature phages inside each host i.e. burst size (BS). A bacterium was considered infected if it contained five or more phages. Burst size in each of the 30 samples was defined as the average number of viral particles in all visibly infected cells. This is likely the minimum burst size, as more viral particles may accumulate within an infected cell before it lyses. To estimate viral induced bacterial mortality as a % of the bacterial production (VIBM), the frequency of infected cells (FIC with data given as percentages) was first calculated from the fre-

quency of FVIC according to Weinbauer et al. (2002):

$$\text{FIC} = 9.524 \times \text{FVIC} - 3.526. \quad (2)$$

The proportion of the total bacterial mortality that was due to virally induced lysis was calculated according to Binder (1999):

$$\text{VIBM} = (\text{FIC} + 0.6 \text{FIC}^2) / (1 - 1.2 \text{FIC}). \quad (3)$$

Viral production (VP) was estimated according to Weinbauer et al. (2003):

$$\text{VP} (\text{VLPL}^{-1} \text{ h}^{-1}) = \text{FIC}(\%) \times \text{BS} \times \text{BP} / 100. \quad (4)$$

The BS used in the equation was the BS of each of the 30 samples observed with the TEM.

2.4 Bacterial production, respiration, and bacterial growth efficiency

The incorporation of ³H leucine was used to estimate BP. Leucine concentrations and incubation times were tested on board and adjusted for different depths in order to obtain a sufficient radioactivity signal and to maintain linear uptake during the incubation. At each depth, 20 mL triplicate samples and a trichloroacetic acid (TCA)-killed control were incubated with a mixture of L-[4,5-³H] leucine (Perkin Elmer, 144 Ci mmol⁻¹) with nonradioactive leucine added at final concentrations of 7 and 13 nM for the upper 150 m, 13 and 7 nM for the 150–300 m depth layer, and 9 nM of ³H leucine alone below 300 m. Samples were incubated in the dark at in situ temperature. Incubation times were 3 h (0–150 m), 6 h (150–300), and 8 h (below 300 m). Incubations were terminated by adding ice-cold TCA (5 % final conc.), filtering through 0.22 mm Millipore cellulose acetate filters, and rinsing three times with 3–4 mL ice-cold 5 % TCA. After dissolving the filters in 1 mL ethyl acetate, 10 mL of Ultima Gold scintillation cocktail (Packard) were added and the samples were radioassayed. The incubation times (3–10 h) were tested to satisfy linear incorporation with time. Three stations inside and outside the bloom were checked at 5, 150, 500, and 750 m depths, by concentration kinetics (4, 6, 12, 20, and 40 nM), to ensure that there was no isotopic dilution. The theoretical conversion factor of 1.55 kg of C mol⁻¹ was used to convert leucine incorporation rates to prokaryotic carbon production (Kirchman, 1993).

Bacterial respiration was determined at all main stations, except R-2 and E-2, at one to three depths within the ML (Table 1) as described in Obernosterer et al. (2008). At station R-2, bacterial respiration rates were estimated from both dark community respiration (unfiltered seawater) and the fraction of dark community respiration accounted for by bacterial respiration as determined for the iron-fertilized sites (see Sect. 3.3). Briefly, rates of respiration were determined from dissolved oxygen consumption in 24 h dark incubations of 0.8 µm filtered samples. Dissolved oxygen was determined

by Winkler titration using a PC-based system with a photometric endpoint detector (Lefèvre et al., 2008). Bacterial growth efficiency was determined from in situ bacterial production and respiration rates as

$$\text{BGE} = (\text{BP} + \text{BR}) / \text{BR} \times 100. \quad (5)$$

We used a respiratory quotient of 1, as determined for the Kerguelen study region during KEOPS1 (Lefèvre et al., 2008) to convert oxygen units to carbon units.

3 Results

3.1 Study sites

The hydrographic conditions during KEOPS 2 are reported in detail in Park et al. (2014). The “historical” A3 station situated above the Kerguelen plateau (Blain et al., 2007, 2008) was characterized by a deep ML (Z_{ML} , depth of the mixed layer from 150 to 170 m) (Fig. 2). Here mean concentrations of Chl *a* in the ML increased from 0.6 to 2.0 $\mu\text{g L}^{-1}$ between the first and second visit three weeks later (Table 2). Stations F-L and E-4W revealed mean concentrations of 4.0 and 2.3 $\mu\text{g L}^{-1}$ Chl *a*, respectively, constrained in shallow ML (~ 40 m at station F-L and 60 m at station E-4W). The closely geographical positioned stations E-1, E-3, E-4E, and E-5 were visited between 30 October and 19 November in a quasi Lagrangian manner, and showed concentrations of Chl *a* between 0.6 and 1.2 $\mu\text{g L}^{-1}$ in the ML (Z_{ML} 40 m to 70 m) (Table 2). Temperature in the ML were highest at station F-L (4.2 °C, Fig. 2), indicating the influence of sub-Antarctic waters. Station E-4W was located at the shelf break in a region with very strong currents. E4-W received Fe-rich waters from the Kerguelen Island and Plateau (A3 station area) which mixed with polar front waters while travelling northeast (Fig. 1). The reference site (station R-2) in high nutrient low chlorophyll waters (HNLC) revealed comparatively low mean concentrations of Chl *a* (0.3 $\mu\text{g L}^{-1}$), and a surface temperature of 2.1 °C (Fig. 2) in the Z_{ML} (~ 100 m). The water mass below 200 m in the sampled area was characterized by winter waters from south of the polar front.

The macronutrient concentrations in the mixed layer were 20–30 μM for nitrate plus nitrite, 1–2 μM for phosphate, 8–24 μM for silicic acid (Table 2, Blain et al., 2014; Closset et al., 2014) and dissolved iron ranged between 0.08 and 0.35 nM (Quéroué et al., 2014). According to KEOPS2’s microscopical observations and pigment analysis, Bacillariophyceae dominated the phytoplankton community in the blooms (Sackett et al., 2014, Lasbleiz et al., 2014). In particular, *Fragilariopsis kerguelensis*, *Pseudonitzschia* spp., *Eucampia antarctica*, and *Chaetoceros* spp. were found to be the four dominant diatom taxa (Sackett et al., 2014).

