Changes in latitude and dominant diazotrophic community alter N₂ fixation

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ABSTRACT: The contribution of planktonic diazotrophs to the overall N budget is a key unknown in the eastern Indian Ocean. Here we investigated the relationships between dissolved inorganic nutrients, phytoplankton pigment composition, microbial community structure, nitrogen fixation rates and the δ¹⁵N of fractionated zooplankton samples along the shelf break of Western Australia (32° to 12° S) in September 2012. Bulk nitrogen fixation rates declined from 4.8 nmol l⁻¹ h⁻¹ in the colder and more saline sub-tropical waters at higher latitudes to 1.5 nmol l⁻¹ h⁻¹ in the warmer and fresher Timor Sea at lower latitudes. A regional bloom of *Trichodesmium* was identified between 13° and 9° S in the Timor Sea. *Trichodesmium*-specific N₂ fixation rates were 0.05 ± 0.01 nmol colony⁻¹ h⁻¹. Highest dissolved inorganic nitrogen (DIN) concentrations occurred at the highest NH₄⁺:NO₃⁻ ratios, thereby deviating from the paradigm that greater DIN concentrations come primarily from increased NO₃⁻ through advection, mixing or upwelling. Both the microplankton and nanoflagellate fraction declined significantly in warmer waters, with higher DIN concentrations but decreasing % NO₃⁻. A clear increase in the prokaryotic diagnostic pigment zeaxanthin was seen with increasing temperatures from higher to lower latitudes. The microbial community, measured using automated ribosomal intergenic spacer analysis (ARISA), clustered strongly according to the water mass biogeochemistry including temperature, salinity, DIN and phosphate concentrations (p < 0.001). Isotope analysis suggested that injections of low δ¹⁵N from N₂ fixation lowered the zooplankton δ¹⁵N signature of animals up to ~500 µm in size and that nearly 47% of the fixed nitrogen was used by zooplankton (≤500 µm fraction) in the Timor Sea.

KEY WORDS: Indian Ocean · Nitrogen fixation · Phytoplankton community · Trophic transfer · ARISA · *Trichodesmium*

INTRODUCTION

Since the 1970s, it has been recognized that the supply of new nitrogen (N) via N₂ fixation to the mixed layer is a key factor regulating the fraction of primary production exported from the surface to the deep ocean (Mague et al. 1974, Bryceson & Fay 1981, Capone & Carpenter 1982). Recent work has emphasised the dominant role of N₂ fixation as a source of new N (and new primary production) in stratified and tropical oceanic ecosystems (Capone & Knapp 2007, Großkopf et al. 2012). Yet, the importance of N₂ fixation in the Indian Ocean remains unclear (Waite et al. 2013). Feedback between N₂ fixation and oceanic...
and atmospheric CO₂ coupling may be globally important, as N₂ fixation rates are likely to increase, due to both enhanced stratification (Hood et al. 2004, Breitbarth et al. 2007) and reduced nutrient fluxes over the pycnoclines (Boyd et al. 2010, Garcia et al. 2011). Estimated N₂ fixation inputs to pelagic ecosystems (Westberry & Siegel 2006, Großkopf et al. 2012) have been predicted to increase above 200 Tg N yr⁻¹ under these scenarios (Hutchins et al. 2007).

Diazotrophs can utilize dissolved N₂ as their inorganic N source, thereby reducing their dependence on oceanic nitrate, urea or ammonium for growth. This unique ability makes them a potential long-term winner under putative climate change driven reductions in inorganic N fluxes (Moore & Doney 2007). Recently, the 2 major oceanic unicellular N₂-fixing cyanobacterial groups UCYN-A and Crocosphaera watsonii (group B) have been postulated to have equal or greater abundances and N₂ fixation rates than those of the well-known filamentous cyanobacterium Trichodesmium (Goebel et al. 2010, Moisander et al. 2010). Elevated pCO₂ concentrations and increased aeolian iron inputs resulting from human induced desertification and climate change could increase N₂ fixation rates both for Trichodesmium (35−65%; Hutchins et al. 2007) and for Crocosphaera (24%; Fu et al. 2008). Higher N₂ fixation rates and an increase in abundance predicted for the future ocean make it clear that the diazotrophic community plays a significant role in ocean biogeochemistry.

The microbial community transforming nitrogen compounds from organic to inorganic forms also maintains the productivity of some open-ocean oligotrophic ecosystems (Zehr & Kudela 2011). N₂ fixation followed by nitrification within the euphotic zone (Wankel et al. 2007) has been shown to be an important biologically mediated pathway in unproductive and oligotrophic regions (Ward et al. 1989, Yool et al. 2007) such as open-ocean sub-tropical gyres (Mather et al. 2008), enclosed seas like the Mediterranean (Rees et al. 2006) and the Indian Ocean (Thompson et al. 2011b, Waite et al. 2013).

The contribution of planktonic diazotrophs to the overall N budget is a key unknown in the eastern Indian Ocean, which is one of the most under-studied but strategically important sectors of the globe (Alexander et al. 2012). Our data are the first to (1) quantify N₂ fixation rates along the entire west coast of Australia and (2) estimate a first-order regional transfer of low δ¹⁵N, derived from N₂ fixation, into higher trophic levels based on local data in the eastern Indian Ocean. We investigate how sub-micro-molar concentrations and ratio shifts (NH₄⁺:NO₃⁻) of dissolved inorganic nitrogen (DIN) mediated by the heterotrophic microbial community impacted the phytoplankton community structure in the relatively warm and oligotrophic waters of the west coast of Australia. In this manuscript, we highlight core biogeo graphic differences between higher- and lower latitude waters around the periphery of an oligotrophic gyre.

**MATERIALS AND METHODS**

Our oceanographic study was conducted in the eastern Indian Ocean aboard the RV ‘Southern Surveyor’ from 6 to 16 September 2012 (early austral spring) along the continental shelf of Western Australia from Fremantle (33°S) to Darwin (13°S), covering 1840 nautical miles (Fig. 1a). Continuous underway measurements of sea surface temperature (T in °C) and salinity (S in PSU) were recorded with a SBE 3T Seabird TSG sensor. Water masses were analysed based on their T and S properties according to Wijffels et al. (2002), Domingues et al. (2007) and Waite et al. (2013) and compiled into 3 groups: (1) Sub-Tropical Waters (STW): from 33° to 26.2°S, with T < 21°C and S > 35; (2) East Gyral + Leeuwin Current Waters (EGC+LC): from 26.2° to 16.3° S, with T between 21 and 25.5°C and S ~34.55; and (3) Timor Sea Waters (TS): from 16° to 12°S, with T > 25.5°C and S < 34.35. The horizontal current velocity along the ship’s track was measured using an RDI 75 kHz Ocean Surveyor acoustic Doppler current profiler (ADCP). The ADCP was set to record to a maximum water column depth of 300 m, and data were averaged in 8 m depth bins. Processing of ADCP data for this voyage was done with the python version of the University of Hawaii’s CODAS system (‘pycurrents’). Underway water samples for dissolved inorganic nutrients, chlorophyll a (chl a) concentrations, pigment analysis using high performance liquid chromatography (HPLC), microplankton taxonomy, microbial diversity using automated ribosomal intergenic spacer analysis (ARISA) and N₂ fixation rates were obtained from a continuous flow-through supply 6 m below the surface.

