

ORIGINAL RESEARCH published: 23 March 2017 doi: 10.3389/fmars.2017.00080



N₂ Fixation in the Eastern Arabian Sea: Probable Role of Heterotrophic Diazotrophs

P. Kiran Kumar¹, Arvind Singh^{1*}, R. Ramesh^{1†} and T. Nallathambi²

¹ Physical Research Laboratory, Geosciences Division, Ahmedabad, India, ² National Institute of Ocean and Technology, Chennai, India

Biogeochemical implications of global imbalance between the rates of marine dinitrogen (N₂) fixation and denitrification have spurred us to understand the former process in the Arabian Sea, which contributes considerably to the global nitrogen budget. Heterotrophic bacteria have gained recent appreciation for their major role in marine N budget by fixing a significant amount of N₂. Accordingly, we hypothesize a probable role of heterotrophic diazotrophs from the ¹⁵N₂ enriched isotope labeling dark incubations that witnessed rates comparable to the light incubations in the eastern Arabian Sea during spring 2010. Maximum areal rates (8 mmol N m⁻² d⁻¹) were the highest ever observed anywhere in world oceans. Our results suggest that the eastern Arabian Sea gains ~92% of its new nitrogen through N₂ fixation. Our results are consistent with the observations made in the same region in preceding year, i.e., during the spring of 2009.

Keywords: dinitrogen fixation, ¹⁵N, ¹³C, nitrogen budget, carbon uptake rate, nutrients, biogeochemistry, Arabian Sea

INTRODUCTION

Reactive nitrogen (e.g., NO_3^-) is an important substrate for marine primary producers because dinitrogen (N₂), though the most abundant gas in the Earth's atmosphere, is unassimilable by most photosynthetic organisms (Middelburg and Nieuwenhuize, 2000). However, marine diazotrophic cyanobacteria (e.g., *Trichodesmium*) have an enzymatic advantage to convert atmospheric N₂ gas to a bioavailable form of nitrogen (such as NH_4^+). These diazotrophs are distributed over the oligotrophic tropical and sub-tropical marine environments, e.g., in the Arabian Sea located in the northwest Indian Ocean (Capone and Carpenter, 1982; Jickells et al., 2017).

The Arabian Sea is a hotspot for studying N_2 fixation. Strong summer monsoonal winds cause intense upwelling over the western Arabian Sea enhancing primary productivity over the Somali coast (Prasannakumar et al., 2001). Further, the northern and central Arabian Seas are known for high productivity during winter, caused by the cooling-driven deep convection (Madhupratap et al., 1996; Singh and Ramesh, 2015). High biological production in the surface layers and its subsequent export leads to oxygen depletion in the subsurface layers (Naqvi and Jayakumar, 2000) that further triggers denitrification process, i.e., the release of N_2 and N_2O back to the atmosphere from nitrate (NO_3^-) reduction. Denitrification in the oxygen minimum zones would deplete only NO_3^- thereby lowering the nitrogen: phosphorus (N:P) ratio in dissolved nutrient pool (Deutsch et al., 2007). An imbalance between N_2 fixation and denitrification rates based on N:P stoichiometry is an imperative problem in the marine nitrogen budget (Codispoti, 2007). N_2 fixation and denitrification dominate ocean N sources and sinks processes, respectively (Codispoti et al., 2001).

OPEN ACCESS

Edited by:

Selvaraj Kandasamy, Xiamen University, China

Reviewed by:

II-Nam Kim, Incheon National University, South Korea Gupta GVM, Centre for Marine Living Resources and Ecology, Ministry of Earth Sciences, India

*Correspondence:

Arvind Singh arvinds@prl.res.in

[†]Present Address:

R. Ramesh, School of Earth and Planetary Sciences, NISER, Jatni, India

Specialty section:

This article was submitted to Marine Biogeochemistry, a section of the journal Frontiers in Marine Science

Received: 22 December 2016 Accepted: 08 March 2017 Published: 23 March 2017

Citation:

Kumar PK, Singh A, Ramesh R and Nallathambi T (2017) N₂ Fixation in the Eastern Arabian Sea: Probable Role of Heterotrophic Diazotrophs. Front. Mar. Sci. 4:80. doi: 10.3389/fmars.2017.00080

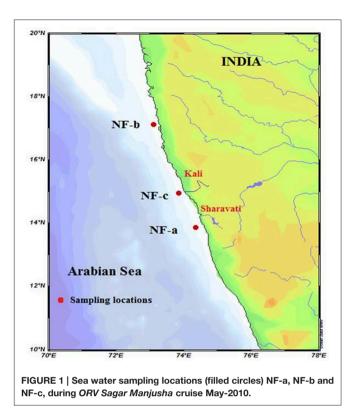
Global nitrogen loss and gain rates are in imbalance, indicating either an overestimation of the loss processes or an underestimation of the gain processes (Codispoti, 2007). The recent recognition of greater diversity (Zehr et al., 2001, 2003) and wider distribution (Hewson et al., 2007; Mulholland, 2007) of marine diazotrophs than had been appreciated hitherto (Mahaffey et al., 2005) suggested an underestimation of N2 fixation. Contribution of atmospheric deposition and riverine nutrients to primary production is minor in the Arabian Sea (Singh and Ramesh, 2011; Singh et al., 2012) which further supports the fact that N₂ fixation is a major process in this region. The Arabian Sea witnesses diverse group of diazotrophs, which may fix N2 at varying rates (Mulholland and Capone, 2009). During spring and autumn seasons, calmer winds, warmer waters, and shallower mixed layers make the Arabian Sea oligotrophic, thus creating a niche for Trichodesmium blooms (Gandhi et al., 2011). The seasonal occurrence of N2 fixation over the Arabian Sea makes it an unique region for studying nitrogen budget (Naqvi, 1987; Capone et al., 1998).

Trichodesmium is not the only species which fixes N₂, as there are some other fixers such as γ - Proteobacteria, and other small heterotrophs also contribute substantially to N₂ fixation rates (Zehr et al., 1995). N₂ fixation might be mediated by a variety of auto and heterotrophic bacterial community in the eastern Arabian Sea, a region of rather limited information. Previously estimated rates (Gandhi et al., 2011) were surprisingly high; so in this study, we revisited the N₂ fixation and carbon uptake rates over the eastern Arabian Sea to verify the veracity of the reported higher rates, using the ¹⁵N₂ gas tracer technique (Montoya et al., 1996). We report measured N₂ fixation rates for dark and light conditions and discuss the possible reasons for this estimated rates.