3.2 Distribution of microbial community components

VLP, HB, and HNF abundances in the upper 200 m were of the order of 10^9 , 10^8 , and 10^5 particles cells L^{-1} , respectively (Fig. 3, Table 3). HB abundance, the %HNA (high nucleic acid) containing bacterial cells, and HNF abundances were significantly higher in the upper 200 m of the fertilized stations compared to the HNLC-site R-2 (Mann–Whitney, $p < 0.05$); this difference disappeared below 200 m (Table 3). The abundance of VLP was not significantly different between sites at any of the depth layers considered (Table 3). The variation between HNLC and Fe-fertilized sites was most pronounced for HNF (range ~ 200 – $900 \text{ S} \times 10^3 \text{ L}^{-1}$), if compared to VLP (range 1.4 – $1.7 \times 10^9 \text{ L}^{-1}$), and HB (range 2.7 – $4.7 \times 10^8 \text{ L}^{-1}$). HNF were distinguished in the three cytometrically identified subpopulations HNF1, HNF2, and HNF3 based on their cytometric signatures (Fig. 1). The individual cell biovolumes determined after cell sorting of each of these populations were $3.9 \pm 1.6 \mu\text{m}^3$ for HNF1, $36.9 \pm 9.3 \mu\text{m}^3$ for HNF2, and $62.2 \pm 41.1 \mu\text{m}^3$ for HNF3 μm^3 (all sites and depths pooled). The calculated clearance rates were 0.4 ± 0.2 , 3.7 ± 0.4 and $6.2 \pm 1.3 \text{ nl HNF}^{-1} \text{ h}^{-1}$ for HNF1, HNF2, and HNF3, respectively. Size-specific clearance rates differed only slightly among bloom sites, and no significant differences between the Fe-fertilized sites and HNLC waters were detected (Mann–Whitney tests, $p > 0.05$). The relative abundances of these cytometrically identified subpopulations were similar among sites and throughout the water column accounting for roughly 46 % for HNF1, 52 % for HNF2, and 2 % for HNF3 of the total HNF abundance (Table 3).

3.3 Bacterial production, growth rates, respiration, and growth efficiency

BP and bacterial growth rates were 5.2 $\text{nmol CL}^{-1} \text{ d}^{-1}$ and 0.018 d^{-1} in the ML at station R-2, and they were overall higher in the Fe-fertilized region. BP ranged between 9.9 and 133.8 $\text{nmol CL}^{-1} \text{ d}^{-1}$ and bacterial growth rates varied between 0.025 and 0.210 d^{-1} (mean ML at stations A3-1 and F, respectively) (Fig. 4 and Table 4). At station A3, BP and bacterial growth rates increased 4- to 5-fold between the first (9.9 $\text{nmol CL}^{-1} \text{ d}^{-1}$ and 0.025 d^{-1} , mean, ML) and the second visit (39.6 $\text{nmol CL}^{-1} \text{ d}^{-1}$ and 0.122 d^{-1}). Within the stationary meander, BP and bacterial growth rates increased over time, from 30 $\text{nmol CL}^{-1} \text{ d}^{-1}$ and 0.068 d^{-1} at station E-1, to 54.7 $\text{nmol CL}^{-1} \text{ d}^{-1}$ and 0.116 d^{-1} at station E-5. The bacterial production below 200 m varied from 0.07 to 2.18 $\text{nmol CL}^{-1} \text{ d}^{-1}$ for all stations (mean $0.55 \pm 0.40 \text{ nmol CL}^{-1} \text{ d}^{-1}$), which represented between 4.6 and 25 % (mean $16.5 \pm 7.0 \%$) in Fe-fertilized stations, and 43 % at station R-2 of the 0–1000 m integrated bacterial production (data not shown). Bacterial respiration rates varied by 8-fold among the Fe-fertilized sites, with lowest and highest rates at station

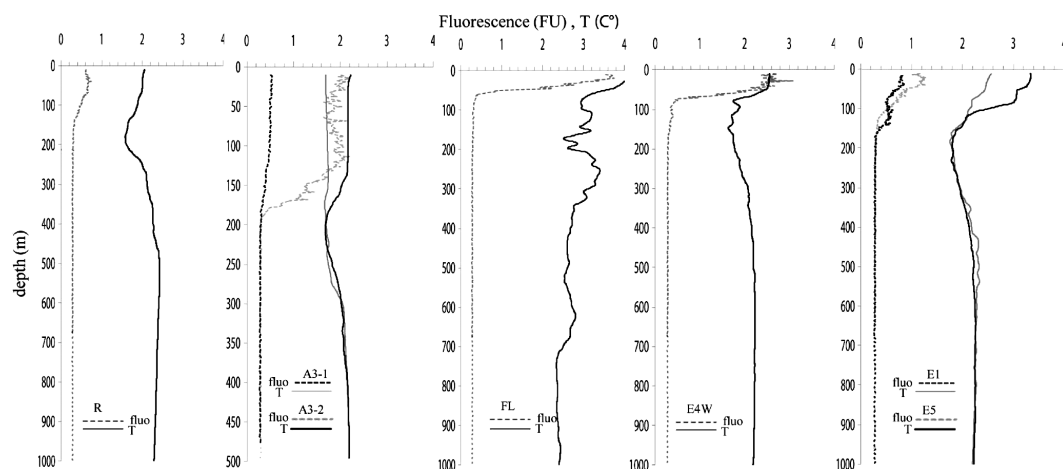


Figure 2. Profiles of temperature and Chl *a* as derived from in vivo fluorescence for the reference HNLC station (R-2); A3 (first and second visit), F-L, E-4W; and E stations first (E-1), and last visit (E-5).

Table 3. Bulk abundances and relative contributions of cytometrically identified subpopulations of virus-like particles (VLPs), heterotrophic bacteria (HB) and heterotrophic nanoflagellates (HNF). Mean values \pm SD of three depth layers for the iron-fertilized sites (Fe) and the HNLC site (R-2) are given. %GF: percent high and low green fluorescence VLPs; %HNA and %LNA: percent high and low DNA containing HB. HNF-1, -2, -3 populations were discriminated by flow cytometry and sized by epifluorescence microscopy after cell-sorting in each cytometric group (see also Fig. S1). ML: mixed layer; Z_{ML} : depth of the mixed layer; Z_{ML-200} : bottom of the mixed layer to 200 m; >200 m: below 200 m to the bottom of each station. Asterisks indicate significant differences between iron-fertilized sites and station R-2 (Mann-Whitney *: $p < 0.05$, **: $p < 0.001$ ***: $p < 0.0001$).

