<table>
<thead>
<tr>
<th>Title</th>
<th>Nitrogen biogeochemistry in the Caribbean sponge <em>Xestospongia muta</em>: A source or sink of dissolved inorganic nitrogen?</th>
</tr>
</thead>
<tbody>
<tr>
<td>Author(s)</td>
<td>Fiore, CL; Baker, DM; Lesser, MP</td>
</tr>
<tr>
<td>Citation</td>
<td>PLOS ONE, 2013, v. 8</td>
</tr>
<tr>
<td>Issued Date</td>
<td>2013</td>
</tr>
<tr>
<td>URL</td>
<td><a href="http://hdl.handle.net/10722/190367">http://hdl.handle.net/10722/190367</a></td>
</tr>
<tr>
<td>Rights</td>
<td>This work is licensed under a Creative Commons Attribution-NonCommercial-NoDerivatives 4.0 International License.</td>
</tr>
</tbody>
</table>
Nitrogen Biogeochemistry in the Caribbean Sponge, *Xestospongia muta*: A Source or Sink of Dissolved Inorganic Nitrogen?

Cara L. Fiore¹, David M. Baker², Michael P. Lesser¹*

¹ University of New Hampshire, Department of Molecular, Cellular and Biomedical Sciences, Durham, New Hampshire, United States of America, ² Geophysical Laboratory, Carnegie Institute of Washington, Washington, District of Columbia, United States of America

Abstract

**Background:** Sponges have long been known to be ecologically important members of the benthic fauna on coral reefs. Recently, it has been shown that sponges are also important contributors to the nitrogen biogeochemistry of coral reefs. The studies that have been done show that most sponges are net sources of dissolved inorganic nitrogen (DIN; NH₄⁺ and NO₃⁻) and that nitrification, mediated by their symbiotic prokaryotes, is the primary process involved in supplying DIN to adjacent reefs.

**Methodology/Principal Findings:** A natural experiment was conducted with the Caribbean sponge *Xestospongia muta* from three different locations (Florida Keys, USA; Lee Stocking Island, Bahamas and Little Cayman, Cayman Islands). The DIN fluxes of sponges were studied using nutrient analysis, stable isotope ratios, and isotope tracer experiments. Results showed that the fluxes of DIN were variable between locations and that *X. muta* can be either a source or sink of DIN. Stable isotope values of sponge and symbiotic bacterial fractions indicate that the prokaryotic community is capable of taking up both NH₄⁺ and NO₃⁻ while the differences in δ¹⁵N between the sponge and bacterial fractions from the NH₄⁺ tracer experiment suggest that there is translocation of labeled N from the symbiotic bacteria to the host.

**Conclusions/Significance:** Nitrogen cycling in *X. muta* appears to be more complex than previous studies have shown and our results suggest that anaerobic processes such as denitrification or anammox occur in these sponges in addition to aerobic nitrification. Furthermore, the metabolism of this sponge and its prokaryotic symbionts may have a significant impact on the nitrogen biogeochemistry on Caribbean coral reefs by releasing large amounts of DIN, including higher NH₄⁺ concentrations that previously reported.


Editor: Melanie R. Mormile, Missouri University of Science and Technology, United States of America

Received December 28, 2012; Accepted July 19, 2013; Published August 26, 2013

Copyright: © 2013 Fiore et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Funding: This project was funded by grants from the National Oceanic and Atmospheric Administration Ocean Exploration Program, Undersea Research Program, National Institute for Undersea Science and Technology as well as the National Science Foundation. The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

Competing Interests: The authors have declared that no competing interests exist.

* E-mail: mpl@unh.edu

Introduction

Sponges are an ecologically dominant component in many marine ecosystems, including coral reefs, where they contribute to the consolidation of reefs, prevent erosion, filter large quantities of seawater and provide habitat and food for many invertebrates and fishes [1–3]. Because of their ability to efficiently filter picoplankton, sponges can also contribute significantly to the coupling of productivity in the overlying water column to the benthos [1,4,5]. More recently, sponges and their prokaryotic symbionts have become an important area of research to quantify the fluxes of DIN by sponges and the biogeochemical cycling of nutrients on coral reefs [4,6–8].

Nitrogen cycling on tropical coral reefs is particularly important, as nitrogen is a limiting nutrient and the success of several coral reef taxa (e.g., corals) is dependent on their symbiotic partners and the efficient re-cycling of nitrogen between host and symbionts [9]. Compared to ambient seawater the excurrent water of actively pumping sponges is often enriched in DIN such as NO₃⁻ (or NO₂⁻ + NO₃⁻) as a result of nitrification [10]. This has been documented for sponges from coral reefs, mangroves, seagrass beds [7,8,11–13], as well as sponges from temperate and cold-water environments [14–16]. In fact, coral reef sponges have been documented to have rates of nitrification that are significantly higher (5.8–16.0 mmol m⁻² d⁻¹ NO₃⁻) [7,13] than what has been reported for benthic habitats such as microbial mats (up to 1.4 mmol m⁻² d⁻¹ NO₃⁻ [17]) or coral reef sediment (1.7 mmol m⁻² d⁻¹ NO₃⁻ [18]).