MATERIALS AND METHODS

Sampling for Incubation Experiments

Water samples were collected using Niskin bottles (bottles were closed by a messenger) from the four different depths (0, 5, 10, and 20 m) within the euphotic zone at three locations (NF-a, NF-b and NF-c that have station depths 42, 37, and 37 m, respectively) over the eastern Arabian Sea, during ORV Sagar Manjusha cruise during 10-14 May 2010 (Figure 1). Duplicate samples were taken from each depth in 1.225 L polycarbonate Nalgene bottles. All the bottles filled without headspace followed by the addition of ¹⁵N₂ gas (bubble method) with the chromatographic gas tight syringe (Montoya et al., 1996). Two milliliters of ¹⁵N₂ gas (99% ¹⁵N enriched gas from Cambridge Isotope Laboratories, Inc. USA) and 1 ml of 0.2 mmol ml⁻¹ NaH¹³CO₃ (99% ¹³C enriched) tracers were injected to each bottle (final enrichment of 16.6% for ¹⁵N and 8.5% for ¹³C) before the start of the incubations, which were performed during 10:00–14:00 h, i.e., symmetric to local noon. Tracer added bottles were covered with the calibrated neutral density filters to simulate the irradiance at the depths from which the samples were taken. After the incubations, the samples were filtered sequentially through pre-combusted (4 h at 400°C) Whatmann GF/F filters (25 mm diameter and 0.7 μ m pore size), washed with filtered sea



water, dried in an oven at 50°C overnight and stored for further mass spectrometric analysis. At each station, 2 L surface seawater was collected for measuring the nitrogen isotopic composition of natural particulate organic nitrogen (PON) and carbon (POC).

Mass Spectrometric Analysis and Calculation of Rates

Elemental analyzer interface with continuous flow mass spectrometer at the Physical Research Laboratory, Ahmedabad was used to measure the PON, POC, atom % 15 N and atom % 13 C in the samples. Volumetric rate of N₂ fixation were calculated following (Montoya et al., 1996).

N uptake rate =
$$(1/t)[(A_{PNf} - AN_0)/(A_{Nenrich} - A_{PN0})]$$

 $\times [PON]_f$ (1)

Where, $A_{PN0} = {}^{15}N$ atom% in PON at the start of experiment, $A_{PNf} = {}^{15}N$ atom% in PON at the end of experiment, t = time of incubation (4 hrs), $[PON]_f =$ concentration of PON at the end of the experiment and $A_{Nenrich} = {}^{15}N$ enrichment in the dissolved form after tracer addition at the start of the incubation, which is estimated as:

$$A_{\text{Nenrich}} = ({}^{15}\text{N}_{\text{tracer}} \times \text{tracer conc} + {}^{15}\text{N}_{\text{natural}} \times \text{ambient}$$

$$\text{conc})/(\text{tracer conc} + \text{natural conc}) \qquad (2)$$

Surface water samples (at 0–1 m depth) were also incubated in complete dark condition and these estimates are attributed to the presence of heterotrophic diazotrophs and their contribution to

N₂ fixation. It has been discovered that in the "bubble method," only 40% enrichment can be achieved in 4 h incubations, which further results in 40% underestimation in the rates (Mohr et al., 2010). Hence, we multiplied N₂ fixation rates by a constant factor of 2.5 to avoid possible underestimation in the bubble gas technique. Later, it was discovered that the underestimation of rates is community dependent-there is less underestimation for Trichodesmium compared to that for other diazotrophs. Our sampling area witnesses Trichodesmium so underestimation in bubble method was less (GroSSkopf et al., 2012; White, 2012; Klawonn et al., 2015). We still multiplied our rates by 2.5 because we compared our results with Gandhi et al. (2011), who multiplied by the same factor. Carbon uptake rate is calculated by substituting N and ¹⁵N by C and ¹³C, respectively, in Equations (1) and (2) (Slawyk et al., 1977). Areal rates were calculated from the volumetric rates using the trapezoidal rule of integration, i.e., mean of volumetric rates were multiplied by the corresponding depth interval and then summed for all the depth intervals. Hundred milliliters of each sample was separately collected for nutrient measurements using a SKALAR auto analyzer at the offshore laboratory.

Spatial distribution of sea surface temperature (SST) was plotted using the Gridded High Resolution Sea Surface Temperature: Operational sea surface temperature and sea ice analysis (GHRSST: OSTIA) satellite data (Donlon et al., 2012). The mixed layer depth (MLD) was estimated based on the temperature criterion (0.2°C difference from the SST; de Boyer Montégut et al., 2004). Salinity values were obtained from the CTD data, values ranged from 35.23 to 35.56 at the sampling locations (**Figure 2**).

RESULTS AND DISCUSSION

Hydrography

Temperature profiles from a portable CTD at sampling locations NF-a, NF-b, and NF-c showed that MLD varied between 17 and 20 m (**Figure 2**). Surface temperature at NF-b was \sim 1.4°C less than that at the other two stations. Temperature measurements obtained from CTD at surface level were reproduced by the

satellite remote sensing data images (**Figure 3**). Salinity values obtained from CTD varied from 35.23 to 35.56 at the surface with maximum at NF-b and minimum at NF-c. NF-b is possibly influenced by Western India Coastal Current (WICC), which brings convecting mixing driven colder water from the north to the south during pre-monsoon (Schott and McCreary, 2001) as evidenced in the temperature profile (**Figure 1**).

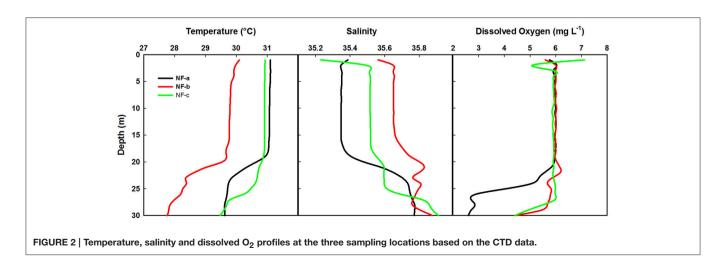
Dissolved oxygen from the same CTD casts showed a dip of 2 mg L⁻¹ at 3 m depth at sampling location NF-c, whereas there was no vertical gradient in oxygen in the upper 20 m at locations NF-a and NF-b (**Figure 2**). Oxygen showed a sudden decline at all the stations below 20 m depth. Lower temperature at NF-b was associated with the detectable SiO₄ values, whereas the dip in the dissolved oxygen was associated with the maximum NO_x (NO₂⁻ + NO₃⁻), phosphate (PO₄³⁻) values and higher N:P ratios at NF-c (**Table 1**).