**Nutrient analysis**

Dissolved inorganic nutrient concentrations including phosphate (PO₄³⁻), nitrate (NO₃⁻), nitrite (NO₂⁻) and ammonium (NH₄⁺) were assayed on a Bran + Luebbe AA3 HR segmented flow analyser, following
standard spectrophotometric methods (Hansen & Koroleff 2009). NO$_3^-$ and NO$_2^-$ were analysed per AA3 method G-172-96 rev15 according to Armstrong et al. (1967) and Grasshoff et al. (2009), with detection limits to 0.015 µmol l$^{-1}$ using 10 mm flow cells and an LED lamp. Unless elsewhere stated, NO$_3^-$ is the sum of NO$_3^-$ + NO$_2^-$. Phosphate was analysed per AA3 method G-172-96 rev15 according to Murphy & Riley (1962), with detection limits to 0.01 µmol l$^{-1}$ using 10 mm flow cell and an LED lamp. NH$_4^+$ was analysed per AA3 method G-172-96 rev15 according to Kérouel & Aminot (1997), with detection limits to 0.004 µmol l$^{-1}$ using a JASCO fluorometer FP-2020. Dilution series of standard stock solutions were made on the day of analysis. All calibration curves had an $r^2$ of 0.999. Coefficients of variation were 0.2% for NO$_3^-$ and NO$_2^-$, 0.4% for PO$_4^{3-}$ and 0.2% for NH$_4^+$.

Pigment analysis

For chl a analysis, 0.525 l of sample water was gently filtered via vacuum filtration (pressure drop <10 kPa) on 25 mm Whatman GF/F filters. Chl a extractions were carried out following acidification according to Parsons et al. (1984). Samples were measured on a Turner Trilogy laboratory fluorometer. For HPLC analysis, 4 l water samples were filtered (pressure drop <10 kPa) on 25 mm Whatman GF/F filters and stored in liquid nitrogen until further analysis. Pigments were determined using HPLC according to the Commonwealth Scientific and Industrial Research Organisation (CSIRO) methods (see Chapter 2 in Hooker et al. 2012). Processed HPLC data were analysed using the diagnostic pigments of the dominant phytoplankton taxa groups (Vidussi et al. 2001, Hirata et al. 2008, Aiken et al. 2009). Pigment data were quality controlled according to Aiken et al. (2009). Our data showed that: (1) the total chl a (TChl a) made up at least 70% of the total pigment concentration and (2) we had a regression between TChl a and the accessory pigments with a slope of 1.1 and $r^2$ > 0.9. One should note that diagnostic pigment analysis has its ambiguities (e.g. fucoxanthin is a precursor for 19'-butanoyl oxyfucoxanthin and 19'-hexanoyloxyfucoxanthin); for more detail, see Aiken et al. (2009) and Uitz et al. (2006). However, this had only minor implications for our data, as the microplankton represented a small fraction of the total phytoplankton functional types (fraction range: 0.04–0.09).

Picoplankton analysis

Seawater samples for picoplankton analysis were collected and preserved with molecular grade paraformaldehyde (final concentration 1%) and stored at −80°C (Marie et al. 1997, Gasol & Giorgio 2000). Microbial populations were separated into heterotrophs, Synechococcus-like and Prochlorococcus-like using a Beckman Coulter Gallios flow cytometer.
equipped with 488 nm and 638 nm lasers (Liu et al. 2004). Emissions were measured using bandpass filters corresponding to emissions for chl a (em: 680; filter: 695/30), phycoerythrobilin (em: 630; filter: 660/30), phycourobilin (em: 577; filter: 575/30) and SYBR® Green I nucleic acid stain (Molecular Probes; em: 520; filter: 525/30). DNA staining was preformed according to Marie et al. (1997), with slight modifications. Samples were amended with 30 mM tripotassium citrate and SYBR® Green I nucleic acid stain (Molecular Probes) 1:10 000 and then incubated for 15 min at 35°C. Beckman Coulter (cat. no. A91346) Flow-Count™ Fluorospheres were added to each sample for determination of the volume sampled as well as normalization of fluorescence intensity. Post processing determination of groups was performed with Kaluza Flow Cytometry software v1.2 (Beckman Coulter) using various gates and bplots of fluorescence to fully separate microbial sub populations. Samples were initially separated into heterotrophs and autotrophs using forward scatter and chl a fluorescence. Prochlorococcus can be difficult to separate from heterotrophs (Marie et al. 1997), but we successfully separated the 2 populations by analysing the initial heterotroph group in a subsequent bplot using phycoerythrin versus SYBR® fluorescence. Phycourobilin (PUB) versus phycoerythrobilin (PEB) bplots were used to separate autotrophs into the following subgroups: Synechococcus of high PUB:PEB, Synechococcus of low PUB:PEB, Synechococcus with PEB only and Prochlorococcus. Subsequent analyses of these subgroups were performed by plotting these on individual bplots using the various combinations of fluorescence; if separation occurred, these were gated and quantified separately.

Microplankton taxonomy

The samples for microplankton taxonomy were collected at midnight and sunrise into 0.5 l high-density polyethylene jars, preserved with a Lugol’s iodine solution of 7.5% (Parsons et al. 1984) and stored in the dark until onshore analysis. Onshore, the samples were homogenised by inverting the jars 50 times, after which a whole subsample (50 or 60 ml) was counted and analysed under a Leica (DM IRB) inverted microscope at 200× magnification after settling for ~24 h (Utermöhl 1958). The cell count varied from 352 to 896 cells per subsample, with a total of 43 different species. Each cell was identified to genus level (or closest taxonomic level possible) using identification manuals and books (e.g. Ricard 1987, Tomas et al. 1997, Suthers & Rissik 2009). Bio-volume of enumerated cells was calculated using basic geometric formulas that were assigned to the different cell shapes in the study of Hillebrand et al. (1999). Where genus-specific calculations were not possible, shapes and measurements were assigned according to taxonomic relatives of the organism and/or to the closest shape according to photos and/or identification books used. Although some cells are known to shrink in Lugol’s fixative, this was not taken into account in the bio-volume calculations due to uncertainties of shrink factors for different cells. Carbon content was calculated from the bio-volume using conversion factors. Conversion factors were obtained from Putt & Stoecker (1989) for ciliates, from Lessard (1991) for dinoflagellates, from Caron et al. (1995) for sarcodines and from Strathmann (1967) for diatoms and silicoflagellate species.

