Dinitrogen fixation in aphotic oxygenated marine environments

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INTRODUCTION

In many of the world’s oceans, the availability of dissolved inorganic nitrogen (N) controls primary production in surface waters (Falkowski, 1997; Karl et al., 2002). This is because the most abundant form of N in the oceans, dinitrogen (N2), is biologically unavailable to the majority of organisms that reside there. Subgroups of prokaryotic organisms (diazotrophs) fix N2 and convert it to ammonium via the nitrogenase enzyme complex. This biological process is an important source of “new” N to oceanic systems that can stimulate production and growth of microbial communities. Current models suggest that new N inputs from N2 fixation by cyanobacterial diazotrophs in the upper photic layers of the oceans is about 130–140 Tg N yr−1 (Gruber, 2008; Eugster and Gruber, 2012).

Globally, on geologic timescales, N inputs via N2 fixation are thought to be balanced by N losses via denitrification (Falkowski, 1997; Deutsch et al., 2007). However, at present, rate measurements suggest that N losses from the ocean via denitrification (∼245–322 Tg N yr−1) far exceed N inputs via N2 fixation and terrestrial sources (Codispoti et al., 2001; Gruber and Galloway, 2008). The balance between N2 fixation and denitrification (including annamox) is considered to critically impact the oceanic inventory of bioavailable nitrogen and thus primary productivity (PP) in the oceans (Gruber and Galloway, 2008).

Current models generally restrict oceanic N2 fixation to the surface, sunlit, photic layer of the oceans where autotrophic cyanobacterial diazotrophs occur (Gruber, 2008). However, recent work suggests that prokaryotic N2 fixers are more widely distributed geographically than previously thought (Moisander et al., 2010; Mulholland et al., 2012) and diazotrophs have been identified from more diverse pelagic environments including aphotic pelagic waters in oxygen minimum zones (OMZ) (Hammersley et al., 2011; Jayakumar et al., 2012). While it is known that diazotrophs inhabit shallow benthic environments and microbial mats (Capone, 1983; Bebout et al., 1987; Herbert, 1999), sea grass communities (McGlathery, 2008), coral reefs (O’Neil and Capone, 2008), and even hydrothermal vent communities (Mehta and Baross, 2006), the small areal extent of
these habitats is thought to limit their contribution to marine N\textsubscript{2} fixation (Brooks et al., 1971; Carpenter and Capone, 2008).

For many years, investigations of marine N\textsubscript{2} fixation also focused on oligotrophic oceanic regions because the presence of fixed N was considered inhibiting to active N\textsubscript{2} fixation (Mulholland and Capone, 1999; Flores and Herrero, 2005; Ward et al., 2009). Oligotrophic regions harbor a variety of cyanobacterial diazotrophs including the filamentous cyanobacteria *Trichodesmium* spp., diatom-diazotroph associations (DDA’s), and unicellular cyanobacteria such as *Crococpsphaera* and group A cyanobacteria (UCYN-A) (Bar-Zeev et al., 2008; Carpenter and Capone, 2008; Zehr et al., 2008). However, even *Trichodesmium*, the most widely studied marine diazotroph, can fix N\textsubscript{2} when ambient inorganic N concentrations are on ~5 \( \mu \)M (Mulholland et al., 2001), conditions rarely reached in most oceanic photic realms. N\textsubscript{2} fixation of marine diazotrophs has been observed in nutrient replete cultures (Mulholland et al., 2001) and systems with high dissolved inorganic N concentrations (Voss et al., 2004; Rees et al., 2009; Mulholland et al., 2012). Data compilation from euphotic, mesopelagic, and benthic studies shows that NO\textsubscript{3} and NH\textsubscript{3} concentrations, up to 30 and 200 \( \mu \)M respectively, can result in N\textsubscript{2} fixation rates similar to those found in nitrate depleted surface waters (reviewed in Knapp, 2012). Moreover, different sensitivities to DIN concentrations, exposure time, and compensation mechanisms all suggest that dissolved N concentrations do not control diazotroph distribution and activities in many marine environments (Knapp, 2012).

In addition to photoautotrophic diazotrophs that have been previously considered the dominant N\textsubscript{2} fixers in marine systems (Carpenter and Capone, 2008), heterotrophic diazotrophs from \( \alpha \), \( \beta \), \( \gamma \), \( \delta \), and \( \varepsilon \)-proteobacterial phylogenies, as well as methanogenic archaea contribute to marine N\textsubscript{2} fixation including benthic and pelagic realms, mesotrophic and eutrophic environments (Chien and Zinder, 1996; Wu et al., 2003; Zehr et al., 2003; Halm et al., 2012). Heterotrophic diazotrophs have been identified from NO\textsubscript{3}–rich deep, aphytic zones of the oceans (Riemann et al., 2010), including OMZ (Jayakumar et al., 2012) and from near hydrothermal vents (Mehta et al., 2005). In the subtropical Atlantic, cyanobacterial *nifH* genes dominated the photic layer (Langlois et al., 2005); while bacterial *nifH* genes dominated the deep aphytic zones (Riemann et al., 2010). In the ultraoligotrophic South Pacific Gyre (SPG) sequence analyses of *nifH* genes and their expressed transcripts revealed a prevalence of heterotrophic diazotrophic \( \gamma \)-proteobacteria (Cluster I—Zehr et al., 2003) and the unicellular phototrophic Group A cyanobacteria (UCYN-A) down to 200 m–the deep chlorophyll maximum (Halm et al., 2012).

While the presence of diazotrophs in oceanic aphytic zones has been demonstrated, to our knowledge, only three published studies have actually measured rates of N\textsubscript{2} fixation in aphytic environments; one study was conducted in the Eastern Tropical Pacific (Fernandez et al., 2011), another in the Southern California Bight (Hamesley et al., 2011), and the third in a hydrothermal vent community (Mehta and Baross, 2006). The first two studies were located in OMZ or hypoxic areas. In the Southern California Bight aphytic N\textsubscript{2} fixation accounted for up to one-third of the total areal rates of N\textsubscript{2} fixation (55 \( \mu \)mol N m\textsuperscript{−2} d\textsuperscript{−1} out of 150 \( \mu \)mol N m\textsuperscript{−2} d\textsuperscript{−1}) (Hamesley et al., 2011). While in the nutrient-rich OMZ of the Eastern Tropical South Pacific (ETSP), N\textsubscript{2} fixation rates from sub-surface oxygen deficient waters contributed about 5 times more N than N\textsubscript{2} fixation from the oxic photic layer (Fernandez et al., 2011). Moreover, N\textsubscript{2} fixation co-occurred spatially and temporally with denitrification in the ETSP (Fernandez et al., 2011), refuting the paradigm of spatial and/or temporal uncoupling between the two processes.