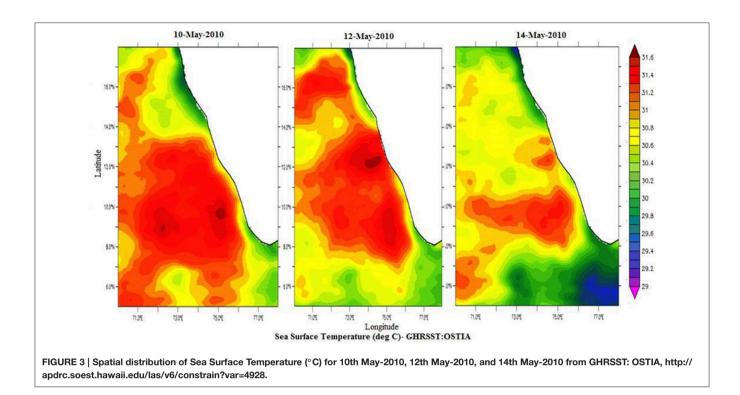
Nutrients

Nutrient concentrations were higher at the surface and decreased with depth, with maximum surface value at NF-c (**Figure 4**). NO₃⁻ varied from 0.37 to 2.42 μ M while PO₄³⁻ ranged from 0 to 0.71 μ M (**Table 1**). At sampling station NF-b, SiO₄ values were readily detectable with maximum at surface (1.69 μ M) and decreased with depth (**Table 1**). We calculated P* that indicates PO₄³⁻ concentration deviations from the Redfield ratio (N:P = 16):

$$P^* = [PO_4^{3-}] - [NO_3^{-}]/16$$
(3)

Positive P* indicates PO_4^{3-} concentrations in excess compared to NO_3^- (Deutsch et al., 2007). Surface water P*-values were mostly in excess (except at 20 m at NF-c) and varied from -0.06to 0.56 μ M (**Table 1**). P*-values at surface showed increasing trend from NF-a to NF-c, and decreased with depth at all the three locations. PO_4^{3-} and NO_3^- maxima at NF-c could be due to the upwelling of subsurface water. Low P* at NF-a could also be either due to the consumption of phosphate by diazotrophy or due to the upwelling, which starts during May (Gupta et al., 2016) as evidenced in the temperature profile at NF-a (**Figure 2**).





Upwelling would create low oxygen just below MLD (Gupta et al., 2016; Sudheesh et al., 2016), as observed at NF-a (**Figure 2**). However, high P*-values would have been expected in low oxygen water but deeper water, as evidenced from the P* profiles (**Table 1**), has low P*—possibly because of the remineralization of diazotrophy dominated organic matter (high N:P ratio) in the deeper depths. PO_4^{3-} was below the detection limit at 20 m depth at NF-c yet there was N₂ fixation. This could again be attributed to the PO_4^{3-} consumption by diazotrophs.

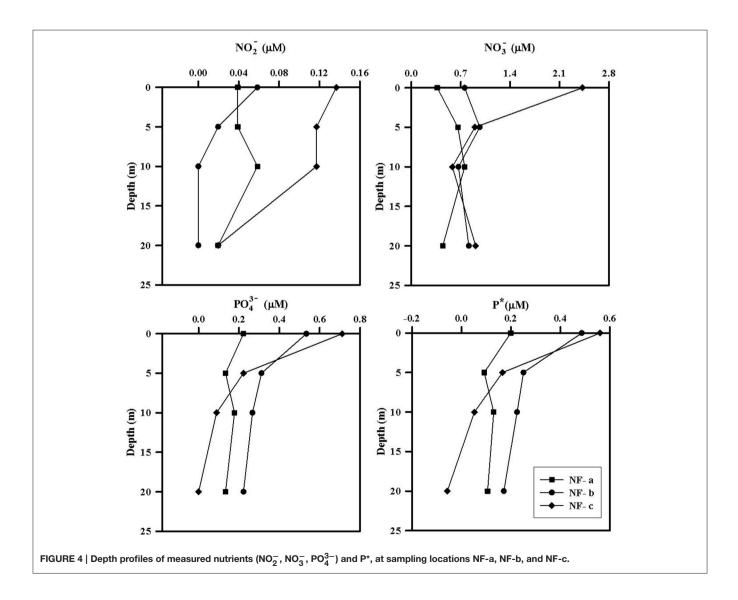
N:P ratio was less than the Redfield ratio (positive P*, Table 1) at all stations with maximum at NF- c, which could have resulted in nitrogen limitation and further would have facilitated of N2 fixation. N2 fixation is controlled by iron and PO_4^{3-} availability (Wu et al., 2000; Capone, 2001; Sañudo-Wilhelmy et al., 2001; Mulholland, 2007). At these sampling locations, N₂ fixation may be regulated by iron input as PO_4^{3-} is not the limiting nutrient in the northern Indian Ocean as suggested by P*-values (Shiozaki et al., 2014). Iron limitation is likely, as some part of the western Arabian Sea has been highlighted as a high nutrient low chlorophyll (HNLC) region (Naqvi et al., 2010; Moffett et al., 2015). Changes in the response of external forcing (e.g., seasonal monsoon and upwelling events) and inputs (e.g., aeolian and river influx) have specific control on N₂ fixation over the basin (Mulholland and Capone, 2009). POC and PON varied between 15 and 73 µM and 3-51 µM, respectively (Table 1), but did not show any correlation with C and N₂ fixation rates. However, the highest values of POC, PON, C, and N₂ fixation rates were observed at the surface at NF-a.

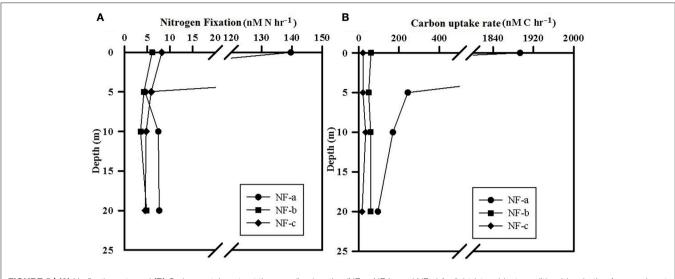
N₂ Fixation Rate and Carbon Uptake

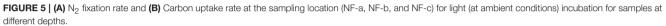
N₂ fixation rates varied from 6.08 to 139.62 nM N h⁻¹, while carbon uptake rate ranged between 14 and 1,894 nM C h^{-1} (Figure 5). Our N₂ fixation rates were higher than those synthesized by Capone et al. (2008) in the world oceans (0-5.4 nM N h⁻¹). The highest values for both carbon uptake and N₂ fixation rate for light incubation fixation (Figure 5) were observed at NF-a lying south of the other two sampling locations (Figure 1). N₂ fixation rates were higher in dark incubations compared to the light incubations at the surface level, except for NF-a (44.93, 63.32, and 164.7 nM N h^{-1} at NF-a, NF-b, and NF-c, respectively, Figure 6). Whereas carbon uptake rates in the light incubations were an order of magnitude higher than in the dark incubations (Figure 6). Higher N₂ fixation rates for the dark incubations compared to light might be attributable to the presence of heterotrophic species at the surface. Heterotrophs contributed up to 64% to the total N₂ fixation (estimated from light and dark incubations, assuming light incubations correspond to phototrophic and dark to heterotrophic). Heterotrophic contribution to N₂ fixation could be even higher in the deeper waters due to favorable conditions for them. These higher values associated with dark incubation suggest that heterotrophic N2 fixers might play an important role in fixing N2 over the eastern Arabian Sea. In the surface waters of the Arabian Sea, the DNA and RNA recovered during the southwest monsoon periods were also categorized as those of heterotrophic bacteria (Jayakumar et al., 2012; Bird and Wyman, 2013). Therefore, our results suggest that an active N2 fixation by heterotrophic bacteria could occur