Microbial diversity

Water samples (2 l each) for ARISA (Fisher & Triplett 1999, Danovaro et al. 2006) were filtered through 0.2 µm Sterivex filters and snap frozen in liquid nitrogen for further analysis. A modified organic (phenol:chloroform:isoamyl based) DNA extraction protocol was used alongside extraction columns from the PowerWater DNA isolation kit (Mo Bio Laboratories) (S. Appleyard pers. comm.), with DNA eluted in 80 µl TE buffer. Following sample extraction, DNA was amplified using universal bacterial primers 16S-1392F (5’-GYA CAC ACC GCC GT-3’) and a 5’HEX-labelled version of the reverse primer 23S-125R (5’-GGG TTB CCC CAT TCR G-3’) which amplified the ITS1 region in the rRNA operon plus approximately 282 bases of the 16S and 23S rRNA genes (Hewson & Fuhrman 2004). PCR amplification conditions included 50 µl volumes with 20 ng of DNA, 25 µl of GoTaq® Green Master Mix (containing Taq DNA polymerase and dNTPs; Promega), 2 µl of bovine serum albumin (Promega) and 1 µl of each of the 10 µmol 1−1 primers and water. To monitor extraneous contamination, negative controls containing the PCR mix but without any DNA template were run during each PCR. PCR amplification consisted of an initial denaturation at 94°C for 3 min, followed by 35 cycles of denaturation at 94°C for 1 min, annealing at 55°C for 1 min 30 s and extension at 72°C for 1 min 30 s, with a final extension of 72°C for 10 min in a GeneAmp PCR system 9700 (Applied Biosystems) thermocycler. PCR products were checked on 2.5%
Determination of were weighed and packed into Sercon tin capsules. using mortar and pestle. Subsamples of 0.5 to 0.7 mg dried a second time overnight and homogenised colonies in any of the fractions. Fractionated plankton samples were dried overnight at 60°C, acidified, and great care was taken to exclude potential Trichodesmium colonies in any of the fractions. Fractionated plankton samples were dried overnight at 60°C, acidified, dried a second time overnight and homogenised using mortar and pestle. Subsamples of 0.5 to 0.7 mg were weighed and packed into Sercon tin capsules. Determination of δ15N and δ13C was carried out using a continuous flow system consisting of a Sercon 20-22 mass spectrometer connected with an Automated Nitrogen Carbon Analyser (ANCA, Sercon). Stable isotope abundances are expressed as the deviation from standards in parts per thousand (%). Multi-point normalization was used in order to reduce raw values to the international scale (Paul et al. 2007). Standard selection for δ13C was as described by Skrzypek et al. (2010).

Zooplankton δ15N and δ13C

Fractionated zooplankton samples were obtained from surface Neuston net tows while steaming at 3 knots for 15 min. The Neuston net had a width of 1 m, height of 0.2 m and a mesh and cod end size of 333 µm. Upon retrieval, samples were sieved sequentially through mesh sizes of 4000, 2000, 1000, 500, 300 and 100 µm. Handpicked samples from individual mesh sizes were thoroughly rinsed, and great care was taken to exclude potential Trichodesmium colonies in any of the fractions. Fractionated plankton samples were dried overnight at 60°C, acidified, dried a second time overnight and homogenised using mortar and pestle. Subsamples of 0.5 to 0.7 mg were weighed and packed into Sercon tin capsules. Determination of δ15N and δ13C was carried out using a continuous flow system consisting of a Sercon 20-22 mass spectrometer connected with an Automated Nitrogen Carbon Analyser (ANCA, Sercon). Stable isotope abundances are expressed as the deviation from standards in parts per thousand (%). Multi-point normalization was used in order to reduce raw values to the international scale (Paul et al. 2007). Standard selection for δ13C was as described by Skrzypek et al. (2010).

N2 fixation rates

We used the ‘dissolution method’ (Mohr et al. 2010) rather than the ‘bubble method’ (Montoya et al. 1996) to estimate N2 fixation rates. Buffered YBCII medium (Chen et al. 1996), without addition of vitamins, was filtered (0.2 µm Sterivex filter) prior to degassing (using a heating and stirring block). YBCII medium was transferred into 3 l gas-tight Tedlar bags. Oxygen concentrations were checked prior (8.24 mg l−1) and post degassing (3.0 mg l−1) to check for completeness of degassing. Tedlar bags were spiked with 0.8 ml 15N2 (98 atom%; Aldrich) gas per 100 ml YBCII media and shaken vigorously until 15N2 gas dissolved into the media. Incubations started by introducing aliquots of 15N2 tracer-enriched seawater (2.6% of the total incubation volume) to the pre-acid washed polycarbonate incubation bottles (4.57 l). Spiked incubation bottles were gently rocked 50 times to enhance mixing of the enriched and incubation water. Incubation bottles were incubated for 24 h in an on-deck incubator under ambient light conditions and sea surface temperatures. Trichodesmium-specific N2 fixation rates were conducted on hand-picked colonies. Fifty isolated colonies were incubated in 1.25 l bottles with 50 ml 15N2 tracer-enriched seawater. Individual Trichodesmium colony sizes were not determined, as our methodology was based on earlier studies of Capone et al. (2005) and Mulholland et al. (2006), where colony-specific N2 fixation rates were reported without size-specific colony measurements. Orcutt et al. (2001) also noted that although the number of trichomes per colony can vary, the average N content of puff and tuft colonies at the Bermuda Atlantic Time-series Study (BATS) site remained around 1.8 µg per colony, a value close to that reported by Letelier & Karl (1996) and McCarthy & Carpenter (1979). Colony-specific N2 fixation rates in our study were calculated based on an average of 200 trichomes per colony, a value that has been used by McCarthy & Carpenter (1979), Carpenter (1983) and Orcutt et al. (2001). Final micro-nutrient additions in the incubation bottles were in the order of nano- and picomol l−1 (Table A1 in the Appendix). Experiments were ended by collecting the suspended particles from each bottle by gentle vacuum filtration (pressure drop <10 kPa) through a 25 mm precombusted GF/F filter. Filters were snap frozen in liquid nitrogen and stored at −80°C while at sea. Back in the laboratory on land, filters were acidified and dried overnight at 60°C. Analysis of the 15N2 incorporation into the particulate organic matter (POM) was carried out using a continuous flow sys-
tem consisting of a Sercon 20-22 mass spectrometer connected with an ANCA (Sercon) at the West Australian Biogeochemistry Centre at the University of Western Australia (Paul et al. 2007, Skrzypek et al. 2010). Final $^{15}\text{N}$ atom enrichment was calculated assuming complete dissolution of the gas bubble. The solubility of the N$_2$ concentration in seawater was calculated according to Hamme & Emerson (2004) for each individual station.

**Statistical analysis and mixing model**

Principal component analysis (PCA) was used to explore the transect dataset and to reduce the number of variables in the data matrix. Statistical analyses were performed in R statistical package v3.0.1 (R Development Core Team 2013). Two- and 1-way ANOVA tests were conducted using the car-package (Fox & Weisberg 2009) for R with a linear regression model, based on a heteroscedasticity consistent covariance matrix, as sample size was smaller than 250 (Long & Ervin 2000).

The structural similarity of ARISA datasets was assessed on standardised and transformed data using Bray-Curtis similarity matrices (Spearman correlation, 9999 permutations) and Mantel tests using Primer 6. Microplankton data were assessed using canonical analysis of principal coordinates (CAP) in Primer 6. Permutational multivariate ANOVA (PERMANOVA; Anderson et al. 2008) tests were conducted in Primer 6.