In oligotrophic environments such as the Levantine Basin (LB) (Eastern Mediterranean Sea, EMS) and the Gulf of Aqaba (northern Red Sea, GA), primary production in surface waters is limited at times by nitrogen (N), phosphorus (P), or co-limited by both (Paytan et al., 2009; Krom et al., 2010; Tanaka et al., 2011). Denitrification in subsurface waters is absent because waters are fully oxygenated (Kress and Herut, 2001; Al-Qutob et al., 2002). In the EMS, new nitrogen inputs via N\textsubscript{2} fixation in the photic zone are generally very low, with rates ranging from 0.01 to 0.3 nmol N L\textsuperscript{−1} d\textsuperscript{−1}. These rates contribute only 1–2% of new production in the LB (Yoge et al., 2011; Berman-Frank and Rahav, 2012; Rahav et al., 2013a). Although occasional small blooms of *Trichodesmium* and DDAs occur in the GA, N\textsubscript{2} fixation rates are still relatively low ranging from ~0.05 to 2 nmol N L\textsuperscript{−1} d\textsuperscript{−1} (Post et al., 2002; Foster et al., 2009). In the photic layer of the EMS and the GA, diazotroph populations include cyanobacteria as well as many heterotrophic microbes (Man-Aharovovich et al., 2007; Foster et al., 2009; Yoge et al., 2011).

In this study we measured N\textsubscript{2} fixation rates in the aphytic, oxygenated waters of the oligotrophic LB and the GA. Our results demonstrate significant N\textsubscript{2} fixation rates equal or greater to those measured from the photic layers and a prevalence of heterotrophic diazotrophs. This initial data suggests the aphytic oxygenated zones can be considerable sources of new N supporting biological production in oligotrophic waters.

**MATERIALS AND METHODS**

**SAMPLING LOCATIONS**

Seawater samples were collected from two representative pelagic stations that were sampled several times during the mixed and stratified periods; one in the GA in the Red Sea and one in the LB in the Eastern Mediterranean Sea (Table 1 and below). Water samples were collected using 10 L Niskin bottles mounted on a rosette equipped with a CTD (Seabird 19 Plus) and fluorometer (Turner designs, Cyclops7) to measure hydrographic properties of the water column, and an *in situ* PAR sensor (QSP-2300, Biospherical Instruments Inc) allowing distinction between the photic and aphytic zones.

**Gulf of Aqaba (GA)**—Water samples from the GA were collected aboard the *R/V Sam Rothberg* at a station located at the northern tip of the Red Sea (Station A 29° 55’N, 34° 28’E) during the winter when the water column was well-mixed (March 2010) and twice during the stratified summer period (September 2010 and July 2012).

**Levantine Basin (LB)**—Samples from the LB were collected during cruises aboard the *R/V Mediterranean Explorer* and the *R/V Shikmona*. A station located west of the Israeli coastline (Station OUT; 32° 57’N, 34° 45’E) was sampled during periods when the water column was well mixed (December 2010,
February 2012, March 2013) and stratified (August 2011, October 2011, April 2012).

**INORGANIC NUTRIENTS**

Duplicate unfiltered water samples were collected in 15-mL acid-washed plastic scintillation vials and immediately frozen at −20°C until they were analyzed with a segmented flow Skalar SAN plus System Instrument as detailed in Kress and Herut (2001). The precisions of nitrate+ nitrite (hereafter NO₃), orthophosphate (PO₄) and silicic acid (Si(OH)₄) measurements were 0.02, 0.003 and 0.06 μM, respectively. The limits of detection were 0.075 μM, 0.008 μM and 0.03 μM for NO₃, PO₄ and Si(OH)₄, respectively.

**DINITROGEN (N₂) FIXATION RATES**

Rates of N₂ fixation were measured in field-collected samples (4.5 L) using the ¹⁵N₂ bubble addition technique described by Montoya et al. (1996) and Mulholland et al. (2006) using highly enriched ¹⁵N₂ gas (99%). The seawater was dispensed into 4.5 L Nalgene incubation bottles, 9 mL of ¹⁵N₂ was added, and bottles were then placed in an incubator with continuously flowing seawater to maintain ambient surface-water temperatures. Triplicate bottles were used for each depth sampled. The incubator was shaded with neutral density screening to simulate ambient irradiance conditions at the depths from which samples were collected. For samples collected from the aphotic layer, bottles were incubated in the dark. Both light and dark incubations were 24–48 h in duration. While the bubble method may underestimate N₂ fixation (Mohr et al., 2010; Großkopf et al., 2012; Wilson et al., 2012), particularly during short incubations (hours), our long incubation times should have minimized this problem (Mulholland et al., 2012; Rahav et al., 2013b). See also the discussion regarding underestimation of rates due to the bubble vs. enriched seawater method (Mohr et al., 2010). To terminate the incubations, water was filtered onto pre-combusted 25 mm GF/F filters (450°C for 4 h). Filters were dried and pelletized in tin disks and analyzed on a Europa 20/20 mass spectrometer equipped with an automated nitrogen and carbon analyzer preparation module. N₂ fixation was calculated using a mixing model (Orcutt et al., 2001) as described by Mulholland et al. (2006) with N solubility factors according to Weiss (1970).

Depth-integrated N₂ fixation rates were based on measurements made at 3 to 6 water column depths spanning both photic and aphotic layers. In months where N₂ fixation measurements were made at only 2 depths, no such calculations were made (i.e., Not Available, N.A; Table 2). Depth-integrations included only the portion of the water column sampled (range was 150–720 m in depth) and did not extend to the bottom except for one case in the GA during July 2012, when N₂ fixation rates were measured also in bottom waters (Table 1). The percent contribution of the aphotic N₂ fixation was calculated by difference after subtracting out the contribution of photic N₂ fixation from the total integrated water column N₂ fixation rate.

**PRIMARY PRODUCTIVITY**

Photosynthetic carbon fixation rates were estimated using ¹³C-labeled bicarbonate (Mulholland and Bernhardt, 2005). Water samples were placed in triplicate clear 4.5 L polycarbonate Nalgene bottles and amended with 99% NaH¹³CO₃ (Sigma) to obtain enrichments of about 1% of the ambient dissolved inorganic carbon pool. Bottles were incubated under the same conditions as for the ¹⁵N₂ incubation bottles described above. Parallel dark bottles were also incubated for photic depths. As for ¹⁵N₂, ¹³C incubations were terminated by immediately filtering the entire contents of incubation bottles onto pre-combusted 25 mm GF/F filters (450°C for 4 h). Filters were stored at −20°C and then dried and pelletized in tin disks prior to their analysis on the Europa 20/20 mass spectrometer. Rates of C uptake were calculated using a mixing model (as described above) and dark carbon fixation was subtracted from the light carbon fixation rates for photic incubations.

**BACTERIAL PRODUCTIVITY**

Bacterial production was estimated using the ³H-leucine (Amersham, Specific activity: 160 Ci mmol⁻¹) incorporation method (Simon et al., 1990). Triplicate 1.7 ml samples were incubated for 4–8 h at *in situ* temperatures in the dark. Triplicate, trichloroacetic acid (TCA) killed samples (Sigma T9159) served as controls. One ml of high ³H affinity scintillation cocktail (ULTIMA-GOLD) was added to samples and they were counted using a TRI-CARB 2100 TR, PACKARD scintillation counter.