Layer	VLP ($\times 10^9 L^{-1}$)		% High GF		% Low GF				
	Fe	R-2	Fe	R-2	Fe	R-2			
ML	1.7 \pm 0.2	1.4 \pm 0.2	19.6 \pm 3.5	19.9 \pm 2.6	79.8 \pm 2.9	80.1 \pm 2.6			
Z_{ML-200} m	1.3 \pm 0.2	1.1 \pm 0.4	22.8 \pm 4.3	21.9 \pm 3.2	76.9 \pm 4.2	78.1 \pm 3.2			
>200 m	0.4 \pm 0.1	0.5 \pm 0.1	22.8 \pm 4.3	24.2 \pm 2.4	77.2 \pm 4.3	75.8 \pm 2.4			
		HB ($\times 10^8 L^{-1}$)		%HNA		%LNA			
		Fe	R-2	Fe	R-2	Fe	R-2		
ML		4.7 \pm 0.9	2.7 \pm 0.3	59.0 \pm 5.3	47.3 \pm 2.5***	41.5 \pm 4.7	52.7 \pm 2.5***		
Z_{ML-200} m		3.1 \pm 0.4	2.1 \pm 0.8 *	55.4 \pm 1.1	53.5 \pm 4.6	44.6 \pm 1.1	46.5 \pm 4.6		
>200 m		1.2 \pm 0.3	1.1 \pm 0.2	56.7 \pm 2.2	55.7 \pm 2.1	43.3 \pm 2.2	44.3 \pm 3.7		
		HNF ($10^3 L^{-1}$)		%HNF-1		%HNF-2		%HNF-3	
		Fe	R-2	Fe	R-2	Fe	R-2	Fe	R-2
ML		903 \pm 549	227 \pm 71 *	49 \pm 6	47 \pm 5	48 \pm 6	51 \pm 4	3 \pm 3	2 \pm 4
Z_{ML-200} m		367 \pm 122	170 \pm 82 *	44 \pm 10	45 \pm 5	54 \pm 10	52 \pm 7	2 \pm 2	3 \pm 3
>200 m		84 \pm 37	60 \pm 26	46 \pm 11	48 \pm 6	53 \pm 11	51 \pm 2	1 \pm 2	1 \pm 1

E-1 (mean ML $0.23 \pm 0.06 \mu\text{mol C L}^{-1} \text{d}^{-1}$, $n = 3$) and E-5 ($1.73 \mu\text{mol C L}^{-1} \text{d}^{-1}$, $n = 1$), respectively (Table 5). At the Fe-fertilized sites bacterial respiration accounted on average for $59 \pm 20\%$ of dark community respiration. We estimated bacterial respiration at station R-2 to vary between $0.25 \pm 12 \mu\text{mol C L}^{-1} \text{d}^{-1}$ ($n = 4$) (dark community respiration in unfiltered seawater) and $0.14 \pm 0.07 \mu\text{mol C L}^{-1} \text{d}^{-1}$ (i.e. the 59% of dark community respiration) (Table 5). Due

to the lower contribution of phytoplankton and HNF to overall microplankton biomass in HNLC compared to Fe-fertilized waters, the contribution of bacterial to dark community respiration is likely to be higher at station R-2 than the mean value determined for the Fe-fertilized sites. We therefore use the respiration rate in unfiltered seawater as an upper estimate for bacterial respiration throughout the manuscript (Table 5). The cell-specific respiration revealed

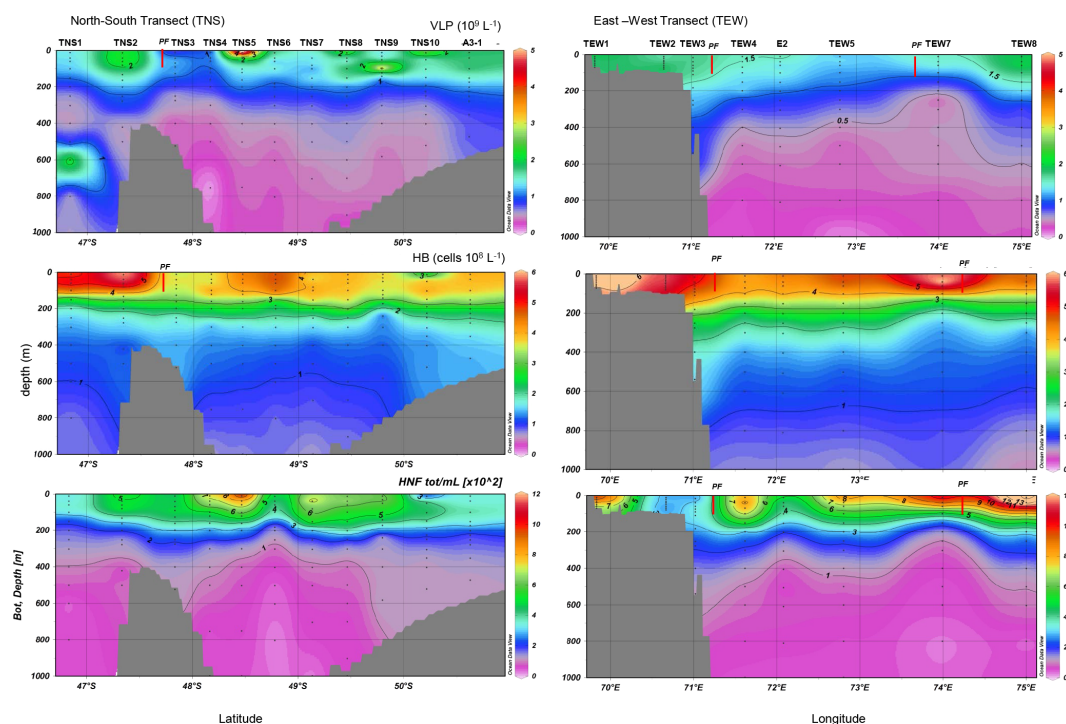


Figure 3. Distribution of the abundance of virus (VLPs – virus-like particles), heterotrophic bacteria (HB), and heterotrophic nanoflagellates (HNF) along the north–south (TNS) and east–west transects (TEW; cf. Fig. 1) of KEOPS2. The red vertical line “PF” denotes the position of the polar front.

Table 4. Bacterial production, cell-specific bacterial production, and bacterial growth rates in surface waters. Mean values \pm SD for the mixed layer are given.