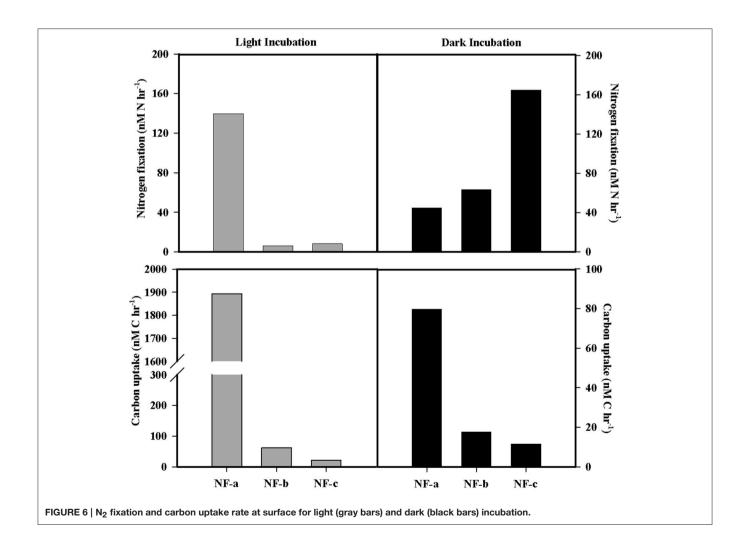
TABLE 1 | Sampling Date, Latitude (°N), Longitude (°E), water depth (m), Nutrients concentrations (μ M), N:P, P^{*} (μ M), particulate organic carbon (POC, μ M) and nitrogen (PON, μ M), N₂ fixation (nM N h⁻¹), and carbon (C) uptake (nM C h⁻¹) at the three locations sampled in the eastern Arabian Sea.

| Date and Station | Lat | Long | Depth | NO ₂ | NO_3 | PO ₄ ³⁻ | SiO ₄ | N:P | P * | POC | PON | N ₂ fixation | C uptake |
|------------------|-------|-------|-------|-----------------|--------|-------------------------------|------------------|------|------------|-----|-----|-------------------------|----------|
| 10-May-10; NF-a | 13.87 | 74.36 | 0 | 0.04 | 0.37 | 0.22 | 0.00 | 1.66 | 0.20 | 73 | 51 | 139.62 | 1894 |
| | | | 5 | 0.04 | 0.66 | 0.13 | 0.00 | 4.94 | 0.10 | 26 | 5 | 4.54 | 244 |
| | | | 10 | 0.06 | 0.76 | 0.18 | 0.00 | 4.25 | 0.13 | 21 | 7 | 7.44 | 170 |
| | | | 20 | 0.02 | 0.45 | 0.13 | 0.19 | 3.34 | 0.10 | 23 | 5 | 7.65 | 96 |
| 12-May-10; NF-b | 17.12 | 73.11 | 0 | 0.06 | 0.76 | 0.53 | 1.69 | 1.42 | 0.49 | 24 | 6 | 6.08 | 61 |
| | | | 5 | 0.02 | 0.97 | 0.31 | 1.44 | 3.11 | 0.25 | 23 | 5 | 4.26 | 50 |
| | | | 10 | 0.00 | 0.67 | 0.27 | 0.63 | 2.51 | 0.22 | 21 | 4 | 3.53 | 60 |
| | | | 20 | 0.00 | 0.81 | 0.22 | 0.94 | 3.66 | 0.17 | 18 | 4 | 4.85 | 60 |
| 14-May-10; NF-c | 14.96 | 73.84 | 0 | 0.14 | 2.42 | 0.71 | 0.00 | 3.40 | 0.56 | 44 | 25 | 8.21 | 22 |
| | | | 5 | 0.12 | 0.90 | 0.22 | 0.00 | 4.05 | 0.17 | 20 | 4 | 5.86 | 22 |
| | | | 10 | 0.12 | 0.58 | 0.09 | 0.00 | 6.53 | 0.05 | 16 | 4 | 4.76 | 34 |
| | | | 20 | 0.02 | 0.91 | 0.00 | 0.00 | NA | -0.06 | 15 | 3 | 4.54 | 18 |









in the surface water of the eastern Arabian Sea (Shiozaki et al., 2014).

Euphotic-depth integrated autotrophic N₂ fixation and carbon uptake rates were highest at NF-a, where N₂ fixation rate was $8.4 \pm 2.8 \text{ mmol N m}^{-2} \text{ d}^{-1}$ (standard deviation of duplicate samples) and carbon uptake rate was $74.7 \pm 17.7 \text{ mmol C m}^{-2} \text{ d}^{-1}$. Our areal rates (8 mmol N m⁻² d⁻¹) were the highest ever observed in anywhere in world oceans (**Table 2**) and comparable to those measured by Gandhi et al. (2011) in the similar region-suggesting that the high rates during the spring 2009 were not

episodic rather it could be a regular phenomenon during the spring in the Arabian Sea. High rates in the eastern Arabian Sea are probably due to the availability of both the essential nutrients for diazotrophs, i.e., iron and PO_4^{3-} . Iron is available because this region unlike the western Arabian Sea is close to the *Thar* desert, whereas the western Arabian Sea is recognized to be an HNLC region (Naqvi et al., 2010). PO_4^{3-} is available because upwelled water brings PO_4^{3-} rich (compared to NO_3^-) water as NO_3^- is lost in denitrification. So the Arabian Sea is unique basin for sustaining diazotrophy and the higher rates of N_2 fixation. On

TABLE 2 | Summary of Photic N₂ fixation rates in the world oceans (Table updated after Singh et al., 2013; Benavides and Voss, 2015).