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**Table 1 | Summary of the sampling dates, locations, and depths.**

| Region        | Sampling location     | Sampling duration | Bottom depth (m) | Photic depths sampled (m) | Aphotic depths sampled (m) | Depth of 0.1% light (m) |
|---------------|-----------------------|-------------------|------------------|---------------------------|---------------------------|-------------------------|
| Red Sea       | Gulf of Aqaba (GA)    | Mar. 2010         | 720              | 0.20,50,100               | 150                       | 105                     |
|               |                       | Aug. 2010         | 0.50, 100        | 200,300,500               | 120                       |                         |
|               |                       | July 2012         | 10,85            | 160, 300, 500, 720        | 100                       |                         |
| Med. Sea      | Levantine basin (LB)  | Dec. 2010         | 1100             | 0.20,50,100               | 300, 500                  | 110                     |
|               |                       | Aug. 2011         | 5                | 250                       | 140                       |                         |
|               |                       | Oct. 2011         | 5                | 300                       | 130                       |                         |
|               |                       | Feb. 2012         | 5                | 300                       | 110                       |                         |
|               |                       | Apr. 2012         | 5                | 300                       | 110                       |                         |
|               |                       | Mar. 2013         | 5, 80            | 250                       | 100                       |                         |

Additional details are available in the Material and methods section. Med. Sea, Mediterranean Sea.
Table 2 | Measured physical, chemical, and biological properties of the photic and aphotic layers from the sampling stations in the Levantine basin and the Gulf of Aqaba averaged seasonally for when the water column was mixed (winter) and during thermal stratification (spring, summer, and fall).

| Location     | Depth    | Period | Temp. (°C) | Salinity | NO₂ + NO₃ (µM) | PO₄ (µM) | BP (µg C L⁻¹ d⁻¹) | PP (µg C L⁻¹ d⁻¹) | N₂ fixation (nmol N L⁻¹ d⁻¹) | Contribution of layer to total N₂ fixation (%) |
|--------------|----------|--------|------------|----------|----------------|----------|------------------|------------------|-----------------------------|---------------------------------|
| Gulf of Aqaba| Mixed    | 22–23  | 40.5–40.7  | 0.09–0.90| BLD-0.04       | 0.4–0.8  | 0.1–3.1          | 0.05–0.11         | 63                          |
|              | Stratified| 23–27  | 40.5–40.8  | 0.04–0.06| BLD-0.04       | 1.5–3.8  | 0.3–0.6          | 0.35–0.63         | 25                          |
| Levantine basin| Mixed    | 17–18  | 39.1–39.2  | 0.06–0.46| BLD-0.20       | 0.4–0.5  | 0.7–2.3          | BLD-0.01          | 44                          |
|              | Stratified| 18–28  | 39.2–39.5  | 0.02–0.10| BLD-0.03       | 0.4–2.5  | 0.4–1.8          | 0.01–0.17         | N.A                          |
| Gulf of Aqaba| Aphotic  | 21–22  | 40.5–40.7  | 0.80–5.61| BLD-0.30       | 0.2–0.4  | 0–0.1            | 0.02–0.20         | 37                          |
|              | Stratified| 21–23  | 40.4–40.7  | 0.06–6.24| BLD-0.30       | 0.5–2.4  | 0               | 0.07–0.38         | 75                          |
| Levantine basin| Mixed    | 14–18  | 38.9–39.1  | 0.20–4.31| BLD-0.20       | 0.2–2.7  | 0.3–0.5          | 0.01–0.02         | 56                          |
|              | Stratified| 15–18  | 38.9–39.1  | 0.10–4.52| BLD-0.20       | 4.6–10.7 | N.A              | 0.01–0.24         | N.A                          |

N.A, not available, BLD, below detection limit.

AMINO-ACIDS ENRICHMENT EXPERIMENT

We assumed that enriching the water with amino acids would promote conditions that were advantageous to heterotrophic bacteria in general and also to heterotrophic diazotrophs. We assumed that if heterotrophic diazotrophic bacteria were limited by energy (carbon), the addition of high-carbon containing amino acids would alleviate this limitation allowing these populations to fix N₂. The idea was to distinguish general bacterial heterotrophic activity (bacterial production rates) from diazotrophic ones (N₂ fixation rates). The response of the microbial community to additions of an amino acid mixture (Sigma A9906) was tested at 4 aphotic (160, 300, 500, 720 m) and two photic (10, 85) depths (Tables 1, 2) during July 2012 in the GA. Amino acids were added to 4.5 L polycarbonate Nalgene bottles filled with whole water from a specified depth at a final concentration of 500 nM C immediately prior to making ¹⁵N₂ additions.

PYROSEQUENCING

DNA was extracted from a representative aphotic depth in the LB stations. Fifteen L of seawater were filtered onto 0.2 µm Supor® filter (PALL) and stored in −80°C until analysis in the lab. DNA was extracted using a genomic DNA kit (Roche) and the extract was purified with a QIAGEN purification kit according to the manufacturer’s instructions. Amplification of nifH genes from genomic DNA was performed as described elsewhere (Zehr and McReynolds, 1989). The nifH sequences retrieved from the pyrosequencing analyses [performed by Research and Testing Laboratories (MRDNA, Texas, USA)] were trimmed and checked for quality using QIIME (Caporaso et al., 2010). The cleaned sequences were compared to a nifH database (Zehr et al., 2003). In the aphotic layer of the LB only 20 sequences with an open reading frame were retrieved. These sequences were further confirmed using BLASTX (Altschul et al., 1990) with a cutoff of 95%. The nifH-like sequences were integrated into a nifH phylogenetic tree created with MEGA 4.0 software described in details by Tamura et al. (2007). nifH gene sequences were submitted to GeneBank and were assigned the accession numbers KC222033–KC222052.

DETERMINATION OF TRANSPARENT EXOPOLYMER PARTICLES (TEP) CONCENTRATION AND SIZE

Triplicate water samples (100 ml) were filtered onto a 0.4 µm polycarbonate filters and stained for 2 s with a 0.02% Alcian Blue solution. TEP concentrations (µg GX equivalents L⁻¹) were measured according to Passow and Alldredge (1995). A conversion factor of 0.51 was used to convert from micrograms of Gum Xanthan (GX) equivalents to micrograms of carbon (Engel and Passow, 2001).

For TEP size (area - µm²) measurements, samples (100 ml) were filtered gently (~100 mbar) onto 0.4 µm polycarbonate filters, stained for 10 s with freshly prepared Alcian Blue solution (0.02%), rinsed with 5 ml of double distilled water (DDW), placed onto microscope slides and kept at 4°C. The filters were analyzed under X40 magnification after 1–2 days and at least 90 pictures per slide were taken. The pictures were analyzed with image J software (http://rsbweb.nih.gov/ij/) to determine the size of TEP.

TEP ENRICHMENT EXPERIMENT

The association of aphotic diazotrophs with TEP was explored by adding 4 different quantities of pure GX (Sigma G1253) to LB seawater only. The bottles were incubated in the dark at ambient temperature (15°C) for 24 h, after which they were sampled to measure TEP concentration and size and rates of N₂ fixation and BP.

STATISTICAL ANALYSES

Data were displayed as means with corresponding standard deviations for correlation and regression analyses between variables. Comparisons between unamended controls (no addition) and the amended microcosms (+Dissolved free amino-acids, DFAA) at the end of the experiment, a One-Way ANOVA followed by Tukey’s multiple comparison post hoc using R software (www.r-project.org). P < 0.05 was considered statistically significant.
RESULTS

PHYSICAL AND CHEMICAL CHARACTERISTICS OF THE APHOTIC LAYER

The aphotic layer was determined at each station as the depth below which irradiance was <0.1% of surface irradiance. The depths of the photic layer varied between stations and seasons, ranging from 105 m to 140 m at both sampling locations (Table 1). Temperature and salinity in the aphotic layer of the GA station were 21.5 ± 0.5°C and 40.6 ± 0.1, respectively, during both the mixed (winter) and stratified (summer) periods. In the photic layer of the GA, temperatures were 22 ± 0.5°C during winter and 25 ± 2°C during the stratified months (Figures 1A,B; Table 2). At the LB station, temperatures and salinities of the aphotic layer were significantly lower than those measured for the GA station, averaging 16.1 ± 1.1°C and 39.1 ± 0.1, respectively (Figures 1C,D; Table 2), while within the photic layer temperature and salinities averaged 23 ± 5°C and 39.3 ± 0.2, respectively, during the stratified period (Figures 1A,B; Table 2).