We applied the concept proposed by Montoya et al. (2002) that injections of low $^{15}\text{N}$ from N$_2$ fixation can lower the zooplankton $^{15}\text{N}$ signature. A simple linear isotope mixing model was used in this study to estimate a first-order transfer of low $^{15}\text{N}$ into higher trophic levels based on local data only. For the 2 end points of our model, we used (1) the $^{15}\text{N} = -1.6\%$ from the *Trichodesmium* colonies to represent cyanobacteria N$_2$ fixation and (2) a regional mean (±SD) of $^{15}\text{NNO}_3^- = 6.6 \pm 0.7\%$ for the deep NO$_3^-$ pool in the STW, EGC+LC and TS waters derived from Waite et al. (2007a) and Waite et al. (2013). The latter is similar to oligotrophic oceanic NO$_3^-$ signatures found in a number of studies globally (Sigman et al. 1997, Brandes et al. 1998, Sigman et al. 2005).

**RESULTS**

The transect northwards along the continental shelf of Western Australia (300–1000 m depth contours) showed an increase in temperature (19.9 to 27.6°C) and a decrease in salinity (35.36 to 34.04) from 31.6° to 12.7°S (Fig. 1a,b). We traversed the colder and more saline STW in the south (high latitudes) and then entered the warmer and fresher TS in the north (low latitudes). ADCP data revealed a southward flow dominating the transect, as expected given the dominance of the Leeuwin Current (LC), with greater current speeds at lowest and highest latitudes (30° and 12°S, 0.5 m s$^{-1}$) and lowest speeds around 21°S (0.2 m s$^{-1}$). Current flows and speeds were as anticipated for the time of the year (Thompson 1984).

**Nutrient analysis**

NH$_4^+$ concentrations increased going to lower latitudes ($r^2 = 0.184$, slope 0.0058, $p = 0.0026$, $n = 47$). NO$_3^-$ (NO$_3^-+\text{NO}_2^-)$ concentrations also increased going to lower latitudes ($r^2 = 0.204$, slope 0.0022, $p = 0.0014$, $n = 47$), although NO$_2^-$ concentrations on their own, with an average of 0.02 ± 0.02 µmol l$^{-1}$ (± SD), showed no significant latitudinal trend ($r^2 = 0.000023$, $p = 0.975$, $n = 47$). Two stations (D2 and D6) showed relative higher NO$_3^-$ and NH$_4^+$ values in the STW (Fig. 2a,b). Both stations also had the highest NO$_2^-$ concentrations (0.11 and 0.07 µmol l$^{-1}$ for D2 and D6, respectively). The steeper slope for NH$_4^+$ ($0.0058 \pm 0.0018$ SE µmol l$^{-1}$ per degree latitude) compared to NO$_3^-$ ($0.0022 \pm 0.0006$ µmol l$^{-1}$ per degree latitude) indicated a significant increase of 21% of NH$_4^+$ to the total DIN pool ($r^2 = 0.261$, $p < 0.001$, $n = 47$). Although the absolute NO$_3^-$ concentrations increased towards lower latitudes, the corresponding % contribution of NO$_3^-$ to total DIN showed a significant decrease by 14% towards lower latitudes ($r^2 = 0.249$, $p < 0.001$, $n = 47$). NH$_4^+$ and NO$_3^-$ are both driving the increase in DIN at lower latitudes; however, NH$_4^+$ does so to a larger extent. As a result, highest DIN concentrations occurred at highest NH$_4^+$/NO$_3^-$ ratios. Total DIN concentrations were significantly different between the STW (higher latitude) and TS (lower latitude) waters (1-way ANOVA $p < 0.05$). No significant differences were found between the EGC+LC and STW or for the EGC+LC and TS (1-way ANOVAs $p > 0.05$, see Table 1). Average ±SE PO$_4^{3-}$ concentrations for the whole transect were 0.11 ± 0.002 µmol l$^{-1}$ ($n = 47$), with no significant latitudinal increase ($r^2 = 0.039$, $p = 0.056$, $n = 47$; Fig. 2c). Overall, the N:P ratio increased going to lower latitudes. Two southern stations (D2 and D6) showed outliers with relative higher N:P ratios in the STW.
N:P values were $1.8 \pm 1.6 < 1.9 \pm 0.8 < 2.9 \pm 1.0$ for the STW, EGC+LC and TS, respectively. N:P ratios were only significantly different between the STW and TS (1-way ANOVA, $p < 0.05$). Although N:P ratios nearly doubled going north, the values were still lower than the Redfield ratio of 6.6:1 (Redfield 1958), such that overall the West Australian coast could still potentially be N limited. A PERMANOVA (testing all the nutrient variables against the individual water masses) showed significant differences between all water masses (STW and EGC+LC, $p = 0.034$; EGC+LC and TS, $p = 0.007$; TS and STW, $p = 0.003$). Temperature, salinity, nutrient and N:P ratio data for the transect are summarized in Table 1.

**Pigment analysis**

A pairwise multiple comparison (Dunn’s Method) between the water masses showed significantly higher chl $a$ concentrations in the STW compared to the TS ($p < 0.05$), higher chl $a$ concentrations in the STW compared to the EGC+LC ($p < 0.05$) but no difference between the EGC+LC and TS ($p > 0.05$; Fig. 2d). Relatively higher inorganic N:P ratios coincided with low chl $a$ values along the transect. Chl $a$ data for the transect are summarized in Table 1.

The photosynthetic microplankton component, defined by the diagnostic pigments of the dominant phytoplankton taxa groups (see 'Materials and methods'), showed a declining trend towards lower latitudes ($r^2 = 0.708$; slope $-0.0047 \pm 0.0007$ SE; $p < 0.0001$), corresponding to warmer waters and declining NO$_3^-$ concentrations (Fig. 3a). The microplankton community declined $-0.4\%$ per 1° latitude north ($-111$ km) in the STW. This declining trend increased farther north to $-0.5\%$ per 1° latitude for the EGC+LC and $-1.6\%$ per 1° latitude for TS. Pairwise multiple comparison (Tukey’s tests) for the microplankton showed significant differences between STW and EGC+LC ($p < 0.001$) and between the STW and TS ($p < 0.001$). No significant differences were found between EGC+LC and TS ($p = 0.234$; Table 2). The nanoflagellate component, although to a much...
greater extent than the microplankton component, showed a similar declining trend towards lower latitudes ($r^2 = 0.851; \text{slope} -0.0279 \pm 0.0028 \text{SE}; p < 0.0001$). This amounted to a $-2.4\%$ per 1° latitude for the STW and $-2.8\%$ per 1° latitude for both the EGC+LC and TS (Fig. 3b). The median nanoflagellate proportions showed significant differences between all water masses (Tukey's tests: $p < 0.0001$ between the STW and TS, $p < 0.0001$ between the STW and EGC+LC and $p = 0.005$ between the EGC+LC and TS; Table 2). A clear increase in the prokaryotic community was seen from higher to lower latitudes ($r^2 = 0.848; \text{slope} 0.0326 \pm 0.0032 \text{SE}; p < 0.0001$), with increasing temperatures and increasing $\text{NH}_4^+$ concentrations. This revealed an increase of $2.8\%$ per 1° latitude for the high-latitude water mass, and a $3.3\%$ increase per 1° latitude for the low-latitude water masses, respectively (Fig. 3c). The median prokaryotic proportions were significantly different between all the water masses (Tukey's tests: $p < 0.0001$ between STW and TS, $p < 0.0001$ for STW and EGC+LC and $p = 0.005$ for EGC+LC and TS; Table 2). A PERMANOVA, combining the microplankton, nanoflagellate and prokaryote abundance, showed significant differences between STW and EGC+LC waters ($p = 0.004$), between STW and TS ($p = 0.001$) and between EGC+LC and TS ($p = 0.004$).