All pathways of nitrogen biogeochemistry have been reported to occur in sponges [10] including nitrogen fixation which was initially measured using the acetylene reduction method [19–21]. Stable isotope tracer studies (i.e., ¹⁵N₂) later confirmed the presence of nitrogen fixation, albeit at low rates, in several species of sponges from coral reefs [21,22]. Recently, the first sponge-derived *nfH* gene sequences and transcripts, which encode for the iron protein component of the nitrogenase enzyme responsible for
nitrogen fixation, were documented in two sponge species from the Florida Keys [23]. nifH genes have also been recovered from *Xestospongia muta* from multiple locations in the Caribbean [Fiore and Lesser unpublished] and hypoxic/anoxic conditions, known to occur in sponges [10], would be required for activity since nitrogenase is inactivated by molecular oxygen and only fixes nitrogen under anaerobic or micro-aerobic conditions [10]. With the presence of anaerobic microhabitats in sponges [10,24], other anaerobic nitrogen transformations including sulfate reduction, denitrification and anaerobic ammonium oxidation (anammox) have been observed and quantified using stable isotopic tracer methods, radiolabeled isotopes and recovery of gene specific sequences for key enzymes [16,24,25]. Interestingly, genomic analysis of the candidate phylum Poribacteria, which is found in sequences for key enzymes [16,24,25]. Interestingly, genomic analysis of the candidate phylum Poribacteria, which is found in sponges from numerous marine habitats [26,27], suggests that Poribacteria may also be capable of denitrification [29].

Nitrification in sponges, which produces the bulk of the DIN released [8,13], has been documented using collection and incubation methods combined with nutrient analyses [12,13,16]. Southwell et al. [7,8] was the first to use an *in situ* method to identify actively nitrifying sponges and estimate the flux of DIN onto the adjacent coral reef, and found no significant difference in flux of DIN between the incubation and *in situ* methods. The interest in nutrient fluxes mediated by sponges and their symbionts as well as the nutrient biogeochemistry of coral reefs has resulted in a surge of research into the prokaryotic community composition of sponges and the processes they mediate. Recently, this has included the use of high throughput sequencing methods, such as 454 pyrosequencing of the 16S rRNA gene [27,29,30,Fiore and Lesser unpublished] which have increased our understanding of the prokaryotic composition of many sponge species from different marine habitats. This genetic information has provided useful complementary insight for characterizing prokaryotic mediated nutrient cycling in these sponges.

On Caribbean coral reefs *Xestospongia muta* is an ecologically dominant member of the benthic community, and on Conch Reef, FL (USA) the number of *X. muta* has been shown to be significantly increasing over time [31]. *Xestospongia muta* is also characterized as a high microbial abundance sponge [32] but little was known about the composition of this community other than it contained Cyanobacteria [33] until recent studies documented a diverse prokaryotic community in *Xestospongia muta* and other members of this genus [34,35,Fiore and Lesser unpublished]. Additionally, *X. muta* outside of the Florida Keys have not been as well studied generally with quantitative 454 pyrosequencing comparing the prokaryotic symbionts of *X. muta* from the Florida Keys, Cayman Islands and Bahamas having been recently completed [Fiore and Lesser unpublished].

The primary goal of this study was to quantify DIN fluxes in *X. muta* from the same three populations where our 16S rRNA 454 pyrosequencing study was done. We ask whether sponges from these same populations in the Caribbean have different fluxes of DIN, and potentially how any differences in DIN fluxes, may be related to the taxonomy of their symbiotic prokaryotes using a comparative approach and a natural experiment [36].

**Materials and Methods**

**Sample Locations**

Replicate sponges (*n* = 6) were sampled at approximately 15 m depth between 9 and 10 AM and again between 4 and 5 PM when indicated from each of three locations: Rock Bottom Reef, Little Cayman, Cayman Islands (LC) (19°42’7.36" N, 80°3’24.94" W), North Perry Reef, Lee Stocking Island, Bahamas (LSI) (23°47’0.03" N, 76°6’5.14" W), and Conch Reef, Key Largo, FL (FL) (24°57’0.03" N, 80°27’11.16" W). All populations were sampled during the late spring and early summer of 2011 where the maximum irradiance of photosynthetically active radiation (PAR; 400–700 nm) at noon for all three locations is ~500–600 μmol quanta m⁻² s⁻¹ [Lesser unpublished]. Necessary permits were obtained for all three locations: the Marine Conservation Board, Cayman Islands; Department of Marine Resources, Bahamas; NOAA ONMS permit number FKNMS-2011-066 for Conch Reef, Florida Keys.