| Methodology | Areal Rates (μ mol N m ⁻² d ⁻¹) | Region | References | |
|---|---|---|----------------------------|--|
| Acetylene Reduction Assay | 239 | Western tropical North Atlantic | Capone et al., 2005 | |
| ¹⁵ N ₂ tracer-bubble method | 850 | Western tropical North Atlantic | Capone et al., 2005 | |
| ¹⁵ N ₂ tracer-bubble method | 2.4–532 | Western North Atlantic | Mulholland et al., 2012 | |
| Extrapolation | 714–3,571 | North Atlantic Ocean | Carpenter and Romans, 1991 | |
| Extrapolation | 160–430 | North Atlantic Ocean | Lipschultz and Owens, 1996 | |
| $N^* = [NO_3^-] - 16 \times [PO_4^{3-}] + 2.72$ | 500-2,500 | North Atlantic Ocean | Michaels et al., 1996 | |
| $N^* = ([NO_3^-] - 16 \times [PO_4^3^-] + 2.90) \times 0.87$ | 197 | North Atlantic Ocean | Gruber and Sarmiento, 1997 | |
| $DIN_{XS} = [NO_3^-] - 16 \times [PO_4^{3-}]$ | 70–208 | North Atlantic Ocean | Hansell et al., 2004 | |
| $P^* = [PO_4^{3-}] - [NO_3^{-}]/16$ | 63 | North Atlantic Ocean | Deutsch et al., 2007 | |
| $P^* = [PO_4^{3-}] - [NO_3^{-}]/16$ | 151–178 | North Atlantic Ocean | Palter et al., 2011 | |
| ¹⁵ N ₂ tracer-bubble method | 1.2–298 | North Atlantic Ocean | FernÁndez et al., 2010 | |
| ¹⁵ N ₂ tracer-bubble method | 1.8–182 | North Atlantic Ocean | Moore et al., 2009 | |
| ¹⁵ N ₂ tracer-dissolution method | 91 ± 4 | Atlantic Ocean | GroSSkopf et al., 2012 | |
| ¹⁵ N-nitrate and ammonium | 4.5-68.1 | Tropical Northwest Atlantic | Goering et al., 1966 | |
| ¹⁵ N ₂ - bubble and Acetylene Reduction Assay | 73–90 | Tropical Northwest Atlantic | Falcón et al., 2004 | |
| ¹⁵ N ₂ tracer-bubble method | 41–93 | Sargasso Sea | Orcutt et al., 2001 | |
| $DIN_{xs} = [NO_3^-] - 16 \times [PO_4^{3-}]$ | 45–259 | Sargasso Sea | Bates and Hansell, 2004 | |
| $N^* = [NO_3^-] - 14.63 \times [PO_4^{3-}]$ | 120 ± 9 | Sargasso Sea | Singh et al., 2013 | |
| Acetylene Reduction Assay | 0.05–540 | Subtropical Northeast Atlantic | Carpenter and Price, 1977 | |
| Acetylene Reduction Assay | 0.001-0.09 | Subtropical Northeast Atlantic | Benavides et al., 2011 | |
| ¹⁵ N ₂ tracer-bubble method | 28–142 | Tropical Northeast Atlantic | Turk et al., 2011 | |
| ¹⁵ N ₂ tracer-bubble method | 56–60 | Tropical Northeast Atlantic | Turk-Kubo et al., 2012 | |
| ¹⁵ N ₂ tracer-bubble method | 15–424 | Eastern equatorial Atlantic | Subramaniam et al., 2013 | |
| ¹⁵ N ₂ tracer-bubble method | 0–148 | Eastern tropical south Pacific tropical South Pacific | Dekaezemacker et al., 2013 | |
| ¹⁵ N ₂ tracer-bubble method | up to 800 | Off Peru-South Pacific | Loescher et al., 2014 | |
| ¹⁵ N ₂ tracer-bubble method | 0–23 | Eastern tropical South Pacific | Knapp et al., 2016 | |
| ¹⁵ N ₂ tracer-bubble method | 520 ± 160 | Eastern North Pacific gyre | Montoya et al., 2004 | |
| ¹⁵ N ₂ tracer-bubble method | 126 ± 47 | Timor–Arafura–Coral seas | Montoya et al., 2004 | |
| ¹⁵ N ₂ tracer-bubble method | 3,995 | Arafura Sea | Montoya et al., 2004 | |
| Acetylene Reduction Assay | 85 | Subtropical North Pacific Ocean | Karl et al., 1997 | |
| ¹⁵ N ₂ tracer-bubble method | 1–13 | South China Sea | Chen et al., 2008 | |
| ¹⁵ N ₂ tracer-bubble method | 0–90 | North Pacific Ocean | Shiozaki et al., 2010 | |
| $^{15}N_2$ tracer-bubble method | 20–310 | North Pacific Ocean | Church et al., 2009 | |
| Acetylene Reduction Assay | 170 | Central Arabian Sea | Capone et al., 1998 | |
| $^{15}N_2$ tracer-bubble method | 24.6-47.1 | Arabian Sea | Shiozaki et al., 2014 | |
| ¹⁵ N ₂ tracer-bubble method | 6.27-16.6 | Equatorial and southern Indian Ocean | Shiozaki et al., 2014 | |
| ¹⁵ N ₂ tracer-bubble method* | 100-34,000 | Eastern Arabian Sea | Gandhi et al., 2011 | |
| ¹⁵ N ₂ tracer-bubble method | 174–238 | Southeastern Arabian Sea | Bhavya et al., 2016 | |
| ¹⁵ N ₂ tracer-bubble method* | 1,047-5,589 | Eastern Arabian Sea | This study | |

*These rates were corrected for bubble in-equilibrium.

the other hand, the other major oceans, like the Atlantic does not have enough phosphate (Wu et al., 2000), while the Pacific does not have enough iron (Behrenfeld and Kolber, 1999).

Gandhi et al. (2011) showed an overall increasing trend in the N_2 fixation rate from south to north, whereas in this present study, it was in the opposite direction. N_2 fixation supplies a large portion of the new N that supports marine productivity (Capone, 1997; Karl et al., 2002; LaRoche and Breitbarth, 2005; Mahaffey et al., 2005). Estimated contribution of new nitrogen to N_2 fixation in the present study is about 92% of new nitrogen, which is the same as that reported by the previous study (Gandhi et al., 2011).

Most observations around the world oceans are indirectly derived from geochemical estimates that are based on sub-surface nutrient distribution (Table 2). We have directly measured N2 fixation using ¹⁵N₂ tracer, which is still the best available method despite its inherent problems of incomplete dissolution of the bubble (GroSSkopf et al., 2012). Our findings of high N₂ fixation in this region have large implications in understanding marine C, N, and cycles. If we extrapolate these high rates over the Arabian Sea, then we might find the missing nitrogen inputs (Codispoti, 2007). But we require to conduct more N₂ fixation experiments using the dissolution method in this region for extrapolation. In addition to seasonal upwelling, N2 fixation might play a role in making the Arabian Sea perennially productive (Singh and Ramesh, 2015). Higher N₂ fixation and its degradation in the subsurface waters may also be useful in understating the phosphorous cycle-low P* at the deeper waters and higher P* at the surface-contrary to what is expected from the denitrified waters.

REFERENCES

- Bates, N. R., and Hansell, D. A. (2004). Temporal variability of excess nitrate in the subtropical mode water of the North Atlantic Ocean. *Mar. Chem.* 84, 225–241. doi: 10.1016/j.marchem.2003.08.003
- Behrenfeld, M. J., and Kolber, Z. S. (1999). Widespread iron limitation of phytoplankton in the South Pacific Ocean. *Science* 283, 840–843. doi: 10.1126/science.283.5403.840
- Benavides, M., Agawin, N. S. R., Arístegui, J., Ferriol, P., and Stal, L. J. (2011). Nitrogen fixation by Trichodesmium and small diazotrophs in the subtropical northeast Atlantic. *Aquat. Microb. Ecol.* 65, 45–53. doi: 10.3354/ame01534
- Benavides, M., and Voss, M. (2015). Five decades of N2 fixation research in the North Atlantic Ocean. Front. Mar. Sci. 2:40. doi: 10.3389/fmars.2015.00040
- Bhavya, P., Kumar, S., Gupta, G., Sudheesh, V., Sudharma, K., Varrier, D., et al. (2016). Nitrogen uptake dynamics in a tropical eutrophic estuary (Cochin, India) and adjacent coastal waters. *Estuar. Coasts* 39, 54–67. doi: 10.1007/s12237-015-9982-y
- Bird, C., and Wyman, M. (2013). Transcriptionally active heterotrophic diazotrophs are widespread in the upper water column of the Arabian Sea. *FEMS Microbiol. Ecol.* 84, 189–200. doi: 10.1111/1574-6941.12049
- Capone, D. G. (1997). Trichodesmium, a globally significant marine Cyanobacterium. *Science* 276, 1221–1229. doi: 10.1126/science.276.5316.1221
- Capone, D. G. (2001). Marine nitrogen fixation: what's the fuss? Curr. Opin. Microbiol. 4, 341–348. doi: 10.1016/S1369-5274(00)00215-0
- Capone, D. G., Bronk, D. A., Mulholland, M. R., and Carpenter, E. J. (2008). *Nitrogen in the Marine Environment*. Burlington, MA: Academic Press.
- Capone, D. G., Burns, J. A., Montoya, J. P., Subramaniam, A., Mahaffey, C., Gunderson, T., et al. (2005). Nitrogen fixation by *Trichodesmium* spp.: an important source of new nitrogen to the tropical and subtropical North Atlantic Ocean. *Glob. Biogeochem. Cycles* 19, 1–17. doi: 10.1029/2004GB002331