### Picoplankton analysis

Flow cytometry data showed that *Prochlorococcus* spp. were least abundant in cooler, sub-tropical waters ($0.5 \times 10^8$ cells l$^{-1}$ at 31° S) but showed a steep increase (to $4 \times 10^8$ cells l$^{-1}$ at 20° S) around DIN concentrations of 0.3 µmol l$^{-1}$ in the EGC+LC. Although DIN concentrations were below 1 µmol l$^{-1}$, *Prochlorococcus* spp. concentrations started to tail off around 0.5 µmol l$^{-1}$. Overall *Synechococcus* spp. abundance was greater than *Prochlorococcus* spp. abundance ($0.5-6.5 \times 10^8$ cells l$^{-1}$). The *Synechococcus*:Prochlorococcus ratio was lowest in STW. This ratio veered further into 2 different relationships, one with a low and one with a high *Synechococcus*:Prochlorococcus ratio. A Bray-Curtis similarity test showed a clear separation of 2 different possible *Synechococcus* and *Prochlorococcus* clades in all water masses (Fig. 4). There was no significant trend for the prokaryotic community, which included all non-phototrophs and was derived from the SYBER® stain, with latitude ($r^2 = 0.0019, p = 0.84, n = 24$) or among the individual water masses (1-way ANOVA: $p = 0.47$). There was also no significant trend in the prokaryotic and autotrophic (*Prochlorococcus* + *Synechococcus* + pico-eukaryotes) ratio with latitude ($r^2 = 1.3 \times 10^{-5}, p = 0.98, n = 24$) or among the water masses (1-way ANOVA: $p = 0.66$).

### Table 1. Temperature, salinity, nutrient, chl $\alpha$ and N:P ratio data for the latitudinal transect summarized according to water masses (STW: Sub-Tropical Waters, EGC+LC: East Gyral Current + Leeuwin Current; TS: Timor Sea). Values are means ± SD. DIN: dissolved inorganic nitrogen.

<table>
<thead>
<tr>
<th>Latitude (°S)</th>
<th>Water mass</th>
<th>Temp (°C)</th>
<th>Salinity (PSU)</th>
<th>NO3$^-$ (µmol l$^{-1}$)</th>
<th>NH4$^+$ (µmol l$^{-1}$)</th>
<th>PO4$^{3-}$ (µmol l$^{-1}$)</th>
<th>Chl $\alpha$ (µg l$^{-1}$)</th>
<th>N:P</th>
<th>DIN (µmol l$^{-1}$)</th>
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<tr>
<td>33.2−26.2 (n = 11)</td>
<td>STW</td>
<td>$&lt;21$</td>
<td>35.18 ± 0.13</td>
<td>0.083 ± 0.033</td>
<td>0.092 ± 0.063</td>
<td>0.864 ± 0.032</td>
<td>0.110 ± 0.014</td>
<td>0.02</td>
<td>0.150 ± 0.130</td>
</tr>
<tr>
<td>26.3−16.3 (n = 23)</td>
<td>EGC+LC</td>
<td>21−25</td>
<td>34.71 ± 0.22</td>
<td>0.092 ± 0.063</td>
<td>0.108 ± 0.023</td>
<td>0.264 ± 0.061</td>
<td>0.100 ± 0.006</td>
<td>0.005</td>
<td>0.171 ± 0.034</td>
</tr>
<tr>
<td>16−12 (n = 13)</td>
<td>TS</td>
<td>$&gt;25.5$</td>
<td>34.25 ± 0.11</td>
<td>0.110 ± 0.022</td>
<td>0.154 ± 0.063</td>
<td>1.84 ± 0.092</td>
<td>0.284 ± 0.063</td>
<td>0.150</td>
<td>0.150 ± 0.150</td>
</tr>
</tbody>
</table>

$^a$ $p < 0.05$, 1-way ANOVA between STW and TS

$^b$ $p < 0.05$, 1-way ANOVA between EGC+LC and TS

$p < 0.005$, 1-way ANOVA between EGC+LC and TS
Microplankton taxonomy

Microscopic analysis revealed that the microplankton community compromised 26 families and 2 higher taxonomic levels from the kingdom Rhizaria. The Dinophyceae including large gymnoids and other Dinophyceae (e.g. Gyrodinium spp., Prorocentrum spp.) were the most abundant microplankton, accounting for more than 50% of the microplankton bio-volume at all stations. A CAP did not reveal a clear clustering of microplankton composition, but a more gradual trend, as the water masses gradually changed in temperature and salinity with latitude (Fig. 5). Gyrodinium spp. clearly correlated with different water masses to their closely related gymnoid species. Although temperature increased and salinity decreased going farther north, there was no significant trend for carbon contribution from microplankton across the latitudinal gradient. Most of the carbon contribution came from the large gymnoids, except for stations at 26 and 24° S, where other microplankton (Rhizaria, Chrysophyceae, Euglenophyceae, Oligotrichida and Choreotrichia) made up the major carbon biomass. This was caused by a higher number of Strombidium spp. there than at other stations. During our study, a regional bloom of the N₂-fixing cyanophyte Trichodesmium was identified as distinctive yellow-orange surface slicks between 13 and 9° S. This was a regional-scale feature during the voyage (~50 × 50 km; Fig. 6a,b).

Microbial diversity

ARISA distinguishes microbial communities based on differences in length heterogeneity in the ribosomal intergenic spacer region between the 16S and
23S rRNA genes in the rRNA operon. The ARISA analyses enabled us to directly and rapidly compare the microbial diversity and community composition along a latitudinal gradient in this study. The structural similarity of ARISA datasets were assessed on standardised (by total) and transformed (square root) data using Bray-Curtis similarity matrices and showed a strong difference between STW, EGC+LC and TS. A Pearson rank vector overlay (0.56) highlighted the importance and close relationship between microbial community structure and NH4+ at low latitudes and NO3− and PO43− concentrations at high latitudes (Fig. 7). A Mantel test confirmed the significant relationship between the microbial community structure and water mass biogeochemistry including temperature, salinity, DIN and phosphate concentrations (Mantel test, p < 0.001). A PERMANOVA, combining all the operational taxonomical units of the ARISA data, showed a significant difference between STW and TS (p = 0.013) but no significant difference between STW and EGC+LC (p = 0.124) or between EGC+LC and TS (p = 0.834).