Station	Number of samples (n)	BP (nmol C L ⁻¹ d ⁻¹)	Cell-specific BP (fmol C cell ⁻¹ d ⁻¹)	μ (d ⁻¹)
R-2	4	5.2 \pm 0.2	0.019 \pm 0.003	0.018 \pm 0.003
A3-1	5	9.9 \pm 0.3	0.025 \pm 0.007	0.025 \pm 0.007
A3-2	5	39.6 \pm 1.3	0.126 \pm 0.015	0.122 \pm 0.015
F-L	2	133.8 \pm 3.3	0.217 \pm 0.005	0.210 \pm 0.005
E-4W	3	58.3 \pm 3.4	0.097 \pm 0.012	0.094 \pm 0.012
E-1	3	30.3 \pm 1.65	0.071 \pm 0.002	0.068 \pm 0.003
E-3	2	52.3 \pm 3.4	0.098 \pm 0.007	0.095 \pm 0.006
E-4E	2	83.0 \pm 2.6	0.147 \pm 0.009	0.143 \pm 0.009
E-5	2	54.7 \pm 4.0	0.119 \pm 0.005	0.116 \pm 0.005

a similar pattern with lowest rates at stations E - 1 (mean ML 0.54 ± 0.13 fmol O₂ cell⁻¹ d⁻¹, $n = 3$), and highest rates at station E-5 (3.76 fmol O₂ cell⁻¹ d⁻¹, $n = 1$) (Table 5). BGE ranged between 3 and 18% in the ML of the fertilized sites (mean 10 ± 5 %, $n = 14$). One exceptionally high value of 28% at the base of the ML at station F-L was registered. At station R-2, BGE was 3 ± 1 % ($n = 4$) based on respiration rates in unfiltered seawater. Assuming BR accounts for 59% of dark community respiration, as described above, would increase the BGE at stations R-2 to 4 ± 2 %.

3.4 Bacterial losses and viral production

The potential grazing capacity estimated by the clearance rates of HNF revealed losses of BP of 50, 70, and 85% for the ML, the Z_{ML}–200 m, and the > 200 m layers, respectively (Fig. 5a). In surface waters (ML) the loss of BP due to grazing varied between ~ 30 and ~ 60 % at the fertilized stations, with the exception of station A3-1 where this value was 80%. The BP loss due to grazing at station R-2 accounted for about 70% of the bacterial production (Fig. 5a). BP loss due to viral lysis was comparatively low, and varied from

Table 5. Bacterial respiration, cell-specific bacterial respiration, and growth efficiency (BGE) in surface waters. Mean values \pm SE are given. At station R-2, respiration rates of unfiltered samples were used. (*) Bacterial respiration rates have been converted from O_2 into C units using a RQ value of 1.

Station	Depth (m)	Bacterial respiration ($\mu\text{mol C L}^{-1} \text{d}^{-1}$)*	Cell-specific bacterial respiration ($\text{fmol C cell}^{-1} \text{d}^{-1}$)	BGE (%)
R-2	20	0.19 ± 0.09	0.77	2 ± 1
	40	0.40 ± 0.06	1.41	2 ± 0
	80	0.27 ± 0.08	0.90	2 ± 1
	100	0.12 ± 0.05	0.43	4 ± 1
A3-2	20	0.61 ± 0.23	2.26	6 ± 2
	50	1.08 ± 0.51	4.20	3 ± 1
	70	0.19 ± 0.07	0.54	18 ± 6
F-L	10	0.91 ± 0.50	1.50	13 ± 6
	20	1.82 ± 0.28	3.00	7 ± 1
	50	0.25 ± 0.23	0.39	28 ± 19
E-4W	30	0.95 ± 0.51	1.57	6 ± 3
	50	0.84 ± 0.22	1.39	7 ± 2
E-1	15	0.27 ± 0.17	0.62	11 ± 6
	20	0.26 ± 0.21	0.60	11 ± 8
	65	0.17 ± 0.30	0.40	15 ± 9
E-3	10	0.25 ± 0.34	0.49	17 ± 19
	20	0.96 ± 0.61	1.89	5 ± 3
	46	1.43 ± 0.32	2.83	3 ± 1
E-4E	30	0.72 ± 0.13	1.28	11 ± 2
	50	0.65 ± 0.44	1.16	11 ± 7
E-5	20	1.73 ± 0.46	3.76	3 ± 1

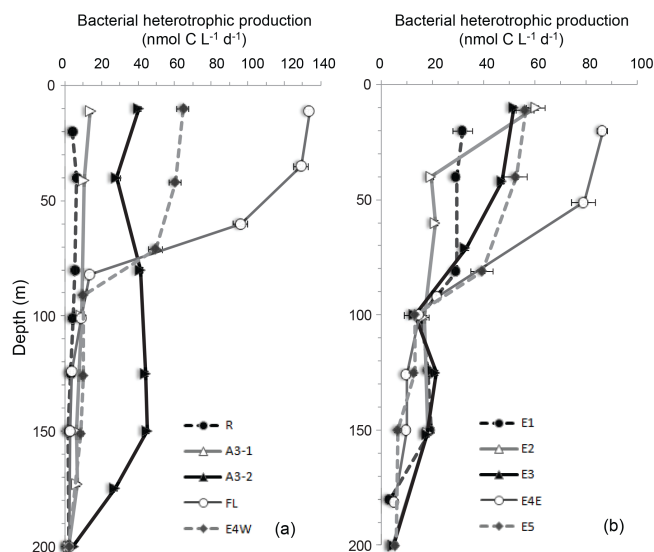


Figure 4. Bacterial production ($\text{nmol C L}^{-1} \text{d}^{-1}$) at the main stations in the 0–200 m layer. Station R; A3-1, and 2; F-L, and E-4W (a); and “E” stations (b).

undetectable to 24 % of BP in the ML (Fig. 5b). The viral induced loss of BP were 11, 2, and 2 % for the ML, $Z_{\text{ML}}-200$ m, and >200 m layers, respectively (Fig. 5b). A higher percent of viral mortality was encountered in the ML at stations E-4E, E-2, E-1, and R-2. Overall there was no difference in the percent loss of BP induced by grazing and viral lysis between the Fe-fertilized sites and the HNLC site R-2 (Fig. 5a, b). The sum of bacterial mortality due to HNF grazing and viral lysis varied from 47 % at station E-1, to 100 % at stations E-3 and E-5. Together, grazing and viral lysis accounted for an average of 83 % bacterial mortality at all stations (Fig. 5a, b). The empirically estimated burst size of bacteria (BS) was 22 ± 15 virus cell^{-1} (mean \pm SD, $n = 30$), and varied from 6 (A2, in the ML) to 88 (F-L, in the $Z_{\text{ML}}-200$ m) without any specific pattern related to Fe-fertilization or depth. Viral production (VP) calculated based on the BS and BP (Eq. 4) was 2 ± 1 and $0.34 \pm 0.07 \times 10^8$ viruses $\text{L}^{-1} \text{d}^{-1}$ in the ML of the fertilized stations, and at station R-2, respectively.