CONCLUSION

Results from the three sampling locations in the eastern Arabian Sea suggest that the Arabian Sea witnessed the highest ever rates of the N₂ fixation among the world ocean for the two consecutive springs. Out of all nitrogen gain processes, about 92% of new nitrogen is gained through N₂ fixation only, with highest areal rates (8 mmol N m⁻² d⁻¹). N₂ fixation rate for the dark was higher than the light incubation at the surface except for NF-a, which alluded to the presence of heterotrophic species. Based on the higher N₂ fixation values at the surface for dark incubation, we hypothesize that heterotrophic fixers dominantly (about 64% of total N₂ fixation is by heterotrophs) play an important role in fixing N₂. There is a consistency between the higher N₂ fixation rates and the carbon uptake rate.

AUTHOR CONTRIBUTIONS

AS and RR designed research; AS, RR, and NT performed research; KK and AS analyzed data; and KK and AS wrote the manuscript with major inputs from all the co-authors.

ACKNOWLEDGMENTS

We thank the Indian Space Research Organisation Geosphere-Biosphere program (ISRO-IGBP) for funding, and all the participants and the crew members of ORV *Sagar Manjusha* for their assistance onboard. We thank Suhas Shetye for nutrients measurements.

- Capone, D. G., and Carpenter, E. J. (1982). Nitrogen fixation in the marine environment. *Science* 217, 1140–1142. doi: 10.1126/science.217.4565.1140
- Capone, D. G., Subramaniam, A., Montoya, J. P., Voss, M., Humborg, C., Johansen, A. M., et al. (1998). An extensive bloom of the N2-fixing cyanobacterium *Trichodesmium erythraeum* in the central Arabian Sea. *Mar. Ecol. Prog. Ser.* 172, 281–292. doi: 10.3354/meps172281
- Carpenter, E. J., and Price, C. C. (1977). Nitrogen fixation, distribution, and production of Oscillatoria (*Trichodesmium*) spp. in the western Sargasso and Caribbean Seas. *Limnol. Oceanogr.* 22, 60–72. doi: 10.4319/lo.1977.22. 1.0060
- Carpenter, E. J., and Romans, K. (1991). Major role of the cyanobacterium Trichodesmium in nutrient cycling in the North Atlantic Ocean. *Science* 254, 1356–1358. doi: 10.1126/science.254.5036.1356
- Chen, Y. L., Chen, H.-Y., Tuo, S., and Ohki, K. (2008). Seasonal dynamics of new production from Trichodesmium N2 fixation and nitrate uptake in the upstream Kuroshio and South China Sea basin. *Limnol. Oceanogr.* 53, 1705–1721. doi: 10.4319/lo.2008.53.5.1705
- Church, M. J., Mahaffey, C., Letelier, R. M., Lukas, R., Zehr, J. P., and Karl, D. M. (2009). Physical forcing of nitrogen fixation and diazotroph community structure in the North Pacific subtropical gyre. *Glob. Biogeochem. Cycles* 23:GB2020. doi: 10.1029/2008gb003418
- Codispoti, L. A. (2007). An oceanic fixed nitrogen sink exceeding 400 TgN a-1 vs the concept of homeostasis in the fixed-nitrogen inventory. *Biogeosciences* 4, 233–253. doi: 10.5194/bg-4-233-2007
- Codispoti, L., Brandes, J. A., Christensen, J., Devol, A., Naqvi, S., Paerl, H. W., et al. (2001). The oceanic fixed nitrogen and nitrous oxide budgets: moving targets as we enter the anthropocene? *Sci. Mar.* 65, 85–105. doi: 10.3989/scimar.2001.65s285
- de Boyer Montégut, C., Madec, G., Fischer, A. S., Lazar, A., and Iudicone, D. (2004). Mixed layer depth over the global ocean: an examination of profile

data and a profile-based climatology. J. Geophys. Res. Oceans 109:C12003. doi: 10.1029/2004jc002378