**N2 fixation rates**

Nitrogen fixation was detected along the entire West Australian coastline. Average ± SD bulk N2 fixation rates in the STW were 4.78 ± 1.16 nmol l−1 h−1 (n = 7), 3.18 ± 1.33 nmol l−1 h−1 (n = 16), in the EGC+LC waters and 1.48 ± 0.65 nmol l−1 h−1 (n = 8) in the TS (Fig. 8a). Median N2 fixation rates showed a significant decline from high to low latitudes. All water masses were distinctively different (1-way ANOVAs: STW and TS, p < 0.001; STW and EGC+LC, p = 0.014; TS and EGC+LC, p = 0.006). Colony-specific mean ± SD N2 fixation rates were 0.05 ± 0.01 nmol colony−1 h−1 (n = 3) in the TS. These rates compare to those reported by Carpenter & Price (1977), McCarthy & Carpenter (1979) and Orcutt et al. (2001). Capone et al. (1998) recorded up to 17 000 trichomes l−1 in surface slicks in the northern Indian Ocean. When each colony is assumed to have 200 filaments (McCarthy & Carpenter 1979), we calculated 85 colonies l−1, resulting in *Trichodesmium*-specific N2 fixation rates of 4.62 ± 1.36 nmol l−1 h−1. These rates are 3 times higher compared to the bulk
N₂ fixation rates (1.48 ± 0.65 nmol l⁻¹ h⁻¹) at the same latitude. Yet bulk N₂ fixation rates in the STW match *Trichodesmium*-specific N₂ fixation rates.

Zooplankton δ¹⁵N and δ¹³C

The zooplankton δ¹⁵N values generally increased with size fraction but decreased in TS in proximity to the *Trichodesmium* bloom (Fig. 8b). To determine whether these differences and their primary covariates were significant, we used PCA. Variables included temperature, phosphate, chl a concentrations, fractionated zooplankton samples and prokaryotic diagnostic pigments. The PCA indicated that the first 2 PCs explained 86.26% of the total variance (67.44% and 18.82%, respectively). The main contributing variables to PC1 included temperature and prokaryotic community abundance (0.512 and 0.503 loadings, respectively). PC2 showed a significant positive loading with the zooplankton δ¹⁵N size fractions (0.878) and a negative loading, although to a much lesser extent, of chl a concentrations (−0.350). PC1 indicated significant negative correlations between temperature and prokaryote concentration (−0.458), and between chl a and phosphate concentrations (−0.469). Conversely, chl a concentrations decreased from high to low latitudes. PC2 highlights the separation between small-sized zooplankton (≤500 µm) caught between 12.4° and 12.7° S and the rest of the data set (Fig. 8a). These samples were found to have negative δ¹⁵N compositions, possibly indicating the presence of an N₂-fixing community that is consumed as a food source.
A simple linear isotope mixing model with 2 end points showed that 80 ± 4% of the 100 µm fraction in the northern latitudes is likely to be grazing, directly or indirectly, on a pool of low δ¹⁵N (derived from N₂ fixation).

The fractional contribution from N₂ fixation declined towards greater size classes (higher trophic levels) to 79 ± 2% for the 300 µm and 47 ± 6% for the 500 µm fraction in the northern latitudes. At the higher latitudes, the zooplankton size classes (up to 500 µm) are all over 94% based on a δ¹⁵N pool derived from deep NO₃⁻ (see Table A2 in the Appendix).

**DISCUSSION**

In our study, relative high DIN concentrations in the south eastern Indian Ocean occurred at the highest NH₄⁺ concentrations, differing from the classic paradigm where large DIN concentrations come primarily from NO₃⁻ via mixing or classic upwelling (Kokkinakis & Wheeler 1987). While new production is always enhanced by the vertical flux of NO₃⁻, the export f-ratio according to Dugdale & Goering (1967) is no longer valid when the highest DIN concentrations occur at highest NH₄⁺:NO₃⁻ ratios. This is likely to occur when N₂ fixation dominates new N input (Gruber & Sarmiento 1997, Großkopf et al. 2012) and/or when nitrification occurs within or near the euphotic zone (here referred to as ‘shallow nitrification’; see Lipschultz et al. 2002, Yool et al. 2007 and Thompson et al. 2011b).

Latitude itself does not control ecological processes, yet in this study we described biological pathways along a spatial gradient as a suite of physical and biogeochemical parameters strongly correlated with latitude. We show that the total DIN concentrations increased significantly at lower latitudes, and that the highest DIN concentrations occurred at the highest NH₄⁺:NO₃⁻ ratios. The latitudinal NO₃⁻ and NH₄⁺ concentration and ratio shift observed during our brief snapshot is accompanied by a gradient in phytoplankton community structure from a micro- and nanoflagellate-dominated ecosystem to a prokaryotic system heading equatorward, despite the apparent increase in available N. The 3.5-fold increase in the prokaryotic diagnostic pigment zeaxanthin (representative of prokaryotic cyanobacteria) from higher to lower latitudes is similar in scale to the 5-fold increase (across similar latitudes) reported by Thompson et al. (2011a). Injection of new N as NH₄⁺ in northern waters is potentially subjected to rapid uptake and recycling. This, in combination with rapid growth rates at higher temperatures (Eppley 1972), is likely to lead to much more rapid processing of N by the lower-latitude planktonic community. The ubiquitous dominance of large non-photosynthetic gymnoids (Dinophyceae) in the microplankton along the entire West Australian shelf suggests that prokaryotes are always subject to high grazing rates (Paterson et al. 2007), although their exact role requires further exploration.

The conceptual model postulated by Thompson et al. (2011b) links the source of NH₄⁺ concentrations at higher latitudes to an increase in grazing pressure after/during the austral autumn bloom when NO₃⁻ concentrations are higher (Alcaraz et al. 1994). Injections of NO₃⁻ occur via erosion of the pycnocline during the autumn acceleration of the Leeuwin Current (Koslow et al. 2008, Rousseaux et al. 2012). Waite et al. (2013) suggested that these increased NH₄⁺ concentrations can be derived, at least in part, from N₂ fixation which is in fact a more likely and continuous source of new N, especially near the surface. Evidently both processes are not mutually exclusive. However, in this study we were able to confirm the findings of high bulk N₂ fixation rates (4.5 nmol l⁻¹ h⁻¹) at higher latitudes (to 32° S). It is interesting to note that our data showed a latitudinal decrease in bulk N₂ fixation rates towards warmer waters (from 4.5 to 1.24 nmol l⁻¹ h⁻¹). Low bulk N₂ fixation rates in the TS were compensated by the injection of new N from a *Trichodesmium* bloom. Our data show that unicellular diazotrophs declined when *Trichodesmium* was present in warmer, tropical waters, suggesting a diazotrophic community shift when *Trichodesmium* is present. To date we have no additional molecular data to show the presence of other diazotrophs such as diatom–diazotroph associations, coccoid forms or UCYN-A that could be responsible for the injections of new N at these lower latitudes from our transect. However, our HPLC and microscopic data showed a declining trend for the photosynthetic microplankton, and we can therefore, carefully, exclude an increasing presence of any diatom–diazotroph associations in the northern latitudes. We therefore summarize cautiously that the increase of fixation at higher latitudes is likely to be caused by unicellular fixation, while *Trichodesmium* is the prime candidate at lower latitudes. Iron conservation strategies by unicellular diazotrophs have been shown to provide an ecological advantage in low-iron environments of the open ocean (Saito et al. 2011), whereas protein level responses of *Trichodesmium* under iron starvation have been shown to
down-regulate N\textsubscript{2} fixation rates (Richier et al. 2012). The trace level iron additions during our experiments ranged between 10 and 17 nmol l\textsuperscript{-1} and although these concentrations are at the lowest end of those reported by Richier et al. (2012), it is important to keep in mind how iron availability influences the physiology and the adjacent fixation of N\textsubscript{2} of diazotrophs (Berman-Frank et al. 2007, Rubin et al. 2011).