Average dissolved NO₃ concentrations in the nitricline and sub-nitricline depths of the GA were 3.0 ± 2.1 μM with the nitricline ranging between 200–400 m during the winter mixing (n = 9, Figure 1A; Table 2). NO₃ concentrations remained similar (3.0 ± 2.5 μM) during the stratified summer for the same depths although the nutricline deepened to 150–500 m (n = 13, Figure 1B; Table 2). PO₄ (DIP) concentrations varied similarly in both periods, averaging 0.2 ± 0.2 μM (Figures 1A,B; Table 2).

Figure 1 | Spatial (depth) and temporal changes of physical and chemical parameters at the sampling stations in the Gulf of Aqaba (A,B) and Levantine basin (C,D) during the mixed (A,C) and stratified (B,D) periods.

Surface NO₃ and DIP concentrations were low and close to their detection limit during both periods (Figures 1A,B; Table 2).

Unlike the GA, no seasonal fluctuations in NO₃ and DIP concentrations were recorded in the aphotic depths of the pelagic LB station (Figures 1C,D). At these depths lower NO₃ and DIP concentrations were 2.2 ± 2.1 μM and 0.1 ± 0.5 μM, respectively (n = 7–9, Figures 1C,D; Table 2).

N₂ FIXATION RATES

Photic N₂ fixation rates compiled from both stations, at all depths and seasons yielded rates usually below 0.1 nmol N L⁻¹ d⁻¹ excluding rates measured in the GA during summer (0.35–0.63 nmol N L⁻¹ d⁻¹) (Table 2). Aphotic N₂ fixation rates ranged from 0.01 to 0.38 nmol N L⁻¹ d⁻¹ at both locations sampled (Table 2). Aphotic rates obtained in the GA during the mixed period ranged from 0.02 to 0.2 nmol N L⁻¹ d⁻¹ and averaged 0.1 ± 0.1 nmol N L⁻¹ d⁻¹ while during the stratified period rates ranged from 0.07 to 0.38 nmol N L⁻¹ d⁻¹ and averaged 0.30 ± 0.10 nmol N L⁻¹ d⁻¹ (Table 2). At the LB station, aphotic N₂ fixation was uniformly low during the mixed period; 0.01 ± 0.01 nmol N L⁻¹ d⁻¹ and increased ~20-fold during the stratified period to 0.24 ± 0.09 nmol N L⁻¹ d⁻¹ (Table 2).

To understand the significance of aphotic N₂ fixation relative to the total areal rates of N₂ fixation, we calculated that during our sampling times, N₂ fixation from aphotic depths sampled accounted for 37 to 75% of total areal N₂ fixation at the GA station and ~50% of the total areal N₂ fixation at the LB station during the mixed period (Table 2).

PHYTOPLANKTON AND BACTERIAL PRODUCTIVITY RATES

PP rates in the GA averaged 3.0 ± 0.1 μg C L⁻¹ d⁻¹ and 0.5 ± 0.1 μg C L⁻¹ d⁻¹ for the mixed and stratified periods, respectively (n = 3, PAR at surface of 1000–1500 μmol quanta m⁻² s⁻¹). PP at the LB station was generally lower; averaging 2.1 ± 0.2 μg C L⁻¹ d⁻¹ and 1.8 μg C L⁻¹ d⁻¹ for the mixed and stratified periods, respectively (n = 3, PAR at surface of 1500–1800 μmol quanta m⁻² s⁻¹) (Table 2). Bicarbonate uptake rates were usually below detection limits within the aphotic zone (below 105–140 m, depending on sampling location) excluding a wintertime measurement in the LB station (200 m) when C uptake averaged 0.4 ± 0.1 μg C L⁻¹ d⁻¹ (Table 2).

Bacterial productivity (BP) rates in the photic waters from both locations ranged from 0.4 to 3.8 μg C L⁻¹ d⁻¹, while in the aphotic zone BP rates were 0.2 to 10.7 μg C L⁻¹ d⁻¹ (Table 2). Rates of N₂ fixation from both the photic and aphotic layers were compared to BP and PP (Figure 2). BP correlated significantly and positively with N₂ fixation rates within the photic zone of both basins (R² = 0.70, P = 0.001, n = 12), while there was no apparent correlation between N₂ fixation rates and PP (Figure 2A). Moreover, BP and N₂ fixation were also positively correlated within the aphotic depths (R² = 0.58, P = 0.003, n = 10) (Figure 2B).

To further understand what limits N₂ fixation by heterotrophic diazotrophs in oligotrophic and aphotic systems we added dissolved free amino acids (DFAA) to microcosm experiments conducted in the oligotrophic stratified period during July 2012 in the GA. In water samples collected from the aphotic layer, DFAA
additions enhanced BP rates by a factor of 2–3.5 and N₂ fixation rates increased by a factor of 1.5–2 (ANOVA, P < 0.05) (Table 3). Within the photic layer, the response of bacteria to DFAA additions was greater than that observed in the aphotic zone; BP increased 2- to 6-fold and N₂ fixation rates increased by a factor of 2 to 4 (ANOVA, P < 0.05) (Table 3).

### Table 3 | The effect of dissolved free amino acid (DFAA) enrichments (+ DFAA) on bacterial productivity (BP) and N₂ fixation during the stratified summer period (GA, July 2012) within different depths of the photic and aphotic layers.

| Sampling depth (m) | BP (µg C L⁻¹ d⁻¹) | N₂ fixation (nmol N L⁻¹ d⁻¹) |
|-------------------|-----------------|----------------|----------------|----------------|
|                   | Control         | + DFAA         | Control         | + DFAA         |
| Photic            |                 |                |                 |                |
| 10                | 0.30 ± 0.40     | 2.40 ± 0.38    | 0.35 ± 0.09     | 0.76 ± 0.10    |
| 85                | 1.15 ± 0.33     | 4.17 ± 0.68    | 0.30 ± 0.11     | 1.12 ± 0.27    |
| Aphotic           |                 |                |                 |                |
| 160               | 2.34 ± 0.69     | 4.68 ± 0.14    | 0.38 ± 0.05     | 0.70 ± 0.20    |
| 300               | 2.20 ± 0.70     | 5.32 ± 0.20    | 0.23 ± 0.05     | 0.34 ± 0.02    |
| 500               | 0.46 ± 0.07     | 1.57 ± 0.19    | 0.07 ± 0.03     | 0.13 ± 0.06    |
| 720               | 2.38 ± 0.43     | 4.86 ± 0.47    | 0.16 ± 0.10     | 0.31 ± 0.06    |

Measurements are based on 3-4 replicates for each sampled depth.

**FIGURE 2** | The relationship between N₂ fixation and primary and bacterial productivity within the photic (A) and aphotic (B) layers of the LB and GA stations. Values are averages and error bars are standard deviations of three independent replicates performed for each control and treatment incubation.