- Dekaezemacker, J., Bonnet, S., Grosso, O., Moutin, T., Bressac, M., and Capone, D. (2013). Evidence of active dinitrogen fixation in surface waters of the eastern tropical South Pacific during El Ni-o and La Ni-a events and evaluation of its potential nutrient controls. *Glob. Biogeochem. Cycles* 27, 768–779. doi: 10.1002/gbc.20063
- Deutsch, C., Sarmiento, J. L., Sigman, D. M., Gruber, N., and Dunne, J. P. (2007). Spatial coupling of nitrogen inputs and losses in the ocean. *Nature* 445, 163–167. doi: 10.1038/nature05392
- Donlon, C. J., Martin, M., Stark, J., Roberts-Jones, J., Fiedler, E., and Wimmer, W. (2012). The operational sea surface temperature and sea ice analysis (OSTIA) system. *Remote Sens. Environ.* 116, 140–158. doi: 10.1016/j.rse.2010.10.017
- Falcón, L. I., Carpenter, E. J., Cipriano, F., Bergman, B., and Capone, D. G. (2004). N2 fixation by unicellular bacterioplankton from the Atlantic and Pacific Oceans: phylogeny and *in situ* rates. *Appl. Environ. Microbiol.* 70, 765–770. doi: 10.1128/AEM.70.2.765-770.2004
- Fernández, A., Mouriño-Carballido, B., Bode, A., Varela, M., and Marañón, E. (2010). Latitudinal distribution of *Trichodesmium* spp. and N 2 fixation in the Atlantic Ocean. *Biogeosciences* 7, 3167–3176. doi: 10.5194/bg-7-3167-2010
- Gandhi, N., Singh, A., Prakash, S., Ramesh, R., Raman, M., Sheshshayee, M., et al. (2011). First direct measurements of N2 fixation during a Trichodesmium bloom in the eastern Arabian Sea. *Glob. Biogeochem. Cycles* 25:GB4014. doi: 10.1029/2010GB003970
- Goering, J., Dugdale, R., and Menzel, D. W. (1966). Estimates of *in situ* rates of nitrogen uptake by *Trichodesmium* sp. in the tropical Atlantic Ocean. *Limnol. Oceanogr.* 11, 614–620. doi: 10.4319/lo.1966.11.4.0614
- GroSSkopf, T., Mohr, W., Baustian, T., Schunck, H., Gill, D., Kuypers, M. M., et al. (2012). Doubling of marine dinitrogen-fixation rates based on direct measurements. *Nature* 488, 361–364. doi: 10.1038/nature11338
- Gruber, N., and Sarmiento, J. L. (1997). Global patterns of marine nitrogen fixation and denitrification. *Glob. Biogeochem. Cycles* 11, 235–266. doi: 10.1029/97GB00077
- Gupta, G., Sudheesh, V., Sudharma, K., Saravanane, N., Dhanya, V., Dhanya, K., et al. (2016). Evolution to decay of upwelling and associated biogeochemistry over the southeastern Arabian Sea shelf. J. Geophys. Res. Biogeosciences 121, 159–175. doi: 10.1002/2015JG003163
- Hansell, D. A., Bates, N. R., and Olson, D. B. (2004). Excess nitrate and nitrogen fixation in the North Atlantic Ocean. *Mar. Chem.* 84, 243–265. doi: 10.1016/j.marchem.2003.08.004
- Hewson, I., Moisander, P. H., Achilles, K. M., Carlson, C. A., Jenkins, B. D., Mondragon, E. A., et al. (2007). Characteristics of diazotrophs in surface to abyssopelagic waters of the Sargasso Sea. *Aquat. Microb. Ecol.* 46, 15–30. doi: 10.3354/ame046015
- Jayakumar, A., Al-Rshaidat, M. M., Ward, B. B., and Mulholland, M. R. (2012). Diversity, distribution, and expression of diazotroph nifH genes in oxygendeficient waters of the Arabian Sea. *FEMS Microbiol. Ecol.* 82, 597–606. doi: 10.1111/j.1574-6941.2012.01430.x
- Jickells, T., Buitenhuis, E., Altieri, K., Baker, A., Capone, D., Duce, R., et al. (2017). A re-evaluation of the magnitude and impacts of anthropogenic atmospheric nitrogen inputs on the ocean. *Glob. Biogeochem. Cycles* 31, 289–305. doi: 10.1002/2016GB005586
- Karl, D., Letelier, R., Tupas, L., Dore, J., Christian, J., and Hebel, D. (1997). The role of nitrogen fixation in biogeochemical cycling in the subtropical North Pacific Ocean. *Nature* 388, 533–538. doi: 10.1038/41474
- Karl, D., Michaels, A., Bergman, B., Capone, D. G., Carpenter, E. J., Letelier, R., et al. (2002). Dinitrogen fixation in the world's oceans. *Biogeochemistry* 57–58, 47–98. doi: 10.1023/A:1015798105851
- Klawonn, I., Lavik, G., Böning, P., Marchant, H. K., Dekaezemacker, J., Mohr, W., et al. (2015). Simple approach for the preparation of 15-15N2enriched water for nitrogen fixation assessments: evaluation, application and recommendations. *Front. Microbiol.* 6:769. doi: 10.3389/fmicb.2015.00769
- Knapp, A. N., Casciotti, K. L., Berelson, W. M., Prokopenko, M. G., and Capone, D. G. (2016). Low rates of nitrogen fixation in eastern tropical South Pacific surface waters. *Proc. Natl. Acad. Sci. U.S.A.* 113, 4398–4403. doi: 10.1073/pnas.1515641113
- LaRoche, J., and Breitbarth, E. (2005). Importance of the diazotrophs as a source of new nitrogen in the ocean. J. Sea Res. 53, 67–91. doi: 10.1016/j.seares.2004.05.005

- Lipschultz, F., and Owens, N. J. (1996). An assessment of nitrogen fixation as a source of nitrogen to the North Atlantic Ocean. *Biogeochemistry* 35, 261–274. doi: 10.1007/BF02179830
- Loescher, C. R., GroSSkopf, T., Desai, F. D., Gill, D., Schunck, H., Croot, P. L., et al. (2014). Facets of diazotrophy in the oxygen minimum zone waters off Peru. *ISME J.* 8, 2180–2192. doi: 10.1038/ismej.2014.71
- Madhupratap, M., Kumar, S. P., Bhattathiri, P., Kumar, M. D., Raghukumar, S., Nair, K., et al. (1996). Mechanism of the biological response to winter cooling in the northeastern Arabian Sea. *Nature* 384, 549–552. doi: 10.1038/384549a0
- Mahaffey, C., Michaels, A. F., and Capone, D. G. (2005). The conundrum of marine N2 fixation. *Am. J. Sci.* 305, 546–595. doi: 10.2475/ajs.305.6-8.546
- Michaels, A. F., Olson, D., Sarmiento, J., Ammerman, J. W., Fanning, K. A., Jahnke, R. A., et al. (1996). Inputs, losses and transformations of nitrogen and phosporus in the pelagic North Atlantic Ocean. *Biogeochemistry* 35, 181–226. doi: 10.1007/BF02179827
- Middelburg, J. J., and Nieuwenhuize, J. (2000). Nitrogen uptake by heterotrophic bacteria and phytoplankton in the nitrate-rich Thames estuary. *Mar. Ecol. Prog.* Ser. 203, 13–21. doi: 10.3354/meps203013
- Moffett, J. W., Vedamati, J., Goepfert, T. J., Pratihary, A., Gauns, M., and Naqvi, S. (2015). Biogeochemistry of iron in the Arabian Sea. *Limnol. Oceanogr.* 60, 1671–1688. doi: 10.1002/lno.10132
- Mohr, W., Grosskopf, T., Wallace, D. W., and LaRoche, J. (2010). Methodological underestimation of oceanic nitrogen fixation rates. *PLoS ONE* 5:e12583. doi: 10.1371/journal.pone.0012583
- Montoya, J. P., Holl, C. M., Zehr, J. P., Hansen, A., Villareal, T. A., and Capone, D. G. (2004). High rates of N2 fixation by unicellular diazotrophs in the oligotrophic Pacific Ocean. *Nature* 430, 1027–1032. doi: 10.1038/nature02824
- Montoya, J. P., Voss, M., Kahler, P., and Capone, D. G. (1996). A simple, highprecision, high-sensitivity tracer assay for N2 fixation. *Appl. Environ. Microbiol.* 62, 986–993.
- Moore, C. M., Mills, M. M., Achterberg, E. P., Geider, R. J., LaRoche, J., Lucas, M. I., et al. (2009). Large-scale distribution of Atlantic nitrogen fixation controlled by iron availability. *Nat. Geosci.* 2, 867–871. doi: 10.1038/ ngeo667
- Mulholland, M., Bernhardt, P., Blanco-Garcia, J., Mannino, A., Hyde, K., Mondragon, E., et al. (2012). Rates of dinitrogen fixation and the abundance of diazotrophs in North American coastal waters between Cape Hatteras and Georges Bank. *Limnol. Oceanogr.* 57, 1067–1083. doi: 10.4319/lo.2012.57.4.1067
- Mulholland, M. R. (2007). The fate of nitrogen fixed by diazotrophs in the ocean. Biogeosciences 4, 37–51. doi: 10.5194/bg-4-37-2007
- Mulholland, M. R., and Capone, D. G. (2009). Dinitrogen fixation in the Indian Ocean. Indian Ocean Biogeochem. Process. Ecol. Var. 185, 167–186. doi: 10.1029/2009GM000850
- Naqvi, S. (1987). Some aspects of the oxygen-deficient conditions and denitrification in the Arabian Sea. J. Mar. Res. 45, 1049–1072. doi: 10.1357/002224087788327118
- Naqvi, S. W. A., and Jayakumar, D. A. (2000). Ocean biogeochemistry and atmospheric composition: significance of the Arabian sea. *Curr. Sci.* 78, 289–299.
- Naqvi, S., Moffett, J. W., Gauns, M., Narvekar, P., Pratihary, A., Naik, H., et al. (2010). The Arabian Sea as a high-nutrient, low-chlorophyll region during the late Southwest Monsoon. *Biogeosciences* 7, 2091–2100. doi: 10.5194/bg-7-2091-2010
- Orcutt, K. M., Lipschultz, F., Gundersen, K., Arimoto, R., Michaels, A. F., Knap, A. H., et al. (2001). A seasonal study of the significance of N 2 fixation by *Trichodesmium* spp. at the Bermuda Atlantic Time-series Study (BATS) site. *Deep Sea Res. Part II Top. Stud. Oceanogr.* 48, 1583–1608. doi: 10.1016/S0967-0645(00)00157-0
- Palter, J. B., Lozier, M. S., Sarmiento, J. L., and Williams, R. G. (2011). The supply of excess phosphate across the Gulf Stream and the maintenance of subtropical nitrogen fixation. *Glob. Biogeochem. Cycles* 25:GB4007. doi: 10.1029/2010gb003955
- PrasannaKumar, S., Madhupratap, M., and DileepKumar, M. (2001). High biological productivity in the central Arabian Sea during the summer monsoon driven by Ekman pumping and lateral advection. *Curr. Sci.* 81, 1633–1638.
- Sañudo-Wilhelmy, S. A., Kustka, A. B., Gobler, C. J., Hutchins, D. A., Yang, M., Lwiza, K., et al. (2001). Phosphorus limitation of nitrogen fixation