Despite the documented prevalence of *Trichodesmium* in the open ocean since the 19th century (Capone et al. 1997), and recent identification from space of vast surface distributions covering hundreds of kilometres (Subramaniam et al. 2001, McKinna et al. 2011, Waite et al. 2013), it remains unclear exactly how much *Trichodesmium* contributes to new production in the global ocean, particularly in the poorly-studied Indian Ocean. Up-scaling the average bulk N\textsubscript{2} fixation rates from our data gives us insight in the relative importance of the biogeochemical impact of the unicellular diazotrophs and *Trichodesmium* along our transect. Rodier & Le Borgne (2008, 2010) measured *Trichodesmium* densities up to 1.2 × 10\textsuperscript{2} trichomes l\textsuperscript{-1}. For our calculations, we used *Trichodesmium* densities of 17 000 trichomes l\textsuperscript{-1} reported by Capone et al. (1998) in the North Indian Ocean. We calculated an N\textsubscript{2} fixation flux for *Trichodesmium* of 6.9 × 10\textsuperscript{10} µmol h\textsuperscript{-1} for a 50 × 50 km bloom (down to 6 m depth). The relative importance of *Trichodesmium* in the TS becomes a smaller proportion of the total diazotrophic community (i.e. *Trichodesmium* is diluted out) when one begins to upscale and the bloom area is reduced relative to bulk rates. Because bulk N\textsubscript{2} fixation rates declined with latitude, *Trichodesmium* remains a prime candidate for N injection in the Tropics (TS). Yet, our data highlight the overall importance of the unicellular diazotrophs across the study region, particularly in the south. Our high bulk N\textsubscript{2} fixation rates support the postulated hypothesis by Goebel et al. (2010) and Moisander et al. (2010) that oceanic unicellular N\textsubscript{2}-fixing cyanobacteria can have equal or greater N\textsubscript{2} fixation rates than *Trichodesmium*. ‘New’ primary productivity, calculated from the bulk N\textsubscript{2} fixation rates and using a C:N ratio of 6.3 from our data (n = 31), declined from 361 µg C m\textsuperscript{-2} h\textsuperscript{-1} (STW) to 240 µg C m\textsuperscript{-2} h\textsuperscript{-1} (EGC+LC) and to 112 µg C m\textsuperscript{-2} h\textsuperscript{-1} (TS). These data compare to the surface production levels reported by Hanson et al. (2005). Our data show that it is likely that N\textsubscript{2} fixation contributes significantly (up to 50%) to new production in the eastern Indian Ocean. However, intense seasonal and spatial sampling is needed to resolve the different temporal and physical scales at which *Trichodesmium* and the unicellular N\textsubscript{2}-fixing cyanobacteria operate. *Trichodesmium* blooms have been reported to be temporally short (Bar-Zeev et al. 2013) and cover substantially large areas (Subramaniam et al. 2001), but unicellular N\textsubscript{2}-fixing cyanobacteria have larger depth distributions and cover higher latitudes (Moisander et al. 2010).

If the NH\textsubscript{4}\textsuperscript{+} source for the DIN we measured is an injection of organic N into the planktonic ecosystem via N\textsubscript{2} fixation, it essentially enters the system as ‘new’ reduced N. We would then expect the low δ\textsuperscript{15}N signatures from the diazotrophic community to move into the organic matter of higher trophic levels (Montoya et al. 2002). *Trichodesmium* has been identified as a potential species involved in passing the low δ\textsuperscript{15}N signature of fixed nitrogen up the food chain. This is thought to occur directly by mesozooplankton grazing (O’Neil & Roman 1992, 1994), which is mostly done by *Macrosetella* (McKinnon et al. 2003, 2008) and indirectly, via cell leakage and microbial processing and through phytoplankton uptake of the regenerated N of N\textsubscript{2} fixed products (Montoya et al. 2002, Wannicke et al. 2010, Waite et al. 2013). Our linear isotope mixing model suggested that nearly 50% of the fixed nitrogen from *Trichodesmium* is used and incorporated into the ≤500 µm fraction at lower latitudes. The data from this northern part of our voyage revealed that biological N\textsubscript{2} fixation appears to be a plausible mechanism for introducing δ\textsuperscript{15}N-depleted compounds into Indian Ocean food webs regionally across the north-west of Australia. In the southern part of the transect, however, where bulk N\textsubscript{2} fixation rates were higher, the direct biogeochemical impact of fixed-nitrogen transfer into mesozooplankton seemed to be negligible. We postulate that the sub-tropical waters have a longer food web and microbial loop associated with the N\textsubscript{2}-fixing pico-cyanobacteria (see Wannicke et al. 2010). This prolonged foodweb structure is likely to move N into microzooplankton grazers, rather than mesozooplankton. Even when mesozooplankton is able to access this N, the large number of intermediate trophic levels, which consequently lead to more fractionation steps, will have increased the δ\textsuperscript{15}N of the organic matter. The lack of clear transfer of low δ\textsuperscript{15}N into mesozooplankton at the higher latitudes could also be due to a more continuous flux of NO\textsubscript{3}\textsuperscript{-} through eddy pumping (Waite et al. 2007b) and higher selective grazing rates on the larger proportion of microphytoplankton (diatoms and dinoflagellates) we observed by the zooplankton community (Malone 1971, Gaul & Antia 2001, Wannicke et al. 2010). In contrast, in the north we support the hypothesis of a shorter food web, where direct feed-
ing by mesozooplankton on the larger *Trichodesmium* colonies consequently lowers the $\delta^{15}N$ of their organic matter.

Although we lack measurements of $\text{NH}_4^+$ oxidation and nitrification rates, we propose a conceptual hypothesis of food web structure, based on our data and literature data (Fig. 9). The eastern Indian Ocean is a region where surface (mixed layer) $\text{NO}_3^-$ is most often undetectable (Pearce & Pattiaratchi 1999) such that shallow nitrification can measurably structure $\text{NO}_3^-$ concentrations above the pycnocline (Yool et al. 2007, Thompson et al. 2011b, Paterson et al. 2013, Waite et al. 2013). The microbial community from our transect shifted from a distinct genetic grouping (Gr1) in the south associated with deep nutrient fluxes (increase in $\%\text{NO}_3^-$ and $\text{PO}_4^{3-}$) and to another genetic grouping (Gr2) in the north associated with a lower $\delta^{15}N$ of organic matter, increase in $\%\text{NH}_4^+$ and increase in total DIN. This change in microbial community structure is likely to mediate shifts in the N processing of the community (Zehr & Kudela 2011).