**TEP AS SOURCE OF CARBON OR MICROHABITAT**

TEP concentrations ranged from 100 to 5000 µg GX L⁻¹ and varied in size; the two-dimensional TEP size ranged from 24 to 40 µm² with the largest particles measured at TEP concentrations of ~3500 µg GX L⁻¹ (Figure 3). BP increased almost 7-fold, from ~0.6 µg C L⁻¹ d⁻¹ in the unamended control bottles to ~4 µg C L⁻¹ d⁻¹ in incubations amended with the highest TEP concentrations (5000 µg GX L⁻¹). N₂ fixation was also stimulated (~5-fold) by TEP additions increasing from 0.06 nmol N L⁻¹ d⁻¹ in the controls to 3.46 nmol N L⁻¹ d⁻¹ in the bottles amended with ~3500 µg GX L⁻¹. N₂ fixation rates were positively correlated with both TEP concentration (R² = 0.49, P = 0.07, n = 15) and TEP size (R² = 0.90, P = 3 x 10⁻⁵, n = 15) (Figure 3).

**MOLECULAR DIVERSITY of nifH**

NifH-like genes were retrieved from the aphotic layer of the LB and were compared with nifH genes from known bacterial isolates. Twenty sequences were retrieved and all clustered together with heterotrophic bacteria and were most similar to *Pseudomonas stutzeri* (Figure 4). All of our sequences were separated from the autotrophic clusters.

**DISCUSSION**

Most marine N₂ fixation studies neglect the potential contribution of biological N₂ fixation in deep, aphotic, NO₃-rich waters that comprise the majority of the world’s oceans (Karl et al., 2002; Zehr and Ward, 2002). This is likely because phototrophic and cyanobacterial diazotrophs have an ecological advantage in surface waters where productivity is often limited by the availability of fixed nitrogen yet there is sufficient energy (irradiance) to drive both photosynthetic C uptake and the costly process of N₂ fixation (Howarth et al., 1988; Postgate, 1998; Karl et al., 2002). Currently these assumptions are being modified as significant N₂ fixation rates are reported from marine areas with high DIN concentrations (review in Knapp, 2012). Here we provide evidence that aphotic, nitrogen-rich waters also harbor active diazotrophs that contribute substantial new N to the oceanic N inventory, and that N₂ fixation by these bacterial diazotrophs may be limited by the supply of organic C.

In this study we have documented for the first time significant N₂ fixation rates from the aphotic, oxidized waters (Figure 1) of...
the oligotrophic LB and the GA. The measured N₂ fixation rates in the aphotic waters of both LB and GA ranged from 0.01 nmol N L⁻¹ d⁻¹ to 0.38 nmol N L⁻¹ d⁻¹. These rates were similar to the N₂ fixation rates reported from aphotic depths in the hypoxic Southern California Bight (Hammersley et al., 2011), yet were ~10–100 fold lower than those measured in the Peruvian upwelling OMZ (Fernandez et al., 2011).

The significance of our measured rates was realized upon integration of water column N₂ fixation from surface to the deepest depth sampled (Table 1). Thus, the contribution of aphotic N₂ fixation accounted for ~50% from the total N₂ fixation in the LB station, where winter mixing is shallow and the aphotic water hardly mixes with the photic layer, and 37 to 75% within the GA station (Table 2).

FIGURE 3 | The relationship between N₂ fixation rates of aphotic diazotrophs and transparent exopolymer particles (TEP) concentrations (A) and size (B) in experimental manipulations of LB waters from 250 m depth to which we added 4 concentrations of pure gum xanthan. Values are averages and error bars are standard deviations of three independent replicates performed for each control and treatment incubation.

FIGURE 4 | nifH gene tree of clones retrieved from the aphotic layer in the Levantine Basin. Numbers in parentheses represent the number of clones in the library that were identical to the clones in the phylogenetic tree. Bootstrap values above 50% are indicated above the nodes.
The substantial role of aphotic \( \text{N}_2 \) fixation in the LB and the GA may be especially important to basin-wide productivity given the low rates of \( \text{N}_2 \) fixation in the photic layers of both these nutrient-poor environments (Foster et al., 2009; Berman-Frank and Rahav, 2012). Within the photic layer, \( \text{N}_2 \) fixation was undetectable in a third of the measurements made in the Eastern Mediterranean Sea during several studies \((n > 250)\) spanning seasons and years (Ibello et al., 2010; Yogev et al., 2011). In contrast, measureable rates of \( \text{N}_2 \) fixation were observed in \( >90\% \) of samples collected from the aphotic zone \((n = 48)\), regardless of season, thermal stability or depth (Table 2), further emphasizing the importance of \( \text{N}_2 \) fixation in these realms and the requirement for subsequent comprehensive investigations that could constrain modeling efforts.

Our measured aphotic \( \text{N}_2 \) fixation rates may actually represent an underestimate of this process as: (1) our sampling did not cover the full depth of the water column—thus integration may yield higher contribution of aphotic to the total \( \text{N}_2 \) fixation (see methods); (2) we used the common gas bubble enrichment method (Montoya et al., 1996 and see Materials and Methods) which may underestimate \( \text{N}_2 \) fixation rates by a factor of 2–6 compared to the enriched seawater method (Großkopf et al., 2012; Wilson et al., 2012). Early field comparisons from the LB photic waters between the two methods showed a 2–3 fold underestimation by the gas bubble technique (Rahav et al., 2013b). Assuming our aphotic rates were underestimated by a similar ratio of \( \sim 50\% \), our measured aphotic \( \text{N}_2 \) fixation would increase accordingly so that maximal aphotic rates would be \( \sim 0.8 \text{ nmol N L}^{-1} \text{ d}^{-1} \), and with DFAA addition up to 1.4 \( \text{ nmol N L}^{-1} \text{ d}^{-1} \). Yet, as the estimated increase by \( 50\% \) would also be accounted for in photic rates of \( \text{N}_2 \) fixation, the \% contribution of aphotic to total \( \text{N}_2 \) fixation would remain the same.

Energetic requirements for diazotrophy are high (Postgate, 1998), thus we initially assumed that phototrophic (mainly cyanobacterial) diazotrophs would dominate the photic zone, while at aphotic depths, \( \text{N}_2 \) fixation rates would be extremely low and accomplished by heterotrophic bacterial populations that would be prone to carbon limitation due to the low overlying productivity. However, we found that in the LB and GA stations heterotrophic diazotrophs were the dominant \( \text{N}_2 \) fixers in both photic and aphotic zones. This was confirmed by examining the relationship between \( \text{N}_2 \) fixation and BP and PP (Figure 2), molecular analysis of \( \text{nifH} \) genes (Figure 4), and the stimulation of \( \text{N}_2 \) fixation after amino acid amendments (Table 3). In the stratified photic zone, a significant correlation was found between BP and \( \text{N}_2 \) fixation \((R^2 = 0.70, P = 0.001, n = 10)\) and both were stimulated by the addition of DFAA (Table 3). In contrast, no correlation was found between \( \text{N}_2 \) fixation and PP (Figure 2A). Although ample solar irradiance was available, heterotrophic bacterial diazotrophs appear to dominate in this system relying on dissolved and particulate organic carbon (DOC and POC, respectively) as energy sources (Goldthwait et al., 2005; Azam and Malfatti, 2007; Eichinger et al., 2009).