**Nutrient analyses and rates of sponge pumping**

Ambient and excurrent water samples for nutrient analysis were collected from individual sponges (*n* = 6) at each location for nutrient analysis by slowly filling 100 ml syringes and placing all water samples on ice for transport to shore. Ambient water was obtained by filling the syringe adjacent to each sponge (within 20 cm of the sponge body wall) and excurrent water was obtained by placing weighted Tygon tubing inside the sponge close to the base of the spongocoel that was attached to a 100 ml syringe and drawing water into the syringe slowly (~1 ml s⁻¹). Syringes were then purged of approximately 10 ml and then 40 ml was saved and frozen for NH₄⁺ and NO₃⁻ and NO₂⁻ analysis (NO₃⁻). For the nutrient analyses the water samples were thawed and filtered (0.22 μm, Whatman, USA) to remove particulate matter then re-frozen and sent to the Nutrient Analytical Facility at Woods Hole Oceanographic Institute (WHOI, Woods Hole, MA, USA) for analysis using a Lachat QuickChem 8000 (flow injection analysis system) according to standard protocols to determine concentration of NH₄⁺ and NO₃⁻. Instrumental errors associated with the measurements were calculated as relative standard deviation (RSD) and includes: NH₄⁺ ~0.6% measured RSD, and NO₃⁻ ~0.59% measured RSD. Sponges were marked near their base with labeled flagging tape attached to nails embedded in the substrate to facilitate repeated measurements on the same sponges.

The volume flow or pumping rates for each individual sponge was determined as previously described [37]. A small amount (~1 ml) of fluorescein dye was injected using a syringe and 16 gauge needle into the sponge just below the base of the spongocoel and the time(s) that the dye front took from its first appearance at the base of the spongocoel to the top of the spongocoel was recorded to obtain the centerline fluid velocity to calculate volume flux or pumping rate. We understand that unlike previous studies on tubular sponges where plug flow can be reasonably assumed (e.g., [37]) the morphology of *X. muta* likely creates more complicated excurrent plumes where the velocity across the osculum is not uniform [38]. This is easily observed using the timing of multiple dye tracks on *X. muta* injected in different locations with dye tracks closer to the sponge wall being slower than the centerline flow [Lesser unpublished]. As a result we recognize that our measurements of volume flow or pumping rates are likely to be an overestimate. That said our estimates of volume flow are in agreement with the results of Southwell et al. [8] using similar techniques for *X. muta*. Additionally, in our hands we have never observed cessation of pumping, or other artifacts, as a result of exposure to fluorescein in both thin walled and thick walled sponges [5,37]. Both spongocoel and total sponge volume were calculated by measuring sponge height, base circumference, osculum diameter, and spongocoel depth and inner diameter with a measuring tape (±1.0 mm) and volume calculated as previously described [31]. The mass (kg) of individual sponges was then calculated by multiplying the individual total sponge volume (l), obtained as described above, by the average density of *X. muta*
sponges (0.617 g cm\(^{-3}\)) which was determined from direct measurements of the displacement volume and mass of pieces (\(n = 5\)) of sponge (including both mesohyl and pinacoderm). The flux of nutrients was then calculated by multiplying the \(\Delta\text{DIN}\) (the difference in nutrient concentration between the ambient and excurrent water in \(\mu\text{mol l}^{-1}\) by the flow rate (\(\text{cm s}^{-1}\)) and normalized to both sponge volume and mass for comparisons between sites.

Pumping rates and nutrient flux data were tested for assumptions of ANOVA and if the data failed either normality or homoscedasticity a constant integer to all values was added following by log transformation. The transformed data passed Bartlett’s test [39] for homoscedasticity but often failed the Shapiro-Wilks [40] test for normality. Because Bartlett’s test is sensitive to deviations in normality [41] we choose to proceed with ANOVA, which is known to be robust to deviations from normality [42], on the transformed data. To determine if time of day was a significant factor in the flux of DIN, a two-way ANOVA with interaction was performed using the statistical program R [43] with time (AM and PM) and location as fixed factors for the flux of NO\(_x\), NH\(_4\), total DIN and pumping rate as the response variables. A repeated measures ANOVA was not performed because the general requirement of this approach is three time points. Since the effect of time and interaction of time with location was not significant the flux of NO\(_x\), NH\(_4\), total DIN and pumping rate, \(\Delta\text{DIN}\) and pumping rates for each location were calculated by averaging the AM and PM values for each individual sponge. Collapsing the design to a single factor analysis to examine differences between locations was then assessed using a one-way ANOVA with location as a fixed factor [41].