Combining these results revealed several significant relations (Table 6). In the ML, HB and BP were significantly related to Chl *a*. BP and the abundance of HNF were significantly related in the three layers considered, with BP

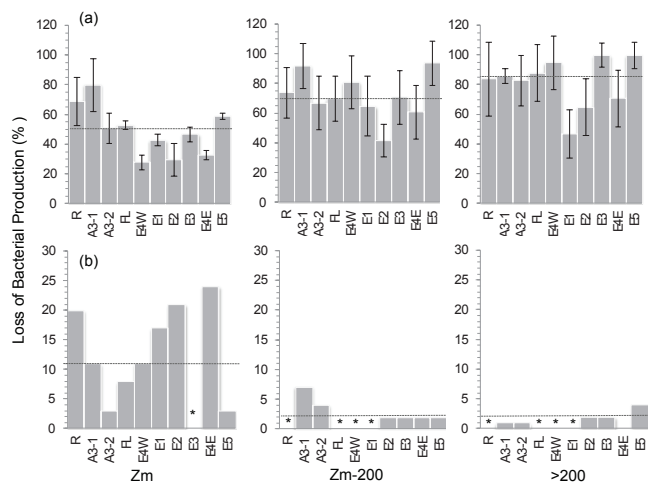


Figure 5. Heterotrophic nanoflagellate (a) and virus (b) mediated mortality of bacteria, as a percentage of bacterial production measured in the mixed layer (ML), from the bottom of the mixed layer to 200 m ($Z_{ML-200m}$), and between 200 and 1000 m (>200 m). The vertical line denotes the median value for each layer. Error bars in (a) indicate the standard deviation of the mean values of each layer. No error bars in (b) since the values are issue of TEM observations of the pooled samples in each layer (see also methods). *: undetectable, the % FVIC was < 0.4 resulting in negative values of %FIC (Eq. 1).

explaining up to 79 % of the HNF variability ($r^2 = 0.79$). While BP and VLP – except in the $Z_{ML-200m}$ layer – showed little relation, BP and VP showed highly significant relations in the first 200 m, and a weaker, but still significant relation in the deep layer. It is worth noting that the relations with VP were insignificant when an overall mean BS of 22 virus cell⁻¹ was applied, but became highly significant when the specific BS for each sample was applied. VP and HB abundance were significantly related in the upper 200 m, however the determination coefficient was relatively low ($r^2 = 0.5$). The relation between BP and BR was significant with BP explaining ~ 50 % of the BR variance ($r^2 = 0.51$). The relation between VP and BR was weaker, but still significant ($r^2 = 0.32$). Notably, there was no detectable relation between VP and VLP at any depth. The stocks of HB were significantly related to the stocks of VLP and HNF at all layers, with strongest relations between HB and HNF below the ML (Table 6).

4 Discussion

4.1 Variability of the response, and comparison to other fertilization studies in the Southern Ocean

To date, six artificial mesoscale, and two detailed natural iron fertilization studies have been conducted in the Southern Ocean and microbial food web dynamics have been con-

Table 6. Determination coefficients (r^2) of log-log linear regressions of bacterial heterotrophic production (BP), viral production (VP), virus-like particles (VLPs), heterotrophic bacterial abundance (HB), heterotrophic nanoflagellates abundances (HNF) and chlorophyll *a* (Chl *a*). ML: mixed layer, *: $p < 0.05$, **: $p < 0.001$ ***: $p < 0.0001$, ns: not significant.

Layer	Parameter 1	Parameter 2	<i>n</i>	r^2
ML	BP	VP	27	0.86***
$Z_{ML-200m}$			41	0.66***
200–1000 m			24	0.25 *
ML	BP	BR	20	0.51 **
ML	BP	HB	27	0.51***
$Z_{ML-200m}$			61	0.83***
200–1000 m			51	0.43***
ML	BP	VLP	27	0.01 ns
$Z_{ML-200m}$			61	0.17**
200–1000 m			51	0.012 ns
ML	BP	HNF	27	0.68***
$Z_{ML-200m}$			61	0.79***
200–1000 m			51	0.20*
ML	BP	Chl <i>a</i>	27	0.57***
ML	VP	HB	27	0.50***
$Z_{ML-200m}$			41	0.47***
200–1000 m			24	0.01 ns
ML	VP	VLP	27	0.001 ns
$Z_{ML-200m}$			41	0.02 ns
200–1000 m			24	0.006 ns
ML	VP	BR	20	0.32*
ML	HB	VLP	27	0.29*
$Z_{ML-200m}$			61	0.27***
200–1000 m			51	0.11*
ML	HB	HNF	27	0.20*
$Z_{ML-200m}$			61	0.84***
200–1000 m			51	0.46***
ML	HB	Chl <i>a</i>	27	0.40*

sidered in part in these studies (Table 7). The most commonly determined microbial parameters, such as HB abundances and BP, are generally enhanced upon iron fertilization. However, remarkable variability in the extent of the response exists among different regions of the Southern Ocean (Table 7). During KEOPS2, a patchwork of blooms induced by large-scale natural iron fertilization above the plateau and in the oceanic region off Kerguelen Island was investigated (Blain et al., 2014). The results obtained from KEOPS2 add to previous studies by providing an extensive description of microbial food web dynamics during the onset of spring phytoplankton blooms under varying hydrographic and biogeochemical conditions.

Table 7. Comparison of bacterial parameters following natural and artificial iron-fertilization. Ranges of bacterial abundance and production are given for the wind mixed layer of the respective study. All rates of bacterial production are derived from [^3H]leucine-incorporation, except for Hall and Safi (2001) who applied [^3H]thymidine-incorporation. Outside – values reported for HNLC waters; inside – values reported for iron-fertilized patches. –: not done, ^a BP values given in this paper as pM leu h^{-1} were converted into $\mu\text{g C L}^{-1} \text{d}^{-1}$ according to Kirchman (1993), ^b not given in the paper, ^c calculated from cell production based on a $23.3 \text{ fg cell}^{-1}$ used by the authors, ^d max. value in the patch, mean value outside the patch extrapolated from Fig. 10, Landry et al. (2000).