Within the aphotic layer, \( \text{N}_2 \) fixation was positively correlated with BP \((R^2 = 0.58, P = 0.003, n = 10)\), (Figure 2). Moreover, a 2–3 fold enhancement of \( \text{N}_2 \) fixation rates and enhanced BP were observed following DFAA amendments (Table 3). DFAA are labile dissolved organic carbon (DOC) and nitrogen (DON) sources for microbes (Middelburg and Nieuwenhuize, 2001) that can be directly utilized by bacterial heterotrophs (Davis et al., 2009). Our data thus suggests that the addition of C and N sources stimulated both diazotrophic and non-diazotrophic heterotrophic populations. Moreover, the increase in N source comitantly with enhanced \( \text{N}_2 \) fixation rates demonstrate that the higher concentration of dissolved N sources (either DON or bacterially catabolized into DIN) were not sufficient to inhibit diazotrophy. These results highlight the significant role of heterotrophic bacteria as important components of the photic and aphotic diazotrophic community and suggest that the availability of organic matter may limit bacterial diazotrophy in these oligotrophic regimes.

Previous molecular characterization of diazotrophs that were actively expressing \( \text{nifH} \) genes from the photic zones of the LB and GA demonstrated a diverse array of both heterotrophic bacteria and cyanobacterial phylotypes from Clusters I, II, and III (Foster et al., 2009; Yogev et al., 2011). In this study, phylogenetic analyses of \( \text{nifH} \) from the aphotic depths indicated that diazotrophs present were within Clusters I and III, phylotypes predominately comprised of heterotrophic bacteria, including \( \alpha \) and \( \gamma \) proteobacteria (Chien and Zinder, 1996) (Figure 4). The \( \text{nifH} \) phylogenetic diversity obtained from the aphotic zone during this study was similar to that recovered from the Southern California Bight (Hammersley et al., 2011) and anaerobic bacteria from the Sargasso Sea (Hewson and Fuhrman, 2007) (Figure 4).

Our results suggest that in the aphotic zones of both LB and GA heterotrophic bacterial \( \text{N}_2 \) fixation is a considerable source of new N. Yet these environments have not heretofore been considered as major sources of new oceanic N inputs, likely because they are dark, oxygenated, and \( \text{NO}_3 \)-rich (Figure 1). Heterotrophic diazotrophs in aphotic areas may be carbon (energy) limited, especially in oligotrophic systems where export of organic particulate matter to depth is low. This is illustrated by the uncoupling between BP and PP observed in our study sites. In this study we demonstrated that \( \text{N}_2 \) fixation and BP rates were enhanced when DFAA were added (Table 3). In addition to labile organic compounds such as DFAA, the mesopelagic aphotic layer is rich in organic carbon in the form of planktonic microgelatinous marine snow (Azam, 1998; Azam and Malfatti, 2007). This includes protobiofilms that are transparent exopolymer particles (TEP), heavily colonized by bacteria (Bar-Zeev et al., 2011, 2012). In oligotrophic systems these protobiofilm shuttles contribute significantly to the organic carbon pool especially at depths where other C sources are scarce (Azam and Malfatti, 2007; Ward et al., 2009; Bar-Zeev et al., 2012). While these particles are rich in C, the high C:N ratios of TEP (Engel and Passow, 2001; Passow, 2002) make them poor N sources and would seem to require the use of some N supplement to foster microbial growth. Heterotrophic \( \text{N}_2 \) fixation could provide an excellent source of N to fuel microbial growth on, within, and near these particles. Indeed, our data demonstrate a positive relationship between TEP concentration \((R^2 = 0.49, P = 0.07, n = 15)\), TEP size \((R^2 = 0.90, P = 3 \times 10^{-5}, n = 15)\), and \( \text{N}_2 \) fixation (Figure 3). The ability to fix \( \text{N}_2 \) could provide diazotrophic
heterotrophs with a competitive advantage over non-diazotrophic heterotrophic bacteria in carbon-rich and nitrogen-poor microenvironments such as these. Moreover, in aggregates such as marine snow, TEP, and other marine microgrids, bacterial colonization causes high metabolic consumption of oxygen and decreasing oxygen gradients from the surface to the anaerobic center (Pael and Prufert, 1987). Reduced oxygen concentrations would theoretically benefit diazotrophic bacteria because nitrogenase is an oxygen-sensitive protein (Postgate, 1998). We speculate these anaerobic, high C, low N microenvironments could promote the competitive success and activity of heterotrophic diazotrophs at depth.

Our novel data thus provides a conservative estimate of an additional 37 to 75% source of N via aphytic N2 fixation in the oligotrophic Levantine basin and the GA. Our measurements substantiate early indications (Pael and Prufert, 1987) and recent phylogenetic characterizations (Riemann et al., 2010) highlighting a pivotal ecological role for diazotrophs in deep, oxygenated, NO3-rich waters. Because these areas comprise the bulk of the world’s oceans they could represent a hitherto unaccounted for source of new N that may contribute to global productivity and could reduce the existing discrepancies [along with other newly described contributions of N2 fixation (reviewed in Knapp, 2012)] between sources and losses of N found in global marine N budgets.

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REFERENCES

Al-Qoutob, M., Hase, C., Tilzer, M. M., and Lazar, B. (2002). Phytotrophon drives nitrite dynamics in the Gulf of Aqaba, Red Sea. Mar. Ecol. Prog. Ser. 239, 233–239. doi: 10.3354/meps239233

Altschul, S. F., Gish, W., Miller, W., Myers, E. W., and Lipman, D. J. (1990). Basic local alignment search tool. J. Mol. Biol. 215, 403–410.

Azam, F. (1998). Microbial control of oceanic carbon flux: the plot thickens. Science 280, 694–696. doi: 10.1126/science.280.5364.694

Azam, F., and Malfatti, F. (2007). Microbial structuring of marine ecosystems. Nat. Rev. Microbiol. 5, 782–791. doi:10.1038/nrmicro1747

Bar-Zeev, E., Berman, T., Rahav, E., Dishon, G., Herut, B., and Berman-Frank, I. (2011). Transparent exopolymer particle (TEP) dynamics in the Eastern Mediterranean Sea. Mar. Ecol. Prog. Ser. 431, 107–118. doi:10.3354/meps09110

Bar-Zeev, E., Berman-Frank, I., Girshavit, O., and Berman, T. (2012). Revised paradigm of aquatic biofilm formation facilitated by microgel transparent exopolymer particles. Proc. Natl. Acad. Sci. U.S.A. 109, 9119–9124. doi:10.1073/pnas.1203708109

Bar-Zeev, E., Yogev, T., Man-Aharonovich, D., Kress, N., Herut, B., Béja, O., et al. (2008). Seasonal dynamics of the endosymbiotic, nitrogen-fixing cyanobacterium Richelia intracellularis in the Eastern Mediterranean Sea. ISME J. 2, 911–923. doi:10.1038/ismej.2008.56

Bebout, B. M., Pael, H. W., Crocker, K. M., and Prufert, L. E. (1987). Diel interactions of oxygen photosynthesis and N2 fixation (acetylene-reduction) in a marine microbially mat community. Appl. Environ. Microbiol. 53, 2353–2362.