Flow Cytometry

Ambient and excurrent water samples were collected as described above for the nutrient analyses for another set of sponges (\(n = 4\)) from LSI only. Approximately 3 ml from each collected water sample were fixed in electron microscopy grade paraformaldehyde at a final concentration of 0.5% in filtered (0.22 \(\mu\text{m}\) seawater and frozen at \(-50^\circ\text{C}\). Frozen water samples were sent to the Bigelow Laboratory for Ocean Sciences J.J. MacIsaac Aquatic Cytometry Facility where they were stored in liquid nitrogen until analysis. Each sample was analyzed for cell abundances using a Becton Dickinson FACScan flow cytometer with a 30 mW, 488 nm laser. Simultaneous measurements of forward light scattering (FSC, relative size), 90\(^\circ\) light scatter (SSC), chlorophyll fluorescence (>650 nm), and phycoerythrin fluorescence (560–590 nm) were made simultaneously on each sample as previously described [5]. Calculations of cyanobacteria, prochlorophytes, and heterotrophic cell concentration and filtering efficiency were performed as previously described [5]. Technical replicates (\(n = 2\)) were averaged for each sample and the cell abundance of heterotrophic bacteria was determined using PicoGreen (Molecular Probes), a dsDNA specific dye, which stains all prokaryotes (emission fluorescence 515–525 nm). Subtraction of the chl \(a\) containing picoplankton from the total prokaryotes yielded the heterotrophic bacterial component of the community while cyanobacterial and prochlorophyte cells were differentiated by the presence or absence, respectively, of phycoerythrin fluorescence. All filtered cells were converted to carbon and nitrogen equivalents using the following conversions; heterotrophic bacteria: 20 fg C cell\(^{-1}\) [44], Prochlorococcus: 53 fg C cell\(^{-1}\) [45], Synechococcus: 470 fg C cell\(^{-1}\) [46], heterotrophic bacteria: 3.3 fg N cell\(^{-1}\) [47], Prochlorococcus: 9.4 fg N cell\(^{-1}\) [48], Synechococcus: 55 fg N cell\(^{-1}\) [48]. Data were log transformed or arcsin transformed as necessary and an ANOVA followed by Tukey’s HSD were performed to test for significant differences in the number of filtered cells between cell types (cyanobacteria, prochlorophytes, heterotrophic bacteria and total cells), filtration efficiency and total particulate carbon (POC) and nitrogen (PON) consumed by sponges.

Stable isotopic analyses and tracer experiments

Sponge samples that were frozen without buffer (\(n = 3\) each location) were later lyophilized, ground to a powder with a mortar and pestle, and then acid treated with 1 M HCl to remove carbonate and rinsed with distilled water and allowed to dry. An analysis of samples from FL, separated into the outer pigmented layers of the sponge and the non-pigmented inner tissues (containing the pinacoderm and outer mesohyl respectively), showed no significant differences in stable isotope signatures [Fiore and Lesser, unpublished data] so whole cross-sections of sponge samples, consisting of both pinacoderm and mesohyl, from all locations were analyzed. Samples were then sent to the Marine Biological Laboratory (MBL) for the analysis of particulate C and N as well as the natural abundance of the stable isotopes \(\delta^{13}\text{C}\) and \(\delta^{15}\text{N}\). Samples were analyzed using a Europa ANCA-SL elemental analyzer-gas chromatograph attached to a continuous-flow Euro- pa 20–20 gas source stable isotope ratio mass spectrometer. The carbon isotope results are reported relative to Vienna Pee Dee Belemnite and nitrogen isotope results are reported relative to atmospheric air and both are expressed using the delta (\(\delta\)) notation in parts per mil (%). The analytical precision of the instrument is \(\pm 0.1\%\), and the mean precision of sample replicates for \(\delta^{13}\text{C}\) was \(\pm 0.4\%\) and \(\delta^{15}\text{N}\) was \(\pm 0.2\%\). A one-way ANOVA was used to test for significant differences between locations for \(\delta^{13}\text{C}, \delta^{15}\text{N}\) and C:N ratios followed by the post hoc multiple comparison Tukey’s HSD test as needed.