In particular, we speculate that nitrification of the $\text{NH}_4^+$ injections is likely to be more important in the northern group (Gr2), as nitrification of POM derived from N$_2$ fixation has been identified in the top 200 m of EGC+LC waters (Waite et al. 2013).

We speculate that the POM, derived from N$_2$ fixation in the north-eastern Indian Ocean, has the potential to be remineralised and transported southward (to higher latitudes) by the Leeuwin Current while undergoing sequential oxidation of $\text{NH}_4^+$ to $\text{NO}_2^-$ and $\text{NO}_3^-$ (Zehr & Ward 2002, Waite et al. 2013). Coupled measurements of $\delta^{14}N$, and $\delta^{18}O$ of $\text{NO}_3^-$ would allow determination of the N cycling processes that occur in the upper ocean, as the process of nitrification is potentially a source of isotopically light N (Wankel et al. 2007). The presence of nitrification genes of the genus *Nitrobacter* sp. at the surface and independent of water masses along the entire visited track (E. J. Raes unpubl.), could support our proposed conceptual model. Remineralisation (Ward 2008) is further optimized when the organic

![Conceptual hypothesis/model showing fixed N with low $\delta^{15}N$ signatures moving south from the original source in the Timor Sea. The dissolved inorganic nitrogen (DIN) data show that highest DIN concentrations occur at highest $\text{NH}_4^+\text{-NO}_3^-$ ratios. Arrows indicating in situ current velocity from Acoustic Doppler Current Profiler (ADCP) data. Colour coding: blue for Sub-Tropical Waters (STW), green for East Gyral + Leeuwin Current (EGC+LC) and red for the Timor Sea (TS). *Trichodesmium* bloom shown by black circle and bathymetry shown on right side of the regional map. Top right panel: Conceptual hypothesis of a possible food web showing the community relationships in the TS. Red boxes highlight major dominant pools. Red arrow highlights the incorporation of low $\delta^{15}N$ into mesozooplankton. Bottom panel shows the conceptual hypothesis of possible relationships in STW. Blue boxes show major pools. Blue arrows from the pico N$_2$-fixing cyanobacteria (Diazos) highlight a possible longer foodweb (more fractionation). Greater nano- and micro-fractions (supported by deep $\text{NO}_3^-$) transfer higher $\delta^{15}N$ into mesozooplankton. Dotted arrow indicates nitrification in the euphotic zone (see Lipschultz et al. 2002, Yool et al. 2007, Thompson et al. 2011b). DON: dissolved organic nitrogen, MLD: mixed layer depth. Note: diagrams are non-exclusive considering there are more than the shown possible N transformations.}
matter has the potential to accumulate at strong interfaces (MacIntyre et al. 1995, Doostmohammadi et al. 2012, Waite et al. 2013) such as low dissolved oxygen high nitrate (LDONH) layers. The deeper boundary of the low salinity layers in these LDONH layers, running along the entire West Australian shelf (Thompson et al. 2011b), could be a hotspot for remineralisation and nitrification of POM. Recent work by Waite et al. (2013) suggested that up to 90% of unassimilated NO$_3^-$ (NO$_3^-$ with isotopically light $\delta^{15}N$) found in surface waters in the northern part of the south-eastern Indian Ocean is likely to have originated from N$_2$ fixation (Waite et al. 2013).

It is clear that more work has to be done to unravel how the diazotrophic community and in situ biogeochemical processes (such as sinking/subduction, decomposition and nitrification of the biomass) interrelate towards primary productivity in the eastern Indian Ocean. It will be important to elucidate the temporal and physical scale of *Trichodesmium* blooms and regional N$_2$ fixation rates of pico-cyanobacteria, as our data showed that NH$_4^+$ injections from N$_2$ fixation and from microbial remineralisation have a potentially profound impact on marine community structure.

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Table A1. Concentration of micronutrients in the YBC II medium and the final micro nutrient additions, when using 2.5% and 4% of the enriched YBC II medium, during the bulk N₂ fixation and *Trichodesmium* incubations

<table>
<thead>
<tr>
<th>Micro-nutrients</th>
<th>YBC II Medium (nmol l⁻¹)</th>
<th>Bulk N₂ fixation (2.6% addition nmol l⁻¹)</th>
<th><em>Trichodesmium</em> N₂ fixation (4% addition nmol l⁻¹)</th>
</tr>
</thead>
<tbody>
<tr>
<td>FeCl₃·6H₂O</td>
<td>4.07 × 10⁻⁴</td>
<td>11.0</td>
<td>17.0</td>
</tr>
<tr>
<td>MnCl₂·4H₂O</td>
<td>2.00 × 10⁻⁵</td>
<td>0.539</td>
<td>0.833</td>
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<tr>
<td>ZnSO₄·7H₂O</td>
<td>4.00 × 10⁻⁶</td>
<td>0.108</td>
<td>0.167</td>
</tr>
<tr>
<td>CoCl₂·6H₂O</td>
<td>2.50 × 10⁻⁶</td>
<td>6.74 × 10⁻²</td>
<td>0.104</td>
</tr>
<tr>
<td>Na₂MoO₄·2H₂O</td>
<td>1.10 × 10⁻⁵</td>
<td>0.297</td>
<td>0.458</td>
</tr>
<tr>
<td>CuSO₄</td>
<td>1.00 × 10⁻⁶</td>
<td>2.70 × 10⁻²</td>
<td>4.17 × 10⁻²</td>
</tr>
</tbody>
</table>

Table A2. Fractional contribution of δ¹⁵N from deep NO₃⁻ (6.6 ± 0.7‰) and the fractional contribution the δ¹⁵N from N₂ fixation (–1.6‰) to the zooplankton size classes. STW: Sub-Tropical Waters, EGC+LC: East Gyral + Leeuwin Current, TS: Timor Sea

<table>
<thead>
<tr>
<th>Water</th>
<th>Size (µm)</th>
<th>δ¹⁵N – Deep NO₃⁻</th>
<th>δ¹⁵N – Diatrophs</th>
</tr>
</thead>
<tbody>
<tr>
<td>STW (n = 1)</td>
<td>100</td>
<td>0.94</td>
<td>0.06</td>
</tr>
<tr>
<td>EGC+LC (n = 3)</td>
<td>100</td>
<td>0.97 ± 0.02</td>
<td>0.02 ± 0.02</td>
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<tr>
<td>TS out (n = 2)</td>
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<td>0.97 ± 0.04</td>
<td>0.03 ± 0.04</td>
</tr>
<tr>
<td>TS in (n = 2)</td>
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<td>0.13 ± 0.04</td>
<td>0.88 ± 0.04</td>
</tr>
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<td>0.97 ± 0.04</td>
<td>0.04 ± 0.04</td>
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<td>EGC+LC (n = 3)</td>
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<td>0.99 ± 0.02</td>
<td>0.01 ± 0.02</td>
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<td>0.92 ± 0.10</td>
<td>0.08 ± 0.10</td>
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<tr>
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<td>0.79 ± 0.02</td>
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<tr>
<td>STW (n = 2)</td>
<td>500</td>
<td>0.94 ± 0.08</td>
<td>0.07 ± 0.08</td>
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<td>EGC+LC (n = 3)</td>
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<td>500</td>
<td>0.53 ± 0.06</td>
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