Berman-Frank, I., and Rahav, E. (2012). “Nitrogen fixation as a source for new production in the Mediterranean Sea: a review,” in Life in the Mediterranean Sea: A Look at Habitat Changes, ed N. E. Stambler (New York, NY: Nova Science Publishers), 199–226.

Brooks, R. H., Brezonik, P. L., Putnam, H. D., and Keim, M. A. (1971). Nitrogen fixation in an estuarine environment: the Waccassassa on Florida Gulf coast. Limnol. Oceanogr. 16, 701–710. doi:10.4319/lo.1971.16.5.0701

Capone, D. G. (1983). N2 fixation in seagrass communities. Mar. Technol. Soc. J. 17, 32–37.

Coparoso, J. G., Kucyzinski, J., Stombaugh, J., Bittinger, K., Bushman, F. D., Costello, E. K., et al. (2010). QIIME allows analysis of high-throughput community sequencing data. Nat. Methods 7, 335–336. doi:10.1038/nmeth.f.1303

Carpenter, E. J., and Capone, D. (2008). “Nitrogen fixation in the marine environment,” in Nitrogen Fixation in the Marine Environment, eds D. Capone, A. D. Bronk, R. M. Mulholland, and E. J. Carpenter (San Diego, CA: Elsevier Academic Press), 141–198.

Chien, Y. T., and Zinder, S. H. (1996). Cloning, functional organization, transcript studies, and phylogenetic analysis of the complete nitrogenase structural genes (nifHDK2) and associated genes in the archaeon Methanococcus bakeri 227. J. Bacteriol. 178, 143–148.

Codispoti, L. A., Brandes, J. A., Christensen, J. P., Devol, A. H., Naqvi, S. W. A., Pael, H. W., et al. (2001). The oceanic fixed nitrogen and nitrous oxide budget: moving targets as we enter the anthropocene. Sci. Mar. 65, 85–105.

Davis, J., Kaiser, K., and Benner, R. (2009). Amino acid and amino sugar yields and compositions as indicators of dissolved organic matter diagenesis. Org. Geochem. 40, 343–352. doi: 10.1016/j.orggeochem.2008.12.003

Deutsch, C., Sarmiento, L. J., 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

Eichinger, M., Kooijman, S. A. L. M., Sempere, R., Lefevre, D., Gregori, G., Carriére, B., et al. (2009). Consumption and release of dissolved organic carbon by marine bacteria in a pulsed-substrate environment: from experiments to modeling. Aquat. Microb. Ecol. 56, 41–54. doi:10.3354/am10132

Engel, A., and Passow, U. (2001). Carbon and nitrogen content of transparent exopolymer particles (TEP) in relation to their Alcan蓝 dye adsorption. Mar. Ecol. Prog. Ser. 219, 1–10. doi: 10.3354/meps219001

Eugster, O., and Gruber, N. (2012). A probabilistic estimate of global marine N-fixation and denitrification. Global Biogeochem. Cycles 26. doi:10.1029/2012GB004300

Falkowski, P. G. (1997). Evolution of the nitrogen cycle and its influence on the biological sequestration of CO2 in the ocean, Nature 387, 272–275. doi:10.1038/387272a0

Fernandez, C., Farias, L., and Ulloa, O. (2011). Nitrogen fixation in denitrified marine waters. PLoS ONE 6:e20539. doi:10.1371/journal.pone.0020539

Flores, E., and Herrero, A. (2005). Nitrogen assimilation and nitrogen control in cyanobacteria. Biochem. Soc. Trans. 33, 164–167. doi:10.1042/BST0330164

Foster, R. A., Paytan, A., and Zehr, J. P. (2009). Seasonality of N2 fixation and nifH gene diversity in the Gulf of Aqaba (Red Sea). Limnol. Oceanogr. 54, 219–233. doi:10.4319/lo.2009.54.1.0219

Goldthwait, S. A., Carlson, C. A., Henderson, G. K., and Allredge, A. L. (2005). Effects of physical fragmentation on remineralization of marine snow. Mar. Ecol. Prog. Ser. 305, 59–65. doi:10.3334/meps305059

Großkopf, T., Mohr, W., Baustian, T., Schunck, H., Gill, D., Kuyper, M. 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. (2008). “The marine nitrogen cycle: overview of distributions and processes,” in Nitrogen in the Marine Environment, 2nd Edn., eds D. Capone, D. Bronk, M. Mulholland, and E. Carpenter (Amsterdam: Elsevier), 1–50.

Gruber, N., and Galloway, J. N. (2008). An Earth-system perspective of the global nitrogen cycle. Nature 451, 293–296. doi:10.1038/nature06592

Halm, H., Lam, P., Ferdelman, T. G., Lavik, G., Dittmar, T., Laroche, J., et al. (2012). Heterotrophic
organisms dominate nitrogen fixation in the South Pacific Gyre. ISME J. 6, 1238–1249. doi: 10.1038/ismej.2011.182

Hammersley, M. R., Turk, K. A., Leinweber, A., Gruber, N., Zehr, J. P., Gunderson, T., et al. (2011). Nitrogen fixation within the water column associated with two hypoxic basins in the Southern California Bight. Aquat. Microb. Ecol. 63, 193–205. doi: 10.3354/ame01494

Herbert, R. A. (1999). Nitrogen cycling in coastal marine ecosystems. FEMS Microbiol. Rev. 23, 563–590. doi: 10.1111/j.1574-6976.1999.tb00414.x

Hewson, I., and Fuhrman, J. A. (2007). Covariation of viral parameters with bacterial assemblage richness and diversity in the water column and sediments. Deep Sea Res. I. 54, 811–830. doi: 10.1016/j.dsr.2007.02.003

Howarth, R. W., Marino, R., Lane, J., and Cole, J. I. (1988). Nitrogen fixation in fresh-water, estuarine, and marine ecosystems. I. rates and importance. Limnol. Oceanogr. 33, 669–687. doi: 10.4319/lo.1988.33.3_part_2.0669

Ilbello, V., Cantoni, C., Cozzi, O., and Civitarese, G. (2010). First basin-wide experimental results on N2 fixation in the open Mediterranean Sea. Geophys. Res. Lett. 37. doi: 10.1029/2009GL041635

Jayakumar, A., Al-Rshaidat, M. M. D., Ward, B. B., and Mulholland, M. R. (2012). Diversity, distribution, and expression of diazotroph nifH genes in oxygen-deficient waters of the Arabian Sea. FEMS Microbiol. Ecol. 82, 597–606. doi: 10.1111/j.1574-6941.2012.01430.x

Karl, D., Michaels, A., Bergman, B., Capone, D., Carpenter, E., Letelier, R., et al. (2002). Dinitrogen fixation in the world’s oceans. Biogeochemistry 57, 47–98. doi: 10.1023/A:1015799105851

Knapp, A. N. (2012). The sensitivity of marine N2 fixation to dissolved inorganic nitrogen. Front. Microbiol. 3:374. doi: 10.3389/fmicb.2012.00374

Kress, N., and Herut, B. (2001). Spatial and seasonal evolution of dissolved oxygen and nutrients in the Southern Levantine Basin (Eastern Mediterranean Sea): chemical characterization of the water masses and inferences on the N: P ratios. Deep Sea Res. I. 48, 2347–2372. doi: 10.1016/S0967-0637(01)00022-X

Krom, M. D., Emeis, K. C., and Van Cappellen, P. (2010). Why is the Eastern Mediterranean phosphorus limited. Prog. Oceanogr. 85, 236–244. doi: 10.1016/j.pocean.2010.03.003

Langlois, R. J., LaRoche, J., and Raab, P. A. (2005). Diazotrophic diversity and distribution in the tropical and subtropical Atlantic Ocean. Appl. Environ. Microbiol. 71, 7910–7919. doi: 10.1128/AEM.71.12.7910-7919.2005

Man-Aharonovich, D., Kress, N., Bar-Zeev, E., Berman-Frank, I., and Beja, O. (2007). Molecular ecology of nifH genes and transcripts in the Eastern Mediterranean Sea. Environ. Microbiol. 9, 2354–2363. doi: 10.1111/j.1462-2920.2007.01353.x

McGlherty, K. J. (2008). “The marine nitrogen cycle: overview of distributions and processes,” in Nitrogen in the Marine Environment, 2nd Edn., eds D. Capone, D. Bronk, M. Mulholland, and E. Carpenter (Amsterdam: Elsevier), 1037–1071.