Two stable isotope tracer experiments were conducted during the summer of 2011 at LSI: the first used Na\(_{15}\text{NO}_3 (5 \text{mg l}^{-1} \text{final concentration}) plus H\(_{13}\text{CO}_3 (50 \text{mg l}^{-1})\) and the second used NH\(_4\) \((0.31 \text{mg l}^{-1})\) as tracers (Sigma-Aldrich, USA). The method was the same for each experiment: 11 individual X. muta (average volume 172±77 ml or mass 0.106±0.048 kg; mean ±SD) were collected by cutting through the bottom of the sponge but keeping the tissue from the base of the spongocoel intact, from approximately 12 m at North and South Perry reefs at LSI and held in a large holding tank with flow through seawater for 5 d to recover from being removed from the reef. Care was taken to ensure that sponges were never exposed to air and that light levels were maintained at the same levels as found at \(-12\) m using neutral density screens over the outdoor flowing seawater tanks. Sponges were checked for pumping activity using fluorescein dye and incubated statically with the tracer compound(s) for 4 h. Subsequently, T\(_0\) sponges (\(n = 3\)) were then removed and stored for analysis.

The remaining sponges were placed in individual aquaria with flow through seawater and sponges were sampled at 3 h (\(n = 2\)), 6 h (\(n = 3\)) and at 12 h (\(n = 3\)) for each fraction. Frozen samples were initially processed by separating the bacteria and sponge fractions following the methods of Freeman and Thacker [49] and Freeman et al. [50] except for two steps: an initial centrifugation was performed at 520\(\times\) g for 4 min, and the resulting sponge pellet was rinsed an additional two times. The purity of the sponge and bacterial fractions were assessed using light and epifluorescence light microscopy as described by Freeman and Thacker [49]. The sponge fractions always contained large cells (8–10 \(\mu\text{m}\) diameter) consisting of at least 85% per microscopic field and exhibiting low natural fluorescence, whereas the bacterial fractions

---

PLOS ONE | www.plosone.org 3 August 2013 | Volume 8 | Issue 8 | e72961

Nitrogen Biogeochemistry in a Caribbean Sponge

---

[41] Freeman et al. [50] except for two steps: an initial centrifugation was performed at 520\(\times\) g for 4 min, and the resulting sponge pellet was rinsed an additional two times. The purity of the sponge and bacterial fractions were assessed using light and epifluorescence light microscopy as described by Freeman and Thacker [49]. The sponge fractions always contained large cells (8–10 \(\mu\text{m}\) diameter) consisting of at least 85% per microscopic field and exhibiting low natural fluorescence, whereas the bacterial fractions...
contained only small cells (<1–2 μm diameter) and high natural fluorescence. While efforts were made to separate and purify the sponge fraction as much as possible from all prokaryotic cells, it is possible that some prokaryotes that were located intracellularly were not detected (non-fluorescent) in the sponge fraction. These methods have been shown to be effective for other sponge species [49] and were optimized for use with *X. muta*. Additionally, a one-way ANOVA of the C:N ratios for the two fractions yielded significant differences (F1,13 = 3.38, p = 0.01) indicating that good separation of these fractions occurred. It is likely, however, that some contamination occurred and was considered when interpreting the results of these experiments. Samples were then lyophilized and ~1.0 mg was weighed and placed into silver capsules (Costech, CA, USA) and acidified three times with 20 μl of 12 M HCl. Samples were allowed to dry in between acidifications, then oven dried at 50°C for 48 h. Samples were combusted in a Carlo-Erba NC2500 elemental analyzer, and the analytical precision of the instrument was ±0.2%, and the mean standard deviation of sample replicates for δ¹³C was ±0.4% and for δ¹⁵N it was ±0.8% for enriched samples and ±0.1% and ±0.1% for natural abundance samples, respectively. For the tracer experiments the data were log transformed as necessary to meet the assumptions of parametric statistics and a two-way ANOVA with interaction, with fraction and time as fixed factors, was used to assess treatment effects.

Results

Stable Isotopic Signatures of Sponges

The values of δ¹³C from each location were not significantly different from each other (ANOVA, F2,6 = 1.16, p = 0.38). The δ¹³C of sponge samples ranged from −19.1 to −18.4% (Fig. 1). The δ¹⁵N of sponge samples ranged from 4.0–4.4% (Fig. 1), and were not significantly different between locations (ANOVA, F2,6 = 0.52, p = 0.62). The ratios of C:N were significantly different between locations (ANOVA, F2,6 = 22.57, p = 0.002), with post hoc pairwise comparisons showing that FL sponges had significantly higher C:N ratios than LSI (Tukey’s HSD, p<0.05), and that LC sponges significantly higher than LSI (Tukey’s HSD, p<0.05). There was no significant difference between LC and FL (Tukey’s HSD, p>0.05). Despite the significant results for C:N ratios the mean values did not vary greatly, with a range of 4.33–4.83.