by Trichodesmium in the central Atlantic Ocean. Nature 411, 66–69. doi: 10.1038/35075041

- Schott, F. A., and McCreary, J. P. (2001). The monsoon circulation of the Indian Ocean. Prog. Oceanogr. 51, 1–123. doi: 10.1016/S0079-6611(01)00083-0
- Shiozaki, T., Furuya, K., Kodama, T., Kitajima, S., Takeda, S., Takemura, T., et al. (2010). New estimation of N2 fixation in the western and central Pacific Ocean and its marginal seas. *Glob. Biogeochem. Cycles* 24:GB1015. doi: 10.1029/2009GB003620
- Shiozaki, T., Ijichi, M., Kodama, T., Takeda, S., and Furuya, K. (2014). Heterotrophic bacteria as major nitrogen fixers in the euphotic zone of the Indian Ocean. *Glob. Biogeochem. Cycles* 28, 1096–1110. doi: 10.1002/2014GB004886
- Singh, A., Gandhi, N., and Ramesh, R. (2012). Contribution of atmospheric nitrogen deposition to new production in the nitrogen limited photic zone of the northern Indian Ocean. J. Geophys. Res. Oceans 117:C06004. doi: 10.1029/2011jc007737
- Singh, A., Lomas, M., and Bates, N. (2013). Revisiting N2 fixation in the North Atlantic Ocean: significance of deviations from the Redfield Ratio, atmospheric deposition and climate variability. *Deep Sea Res. Part II Top. Stud. Oceanogr.* 93, 148–158. doi: 10.1016/j.dsr2.2013.04.008
- Singh, A., and Ramesh, R. (2011). Contribution of riverine dissolved inorganic nitrogen flux to new production in the coastal northern Indian Ocean: an assessment. *Int. J. Oceanogr.* 2011:983561. doi: 10.1155/2011/983561
- Singh, A., and Ramesh, R. (2015). Environmental controls on new and primary production in the northern Indian Ocean. *Prog. Oceanogr.* 131, 138–145. doi: 10.1016/j.pocean.2014.12.006
- Slawyk, G., Collos, Y., and Auclair, J. (1977). The use of the 13C and 15N isotopes for the simultaneous measurement of carbon and nitrogen turnover rates in marine phytoplankton. *Limnol. Oceanogr.* 22, 925–932. doi: 10.4319/lo.1977.22.5.0925
- Subramaniam, A., Mahaffey, C., Johns, W., and Mahowald, N. (2013). Equatorial upwelling enhances nitrogen fixation in the Atlantic Ocean. *Geophys. Res. Lett.* 40, 1766–1771. doi: 10.1002/grl.50250
- Sudheesh, V., Gupta, G., Sudharma, K., Naik, H., Shenoy, D., Sudhakar, M., et al. (2016). Upwelling intensity modulates N2O concentrations over the western

Indian shelf. J. Geophys. Res. Oceans 121, 8551-8565. doi: 10.1002/2016JC0 12166

- Turk, K. A., Rees, A. P., Zehr, J. P., Pereira, N., Swift, P., Shelley, R., et al. (2011). Nitrogen fixation and nitrogenase (nifH) expression in tropical waters of the eastern North Atlantic. *ISME J.* 5, 1201–1212. doi: 10.1038/ismej.2010.205
- Turk-Kubo, K. A., Achilles, K. M., Serros, T. R., Ochiai, M., Montoya, J. P., and Zehr, J. P. (2012). Nitrogenase (nifH) gene expression in diazotrophic cyanobacteria in the Tropical North Atlantic in response to nutrient amendments. *Front. Microbiol.* 3:386. doi: 10.3389/fmicb.2012.00386
- White, A. E. (2012). Oceanography: the trouble with the bubble. *Nature* 488, 290–291. doi: 10.1038/nature11481
- Wu, J., Sunda, W., Boyle, E. A., and Karl, D. M. (2000). Phosphate depletion in the western North Atlantic Ocean. *Science* 289, 759–762. doi: 10.1126/science.289.5480.759
- Zehr, J. P., Jenkins, B. D., Short, S. M., and Steward, G. F. (2003). Nitrogenase gene diversity and microbial community structure: a cross-system comparison. *Environ. Microbiol.* 5, 539–554. doi: 10.1046/j.1462-2920.2003. 00451.x
- Zehr, J. P., Mellon, M., Braun, S., Litaker, W., Steppe, T., and Paerl, H. W. (1995). Diversity of heterotrophic nitrogen fixation genes in a marine cyanobacterial mat. Appl. Environ. Microbiol. 61, 2527–2532.
- Zehr, J. P., Waterbury, J. B., Turner, P. J., Montoya, J. P., Omoregie, E., Steward, G. F., et al. (2001). Unicellular cyanobacteria fix N2 in the subtropical North Pacific Ocean. *Nature* 412, 635–638. doi: 10.1038/35088063

Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Copyright © 2017 Kumar, Singh, Ramesh and Nallathambi. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) or licensor are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.