Mehta, M. P., and Baross, J. A. (2006). Nitrogen fixation at 92 degrees C by a hydrothermal vent archaeon. Science 314, 1783–1786. doi: 10.1126/science.1134772

Mehta, M. P., Huber, J. A., and Baross, J. A. (2005). Incidence of novel and potentially archaean nitrogenase genes in the deep Northeast Pacific Ocean. Environ. Microbiol. 7, 1525–1534. doi: 10.1111/j.1462-2920.2005.00836.x

Middelburg, J. J., and Nieuwenhuize, J. (2001). Nitrogen isotope tracer assay for N2 fixation. Limnol. Oceanogr. 46, 1001–1009. doi: 10.4319/lo.2001.46.2.01352

O’Neil, J. M., and Capone, D. G. (2008). “Nitrogen Cycling in Coral Reef Environments,” in Nitrogen in the Marine Environment, 2nd Edn., eds D. Capone, D. Bronk, M. Mulholland, and E. Carpenter. (Amsterdam: Elsevier), 949–989.

Orr, K. M., Lipchultz, F., Gundersen, K., Arimoto, R., Michaels, A. F., Knapp, A. H., et al. (2001). A seasonal study of the significance of N2 fixation by Trichodesmium spp. at the Bermuda Atlantic Time-series Study (BATS) site. Deep Sea Res. I. 48, 1583–1608.

Paerl, H., and Prufert, L. E. (1987). Organic nitrogen in estuarine and seasonal evolution of dissolved inorganic nitrogen. Appl. Environ. Microbiol. 53, 1080–1087.

Passow, U. (2002). Production of transparent exopolymer particles (TEP) by phyto- and bacterioplankton. Mar. Ecol. Progr. Ser. 236, 1–12. doi: 10.3354/meps23601

Passow, U., and Allredge, A. L. (1995). A dye-binding assay for the spectrophotometric measurement of transparent exopolymer particles (TEP). Limnol. Oceanogr. 40, 1326–1335. doi: 10.4319/lo.1995.40.7.1326

Payań, A., Mackey, K. R. M., Chen, Y., Lima, I. D., Doney, S. C., Mahowald, N., et al. (2009). Toxicity of atmospheric aerosols on marine phytoplankton. Proc. Natl. Acad. Sci. U.S.A. 106, 4601–4605. doi: 10.1073/pnas.0811486106

Post, A. E., Dedej, Z., Gottlieb, R., Li, H., Thomas, D. N., El-Absiawi, M., et al. (2002). Spatial and temporal distribution of Trichodesmium spp in the stratified Gulf of Aqaba, Red Sea. Mar. Ecol. Prog. Ser. 239, 241–250. doi: 10.3354/meps239241

Postgate, J. (1998). “Nitrogen fixation,” in Nitrogen Fixation, 3rd Edn., ed J. Postgate (London: Cambridge University Press), 124.

Rahav, E., Herut, B., Stambler, N., Bar-Zeev, E., Mulholland, M. R., and Berman-Frank, I. (2013a). Coupling between dinitrogen fixation and primary productivity in the Eastern Mediterranean Sea. J. Geophys. Res. 118, 1–8. doi: 10.1002/jgrc.20023

Rahav, E., Herut, B., Levi, A., Mulholland, M. R., and Berman-Frank, I. (2013b). Springtime contribution of dinitrogen fixation to primary production across the Mediterranean Sea. Ocean Sci. 9, 1–10. doi: 10.5194/os-9-489-2013

Rees, A. P., Gilbert, J. A., and Kelly-Gerreyn, B. A. (2009). Nitrogen fixation in the western English Channel (NE Atlantic Ocean). Mar. Ecol. Prog. Ser. 374, 7–12. doi: 10.3354/meps07771

Riemann, L., Farnelid, H., and Steward, G. E. (2010). Nitrogenase genes in non-cyanobacterial plankton: prevalence, diversity and regulation in marine waters. Aquat. Microb. Ecol. 61, 225–237. doi: 10.3354/meps09131

Simon, M., Allredge, A. L., and Azam, F. (1990). Bacterial carbon dynamics on marine snow. Mar. Ecol. Prog. Ser. 65, 205–211. doi: 10.3354/meps065205

Tamura, K., Dudley, J., Nei, M., and Kumar, S. (2007). MEGA4: molecular evolutionary genetics analysis (MEGA) software version 4.0. Mol. Biol. Evol. 24, 1596–1599. doi: 10.1093/molbev/msm092

Takahata, T., Thingstad, T. F., Christaki, U., Colombo, J., Cornet-Barthaux, V., Courties, C., et al. (2011). Lack of P-limitation of phytoplankton and heterotrophic prokaryotes in surface waters of three anticyclonic eddies in the stratified Mediterranean Sea. Biogeosciences 8, 525–538. doi: 10.5194/bg-8-525-2011
Voss, M., Croot, P., Lochte, K., Mills, M., and Peeken, I. (2004). Patterns of nitrogen fixation along 10N in the tropical Atlantic. *Geophys. Res. Lett.* 31. doi: 10.1029/2004gl020127

Ward, B. B., Devol, A. H., Rich, J. J., Chang, B. X., Bulow, S. E., Naik, H., et al. (2009). Denitrification as the dominant nitrogen loss process in the Arabian Sea. *Nature* 461, 78–81. doi: 10.1038/nature08276

Weiss, R. F. (1970). Solubility of nitrogen, oxygen and argon in water and seawater. *Deep-Sea Res.* 17, 721–735.

Wilson, S. T., Boettjer, D., Church, M. J., and Karl, D. M. (2012). Comparative assessment of nitrogen fixation methodologies, conducted in the oligotrophic North Pacific Ocean. *Appl. Environ. Microbiol.* 78, 6516–6523. doi: 10.1128/AEM.01146-12

Wu, J. F., Chung, S.-W., Wen, L.-S., Liu, K.-K., Chen, Y.-L., Chen, H.-Y., et al. (2003). Dissolved inorganic phosphorus, dissolved iron, and *Trichodesmium* in the oligotrophic South China Sea. *Global Biogeochem. Cycles* 17:1008. doi: 10.1029/2002GB001924

Zehr, J. P., and Ward, B. B. (2002). Nitrogen cycling in the ocean: new perspectives on processes and paradigms. *Appl. Environ. Microbiol.* 68, 1015–1024. doi: 10.1128/AEM.68.3.1015-1024.2002

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