Inorganic Nitrogen Fluxes in *Xestospongia muta*

The difference in nutrient concentration between the ambient and excurrent of NH₄⁺, NO₂⁻ + NO₃⁻ (NOₓ⁻) and total DIN for *X. muta* varied considerably between individual sponges as expected for sponges over a large size range (Table 1). The ∆NH₄⁺ values were not significantly different between locations (ANOVA, F2,13 = 3.19, p = 0.07) as were the ∆NO₂⁻ values (ANOVA, F2,13 = 2.50, p = 0.11). ∆DIN values, however, were significantly different between locations (ANOVA, F2,13 = 6.82, p = 0.008) with post hoc pairwise comparisons showing that FL sponges were significantly lower than both LSI and LC sponges LSI (Tukey’s HSD, p<0.05) which were not significantly different than each other (Tukey’s HSD, p>0.05). No measurements of ambient NO₃⁻ exceeded 4 μM eliminating the potential for ambient nutrient concentrations to be confounded by oceanographic features such as internal waves [51,52]. Sponge pumping rates varied with size (Table 1) and did not differ significantly with location (ANOVA, F2,13 = 1.61, p = 0.23).

The volume and mass normalized fluxes of NH₄⁺ (Table 1, Fig. 2a) were not significantly different between locations (ANOVA, F2,13 = 0.45, p = 0.65 (volume); F2,13 = 0.85, p = 0.45 (mass)). The fluxes of NO₂⁻ normalized to sponge volume (Table 1, Fig. 2b) were significantly different between locations (ANOVA, F2,13 = 4.09, p = 0.02) with FL sponges significantly lower than LSI and LC not significantly different than either FL or LSI (Fig. 2A) but when normalized to mass did not show a significant effect of location (ANOVA, F2,13 = 3.56, p = 0.054). The flux of total DIN (NO₂⁻ + NH₄⁺) normalized to volume and to mass were not significantly different among locations (ANOVA, F2,13 = 1.24, p = 0.32 (volume); F2,13 = 2.09, p = 0.16 (mass)).

Feeding Study

*Xestospongia muta* (n = 4) from LSI instantaneously filtered an average of 1.5×10⁷ cells ml⁻¹ and there was a significant effect of cell type (ANOVA, F3,12 = 4.45, p = 0.03) (Fig. 3A). The number of both total cells and heterotrophic bacteria filtered was significantly higher than that of prochlorophytes (Tukey’s HSD, p<0.05) but not cyanobacteria (Tukey’s HSD, p>0.05), while the number of cyanoabacterial cells filtered was indistinguishable (Tukey’s HSD, p>0.05) from the prochlorophyte or total cell and heterotrophic cell groupings (Fig. 3A). For the filtration efficiency of each cell type there were no significant differences (ANOVA, F3,12 = 0.45, p = 0.72) (Fig. 3B). The total amount of PON for each retained cell type was greatest for cyanoabacteria, but there was no significant difference between cell types (ANOVA, F3,12 = 2.77, p = 0.09). Differences between the amount of PON for each retained cell type was significant (ANOVA, F3,12 = 4.68, p = 0.02) and greatest for total cells and heterotrophic bacteria compared to prochlorophytes (Tukey’s HSD, p>0.05) but not cyanobacteria (Tukey’s HSD, p>0.05), while the PON of cyanoabacterial cells was indistinguishable (Tukey’s HSD, p>0.05) from the prochlorophyte or total cell and heterotrophic cell groupings (Fig. 3C).

Nitrogen tracer experiment: Nitrate and Bicarbonate

While sponge and bacterial fractions became more enriched from 3 to 6 hours there was no significant effect of enrichment of ¹⁵N from the NO₃⁻ tracer in those fractions (ANOVA, F2,14 = 0.84, p = 0.57). There was also no significant difference between sponge and bacterial fractions or over time or the interaction of fraction and time (fraction, F1,3 = 0.89, p = 0.36; time, F1,3 = 1.16, p = 0.36; interaction term, F1,3 = 0.28, p = 0.84) (Fig. 4A). Additionally, the enrichment of ¹³C from the bicarbonate tracer experiment was significant (ANOVA, F1,3 = 20.9, p<0.0001) with fraction being non-significant (F1,3 = 0.13, p = 0.13) and time being significant (F1,3 = 44.3, p<0.001) with a non-significant interaction term, (F1,3 = 3.2, p = 0.056) (Fig. 4B). As a result post-hoc multiple comparison tests were only performed for time. Both the sponge and bacterial fractions became more enriched in ¹³C the sponge fraction over time with all sampling periods being significantly different than T₀ (Tukey’s HSD p<0.05) and not significantly different (Tukey’s HSD p>0.05) from each other (Fig. 4B).