		Temp. (°C)	Time elapsed since start of bloom (d)	Bacterial abundance ($\times 10^8 \text{ cells L}^{-1}$)		HNF abundance ($\times 10^6 \text{ cells L}^{-1}$)		Bacterial production ($\mu\text{g C L}^{-1} \text{d}^{-1}$)		Viral abundance ($\times 10^9 \text{ cells L}^{-1}$)		Reference
				outside	inside	outside	inside	inside	inside	outside	inside	
Natural Southern Ocean	KEOPS2 (50° S, 72° E)	2	≈ 28	2.7	4.7	0.2	0.9	0.1	0.2–1.6	1.4	1.7	This study
	KEOPS1 (50° S, 72° E)	3–4	≈ 60	2	6	0.8	1.1	0.5	1.1–3	3.9	12.5	Christaki et al. (2008); Bonilla-Findji et al. (2008)
	CROZEX (45° S, 50° E)	6	≈ 60	5	9	–	–	0.42	3.6	–	–	Zubkov et al. (2007)
Artificial Southern Ocean	EisenEx (48° S, 21° E)	3–4	22	3–4	2–6	–	–	0.2–0.5	0.2–1	2.1	4.3	Arrieta et al. (2004); Weinbauer et al. (2009)
	SOIREE (61° S, 140° E)	2	13	2–4	2–4	0.4	1.1	<0.5	0.4–1.1	–	–	Hall and Safi (2001)
	SOFEX-North (56° S, 172° W)	7	40	4–5	7–10	–	–	0.2	0.4	–	–	Oliver et al. (2004)
	SOFEX-South (66° S, 171° W)	–1	22	3	3–5	–	–	0.2	0.2–0.3	–	–	Oliver et al. (2004)
	SAGE (46.5° S, 172.5° W)	11.5	15	0.9–3.2	1.2–1.7	–	–	0.9	0.4–1.4	–	–	Kuparinen et al. (2011)
	LOHAFEX (47° S, 16° W)	7	18	1.1–1.4	1.1–1.4	–	–	1	1–2 ^a	–	–	Thiele et al. (2012)
Subarctic Pacific	SEEDS I (49° N, 165° E)	6–9	13	5	8	2.1	5.8	–	–	–	–	Suzuki et al. (2005); Saito et al. (2005)
	SEEDS II (48° N, 166° E)	8–12	12	4–7	5–10	–	–	b	2–7 ^c	23–45	22–64	Higgins et al. (2009)
Equatorial Pacific	IronEx II (5° S 105° W)	25	14	7	11	~2	3.9 ^d	2–3	2–8	–	–	Landry et al. (2000); Cochlan (2001)

Natural iron fertilization of the Southern Ocean induced rapid responses of members and fluxes of the microbial food web. The intensity of the response and its variability among sites were most pronounced in the 200 m surface layer, and this likely reflects the hydrographic and biogeochemical characteristics at the given sites. The depth of the mixed layer (Z_{ML}) varied considerably, extending down to 170 m above the plateau at station A3, and to 40 m north of the polar front at station F-L. To note that although surface Chl *a* concentration, which is a proxy for phytoplankton biomass, varied by a factor of 6 in the ML among sites, the integrated Chl *a* at A3-2 and F-L stations were similar (Lasbleiz et al., 2014). Noticeable differences in surface temperature, a key factor that drives microbial metabolism, were also detected in the ML (1.6–4.2 °C). Our results showed the BP response to be strong and varied, accounting for increases of mean values in the ML of up to 13-fold between bloom stations (e.g. A3-1 compared to F-L, Table 4), and up to 26-fold between bloom stations and HNLC station R-2 (e.g. F-L compared to R-2, Table 4). This enhancement in BP, together with that observed during CROZEX (9-fold; Zubkov et al., 2007) and KEOPS1 (6-fold; Christaki et al., 2008), was much higher than the enhancement reported from artificial fertilization experiments (roughly 2-fold; Table 7).

During KEOPS2, BR was up to 8-fold enhanced by iron fertilization and varied by a factor of 8 among sites among bloom stations. During the late bloom phase, above the

plateau, bacterial respiration was about 3-fold higher in the bloom than in HNLC waters (Obernosterer et al., 2008), but to the best of our knowledge, no BR rates are available for comparison from other fertilization studies in the Southern Ocean.

In contrast to bacterial metabolism, the abundance of HB increased overall to a lesser extent (roughly 2-fold), and thus similarly to those reported previously in natural and artificial iron fertilization studies (Table 7). Besides bulk abundance, the %HNA bacteria were also significantly higher in the ML of fertilized stations (59 %) relative to the HNLC reference site R-2 (47 %; Table 3). Oliver et al. (2004) reported a relatively minor increase of the fraction of HNA cells (up to 45 %) within the iron-fertilized patch during SOFeX. By contrast, the % HNA bacteria accounted for up to 80 % of total bacterial abundance at the late stage of the Kerguelen bloom at station A3 (Obernosterer et al., 2008). The smaller increase in HB abundance indicates efficient top-down control of these members of the microbial food web that are markedly stimulated by natural iron fertilization.

Although HNF showed significantly higher abundances in the 200 m surface layer in the fertilized stations compared to station R-2, the HNF size distribution, resulting from sorted cell observations, was remarkably stable, with the two smaller sized populations contributing > 45 % each, at all stations and depths (Table 3). The few existing studies reporting on viral abundance during iron fertilization experiments

(Higgins et al., 2009; Weinbauer et al., 2009) have pointed to a higher viral stock in fertilized stations. However, this was not the case in the present study, where the total number of VLPs was similar in fertilized sites and HNLC waters. On average $\sim 80\%$ of the VLPs during KEOPS2 showed low green fluorescence, and were most probably bacteriophages. The high green fluorescence VLPs represented $\sim 20\%$ of total VLPs in the study area, and according to Brussaard et al. (2008); this group may represent larger algal viruses.

Although viral stocks did not differ across stations in this study, VP was about 6 times higher in the ML of the fertilized stations compared to HNLC waters. This result is in line with observations from the late bloom phase (KEOPS1, Malits et al., 2014) and artificial iron fertilization experiments (Higgins et al., 2009; Weinbauer et al., 2009). Contrary to previous observations (Weinbauer et al., 2009), the trend of higher bacterial and viral production in iron-fertilized waters was not accompanied by a higher virus induced loss of BP. Lysogeny was not considered in the present study. The proportion of the lysogenized bacterial population can vary extensively from undetectable to 100% (Weinbauer and Suttle, 1996; Williamson et al., 2002). Across a system study, lysogeny was highest in deep sea waters (Weinbauer et al., 2003) where the contact rate between infective phages and hosts is too low to sustain the lytic lifestyle (Paul et al., 2002). The only study on lysogeny during artificial Fe-fertilization experiments in the Southern Ocean did not observe differences inside and outside the patch (Weinbauer et al., 2009). During KEOPS1 the fraction of lysogenic cells was 8 and 6% of the total bacterial cells infected by viruses in the bloom, and in HNLC waters, respectively (Malits et al., 2014). Based on these observations (Weinbauer, 2009; Malits et al., 2014), and this paper's independently obtained results (such as the low viral abundance and the low frequency of infected cells) supports the idea that the loss of BP due to lysis was low at the onset of the phytoplankton bloom. Notably, viruses and HNF revealed opposite vertical trends, with HNF grazing increasing and viral lysis decreasing with depth.

4.2 Microbial food web dynamics in response to iron fertilization

The extent of change of the microbial parameters considered in the present study at different fertilized sites across variable hydrographic and biological regimes appears to have been induced primarily by changes in heterotrophic bacterial activity. Based on the microbial parameters obtained during this study, the following scenario is proposed: typical Southern Ocean characteristics are low phytoplankton biomass and low rates of BP, as was the case at stations R-2 and A3-1. Fe-fertilization stimulates BP; either directly or indirectly, through phytoplankton derived DOM. For the KEOPS2 study region, bottle incubation experiments revealed that both single additions of Fe and C, in the form of glucose, stimulated bacterial heterotrophic production and

growth, suggesting co-limitation by these elements (Obernosterer et al., 2014).

The strong correlation between BP, VP, and HNF abundance, and the correlation between VP and HB, suggests that the enhanced BP drives HNF and viral activity. However, the overall loss of BP due to grazing and lysis decreases upon Fe fertilization (to about 50% of BP), indicating that bacterial growth rates are higher than loss rates, allowing the build-up of bacterial biomass. A notable aspect is that HNF responded more rapidly, and grazing became the dominant top-down factor at the early bloom stage. This pattern contrasts to observations from the late bloom stage, where viral-induced lysis accounted for most of the bacterial loss (Malits et al., 2014). This scenario agrees with observations of rapid growth of HNF at low temperatures (Duarte et al., 2005), and the “kill the winner” model (Thingstad, 2000) that predicts viruses to rise following blooms of their specific bacterial hosts. Indeed, the composition of the bacterial community changed markedly during the bloom above the plateau (West et al., 2008; Obernosterer et al., 2011). Taken together, these results suggest that the mechanism of natural fertilization through the continuous supply of iron strongly affects bacterial heterotrophic metabolism, and as a consequence HNF grazing, with important consequences for the cycling of carbon through the microbial food web.

4.3 Implications of microbial food web dynamics for carbon cycling in the bloom above the Kerguelen plateau

The investigation of the spring phytoplankton bloom located in the southeastern part above the Kerguelen plateau during two distinct phases has provided, for the first time, insight into seasonal dynamics of the microbial food web functioning in the context of natural Fe fertilization of the Southern Ocean. Combining our results with rates of primary production, mesozooplankton activity, and export determined during the KEOPS project allowed us to propose a budget for the flow of carbon through microbial and higher trophic levels in the southeastern bloom (A3 station, Fig. 6). The main purpose of the carbon budget presented here (Fig. 6) is to place the microbial loop in the context of the food web and to compare the fluxes to the potential export and/or accumulation of phytoplankton biomass during the early and late phases of the bloom.

Gross community production (GCP) integrated over the ML was about 3.6-fold higher during the early ($344 \text{ mmol C m}^{-2} \text{ d}^{-1}$) than the late bloom phase ($95 \text{ mmol C m}^{-2} \text{ d}^{-1}$, Fig. 6), however, the fraction of GCP that passed through heterotrophic bacteria, that is the bacterial carbon demand (BCD), increased from 21% at the onset of the bloom to 44% at the late bloom stage (BCD absolute values, 72 and $42 \text{ mmol C m}^{-2} \text{ d}^{-1}$, Fig. 6). BR revealed also marked seasonal differences (Fig. 6). The higher rates of respiration at the onset of the bloom

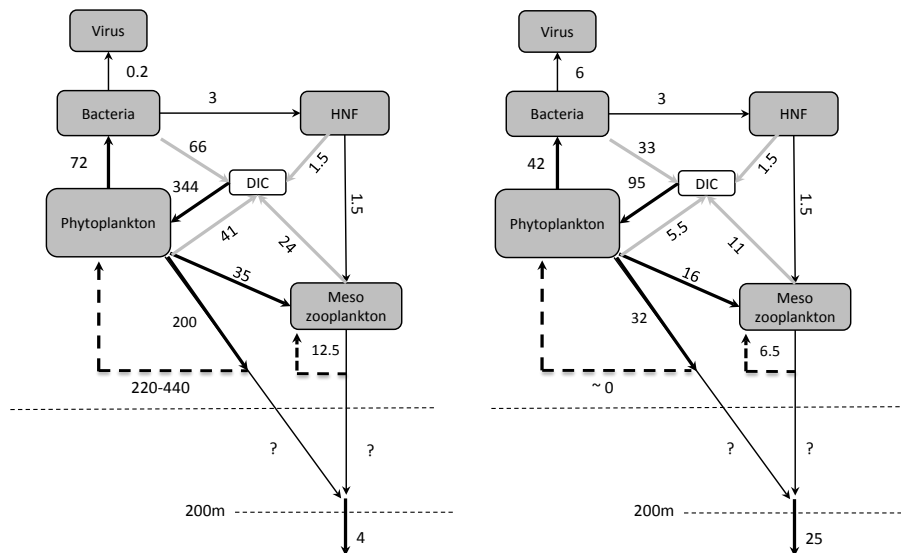


Figure 6. Schematic presentation of the carbon flow through the microbial and higher trophic level food webs during the early (KEOPS2) and late stage of the bloom (KEOPS1) at the southeast bloom above the Kerguelen plateau (station A3). KEOPS1 values are the mean obtained from four consecutive visits at station A3, except for dark community respiration for which only the first three visits were considered. Black arrows indicate the transfer of carbon between microbial and/or higher trophic level food web components, and export. Dashed black arrows going back to “phytoplankton” and “mesozooplankton” indicate possible accumulation of their biomass per day once the respiration and export were subtracted (for details cf. Sect. 4.3). Arrows in grey indicate rates of respiration. Numbers in italics are based on the following estimations: (i) the transfer from HNF to mesozooplankton is calculated from bacterial grazing assuming a 50 % transfer efficiency. (ii) the mesozooplankton ingestion and respiration rates were calculated from the respective rates during the late bloom phase and extrapolated according to the depth of the mixed layer (Z_{ML}). All fluxes are integrated over the ML and units are $\text{mmol C m}^{-2} \text{d}^{-1}$. Late bloom data from Carlotti et al. (2008), Christaki et al. (2008), Lefèvre et al. (2008), Obernosterer et al. (2008), Savoye et al. (2008) and Malits et al. (2014). Early bloom data from Planchon et al. (2014) and Carlotti et al. (2014) for GCP the O_2 -measured fluxes have been converted into C-fluxes using $PQ = 1.8$ KEOPS2 unpublished data. DIC: dissolved inorganic carbon, Z_{ML} : depth of the mixed layer.