Nitrogen tracer experiment: Ammonium

There was significant enrichment of ¹⁵N from the NH₄⁺ tracer in the experimental sponges (ANOVA, F7,13 = 3.38, p = 0.03) with the effects tests for fraction (F1,3 = 6.4, p = 0.03) and time (F1,3 = 5.0, p = 0.02) being significant and the interaction term non-significant (F1,3 = 1.8, p = 0.19) (Fig. 4C). Post-hoc multiple comparison testing for time revealed a significant (Tukey’s HSD
Figure 1. δ¹⁵N and δ¹³C values (mean ± SE) for Xestospongia muta (n = 3) for each location. FL = Florida Keys LC = Little Cayman, LSI = Lee Stocking Island, Bahamas. doi:10.1371/journal.pone.0072961.g001

p < 0.05 increase in enrichment over time with the bacterial fraction exhibiting greater enrichment (Fig. 4C).

Discussion

This study of the inorganic nitrogen fluxes in Xestospongia muta is the first to show differences in net fluxes of DIN both within and between populations of this ecologically dominant sponge on Caribbean coral reefs. Xestospongia muta densities are increasing [31], so quantifying both the fluxes of DIN and understanding the underlying processes driving the nitrogen biogeochemistry in this sponge is important for understanding the DIN availability on reefs in the Caribbean.

In a survey of nitrification in sponges on Conch Reef, Florida Southwell et al. [7,8] reported evidence of nitrification in nine out of twelve sponge species, including X. muta. The rates of nitrification measured in these sponges varied, but overall they were at least two orders of magnitude higher than other habitats (e.g., benthos, coral rubble). In the current study, we observed similar rates of X. muta pumping activity reported for Conch Reef sponges by Southwell et al. [8]. However, unlike Southwell et al. [7,8], X. muta from Conch Reef exhibited a negative flux of NO₃⁻, indicating that either denitrification or anammox processes were taking place or possibly dissimilatory nitrate reduction. Our results from LSI and LC are consistent with Southwell et al. [7,8] and for LSI the fluxes of NO₃⁻ are significantly higher than fluxes of NO₂⁻ from FL compared to Southwell et al. [8].

Xestospongia muta were actively pumping for all measurements taken during this study, which was not significantly different over time of day or between locations. While pumping rates were not significantly different, the observed variability in the unidirectional pumping of sponges has the potential to create microhabitats where both anaerobic nitrogen transformations (e.g., denitrification) and aerobic nitrogen transformations (e.g., nitrification) could occur [10,16,24,53]. Additionally, recent studies of the bacterial communities of X. muta using 16S rRNA sequencing [35, Fiore and Lesser unpublished] have reported many bacterial groups that are capable of denitrification and anammox (i.e., Burkholderiales, Pseudoalteromonadaceae, Poribacteria, Planctomycetes).

Interestingly, Southwell et al. [8] found that NO₃⁻ made up the majority of the DIN pool from X. muta and that NO₃⁻ was almost entirely NO₂⁻. In this study we observed, in addition to positive net NO₃⁻ fluxes, a greater net efflux of NH₄⁺ for all samples of X. muta. For some X. muta populations (i.e., LSI and LC) the flux of NH₄⁺ had a significant impact on total DIN fluxes. These differences in the fluxes of NH₄⁺, probably generated from the utilization of nitrogen rich POM by the sponge host, are unusual given there is an active nitrifying community [7,8], and a prokaryotic photosynthetic community [22] that could readily utilize NH₄⁺ in this sponge [54].

The fluxes of DIN from sponges such as X. muta can have a significant impact on the availability and composition of DIN on coral reefs [7,8]. Results of the current study indicate that fluxes of DIN from X. muta, the primary contributor to DIN on Caribbean coral reefs [7,8], is more complex than previously thought and is significantly different between locations. Additionally, these fluxes vary over time as a previous study of the same population of X. muta from LSI in 2010 showed both positive and negative fluxes of NO₃⁻ with a net ΔNO₃⁻ of −0.27 μM ± 0.06 (mean ± SE) [Fiore and Lesser unpublished].