($66 \text{ mmol C m}^{-2} \text{d}^{-1}$, Fig. 6) as compared to the late bloom stage ($33 \text{ mmol C m}^{-2} \text{d}^{-1}$, Fig. 6) result mainly from the deeper Z_{ML} (153 m vs. 67 m), because on a volumetric basis, rates were in a similar range during both bloom phases (cf. Obernosterer et al., 2008).

The ratios between the BCD and GCP were in the same range at the different phytoplankton bloom sites (BCD : GCP ~ 0.1 – 0.5), and overall lower than those observed during the late bloom phase (BCD : GCP ~ 0.3 – 0.7) (Table 1 in the Supplement). These latter estimates are similar to those reported for spring phytoplankton blooms in the polar frontal zone, the marginal ice zone, and the southern Antarctic Circumpolar Current (0.3 to 0.6; Lochte et al., 1997). To estimate the fraction of primary production that is processed by heterotrophic bacteria most studies report on the ratio BP : PP. For KEOPS, this ratio was on average 0.04 ± 0.02 for the early phytoplankton blooms compared to an average of 0.10 ± 0.03 for the late bloom phase at station A3 (Table 1 in the Supplement). At the HNLC site R-2 the BP : PP ratio was 0.045. Except for the late bloom stage at A3, the ratios reported here are in the same range as those reported for the Ross Sea (0.04; Ducklow, 2000) and the Arctic Ocean, when pri-

mary production was greater than $10 \text{ mmol C m}^{-2} \text{d}^{-1}$ (0.06, Kirchman et al., 2009).

Even though the absolute values of BP transferred to HNF and to mesozooplankton were the same during the early and the late bloom (3 and $1.5 \text{ mmol C m}^{-2} \text{d}^{-1}$, respectively, Fig. 6) the grazing induced consumption of BP was more important during the early than the late phase of the bloom (50 and 36 %, respectively). Interestingly, viral activity showed the opposite pattern, since viral induced mortality changed significantly over time, becoming the dominant top-down control of BP during the late bloom phase (Fig. 6, Malits et al., 2014). This suggests that a larger proportion of BP was channelled to HNF and to higher trophic levels during the early bloom, while most of BP returns to DOM through the viral shunt during the late bloom (Wilhelm and Suttle, 1999). These contrasting scenarios have important implications for organic matter and nutrient cycling.

During the early stage of the bloom, the different loss terms of GCP were bacterial: (i) ($66 \text{ mmol C m}^{-2} \text{d}^{-1}$), other microplankton ($41 + 1.5 \text{ mmol C m}^{-2} \text{d}^{-1}$) and mesozooplankton ($24 \text{ mmol C m}^{-2} \text{d}^{-1}$) respiration, (ii) $0.2 \text{ mmol C m}^{-2} \text{d}^{-1}$ to dissolved pool through bacteria lysis and, (iii) while 11 (i.e. ingestion – respiration of

microzooplankton) and $1.5 \text{ mmol C m}^{-2} \text{ d}^{-1}$, respectively, were transferred directly or through the microbial food web to mesozooplankton for biomass accumulation or export through faecal pellet production. Subtracting the sum of the above terms from GCP yields $200 \text{ mmol C m}^{-2}$ that remained available in the water column for phytoplankton biomass accumulation and/or export (Fig. 6). Comparison to the POC flux ($4 \text{ mmol C m}^{-2} \text{ d}^{-1}$ at 200 m, Planchon et al., 2014), indicates that a minor fraction of this GCP was exported during the early stage of the bloom. Assuming the accumulation of the POC stock ($2190 \text{ mmol C m}^{-2}$, Lasbleiz et al., 2014) over 5–10 days provides an estimate for phytoplankton biomass accumulation of $220\text{--}440 \text{ mmol C m}^{-2} \text{ d}^{-1}$ (Fig. 6).

During the late stage of the bloom, organic carbon lost through bacterial, micro- and mesozooplankton respiration amounted to $51 \text{ mmol C m}^{-2} \text{ d}^{-1}$, while 5 and $1.5 \text{ mmol C m}^{-2} \text{ d}^{-1}$, respectively, were transferred directly or indirectly to mesozooplankton. Thus, 32 mmol C m^{-2} of GCP were not processed in the water column (Fig. 6). This value is similar to the POC flux of $25 \text{ mmol C m}^{-2} \text{ d}^{-1}$ at 200 m (Savoie et al., 2008), and thus our carbon budget indicates a negligible potential for phytoplankton biomass accumulation during the late bloom phase (Fig. 6).

Primary production was measured at one time point during the onset of the bloom at A3 station, and the error associated to the integrated flux is 10%. Daily fluxes are more likely variable due to the influence of environmental factors, such as light intensity, which could in part account for the observed discrepancy.

Concluding, the concurrent investigation of several parameters has provided insight into two key roles of heterotrophic bacteria, and the microbial food web functioning, at the onset and late phase of the spring phytoplankton bloom induced by natural iron fertilization in the Southern Ocean. First, during the early bloom phase the tight link between HNF and heterotrophic bacteria resulted in a more efficient transfer of the BP towards the higher trophic levels. Even though the amount of bacterial carbon transferred did not represent a major flux, it accounted for a larger percentage of BP. Second, the bacterial response was dominated by an increase in respiration that was not accompanied by a respective enhancement of BP, pointing to an inefficient utilization of DOM at the onset of the bloom. Due to the low BGE, and the deep mixed layer above the plateau, heterotrophic bacteria represent a major source of CO_2 at the early bloom phase. This picture differs from that obtained during the late bloom phase, where BGE was higher, but the viral shunt prevented an efficient transfer of the BP to higher trophic levels. This study highlights the variability of the bacterial contribution to organic matter remineralization and export according to the trophic dynamics at different seasons, underlying varying mechanisms channelling organic matter. To improve our understanding of microbial dynamics, further investigation into time series over the season will be necessary in order to

determine the timing and factors that induce the changes in channelling organic matter in the upper ocean.

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