Natural abundance stable isotope values have been commonly used to trace sources of C and N through the food chain [55]. Nitrogen fixation yields an average δ¹⁵N signature of approximately 0.0% [36,37], and trophic enrichment typically results in a +2.2% to +3.5% increase per trophic level for δ¹⁵N [58,59]. Therefore, several studies have used a cutoff of ≤2.0% to indicate N from a fixed source [23,60,61]. Additionally, carbon fixation by marine phytoplankton typically results in δ¹³C values of about −19 to −24% [55], with an average of +0.5 to +1.0% enrichment.
<table>
<thead>
<tr>
<th>Location</th>
<th>Spongocoel (L)</th>
<th>Volume (L)</th>
<th>Mass (kg)</th>
<th>Flow rate (\text{L h}^{-1})</th>
<th>Flux NH4+ (\text{mmol h}^{-1} \text{L}^{-1})</th>
<th>Flux NOx (\text{mmol h}^{-1} \text{L}^{-1})</th>
<th>Flux DIN (\text{mmol h}^{-1} \text{L}^{-1})</th>
<th>Flux NH4+ (\text{mmol h}^{-1} \text{kg}^{-1})</th>
<th>Flux NOx (\text{mmol h}^{-1} \text{kg}^{-1})</th>
<th>Flux DIN (\text{mmol h}^{-1} \text{kg}^{-1})</th>
</tr>
</thead>
<tbody>
<tr>
<td>FL 4</td>
<td>26.8</td>
<td>43</td>
<td>0.15</td>
<td>-0.18</td>
<td>0.26</td>
<td>-0.17</td>
<td>1.17</td>
<td>-0.31</td>
<td>0.65</td>
<td>0.06</td>
</tr>
<tr>
<td>FL 20</td>
<td>101</td>
<td>62.2</td>
<td>0.00</td>
<td>0.65</td>
<td>0.00</td>
<td>0.68</td>
<td>0.94</td>
<td>0.00</td>
<td>0.65</td>
<td>0.00</td>
</tr>
<tr>
<td>FL 39</td>
<td>36</td>
<td>22.4</td>
<td>0.35</td>
<td>-0.30</td>
<td>0.30</td>
<td>1.22</td>
<td>1.22</td>
<td>0.00</td>
<td>0.65</td>
<td>0.00</td>
</tr>
<tr>
<td>FL 33</td>
<td>111</td>
<td>69.2</td>
<td>0.10</td>
<td>-0.09</td>
<td>0.26</td>
<td>-0.17</td>
<td>0.94</td>
<td>0.00</td>
<td>0.65</td>
<td>0.00</td>
</tr>
<tr>
<td>FL 14</td>
<td>47.0</td>
<td>47.0</td>
<td>0.00</td>
<td>0.00</td>
<td>0.26</td>
<td>-0.17</td>
<td>0.94</td>
<td>0.00</td>
<td>0.65</td>
<td>0.00</td>
</tr>
<tr>
<td>FL 7</td>
<td>37</td>
<td>22.9</td>
<td>0.00</td>
<td>0.00</td>
<td>0.26</td>
<td>-0.17</td>
<td>0.94</td>
<td>0.00</td>
<td>0.65</td>
<td>0.00</td>
</tr>
<tr>
<td>LC 12</td>
<td>64</td>
<td>39.3</td>
<td>0.00</td>
<td>0.00</td>
<td>0.26</td>
<td>-0.17</td>
<td>0.94</td>
<td>0.00</td>
<td>0.65</td>
<td>0.00</td>
</tr>
<tr>
<td>LC 5</td>
<td>12</td>
<td>74</td>
<td>0.00</td>
<td>0.00</td>
<td>0.26</td>
<td>-0.17</td>
<td>0.94</td>
<td>0.00</td>
<td>0.65</td>
<td>0.00</td>
</tr>
<tr>
<td>LC 4</td>
<td>5</td>
<td>7.4</td>
<td>0.00</td>
<td>0.00</td>
<td>0.26</td>
<td>-0.17</td>
<td>0.94</td>
<td>0.00</td>
<td>0.65</td>
<td>0.00</td>
</tr>
<tr>
<td>LC 6</td>
<td>5</td>
<td>129</td>
<td>0.00</td>
<td>0.00</td>
<td>0.26</td>
<td>-0.17</td>
<td>0.94</td>
<td>0.00</td>
<td>0.65</td>
<td>0.00</td>
</tr>
<tr>
<td>LC 14</td>
<td>4</td>
<td>49</td>
<td>0.00</td>
<td>0.00</td>
<td>0.26</td>
<td>-0.17</td>
<td>0.94</td>
<td>0.00</td>
<td>0.65</td>
<td>0.00</td>
</tr>
<tr>
<td>LC 10</td>
<td>5</td>
<td>13.6</td>
<td>0.36</td>
<td>1.33</td>
<td>0.36</td>
<td>0.36</td>
<td>0.36</td>
<td>0.36</td>
<td>0.36</td>
<td>0.36</td>
</tr>
<tr>
<td>LC 29</td>
<td>81</td>
<td>90.2</td>
<td>0.50</td>
<td>0.63</td>
<td>0.63</td>
<td>0.63</td>
<td>0.63</td>
<td>0.63</td>
<td>0.63</td>
<td>0.63</td>
</tr>
<tr>
<td>LC 2</td>
<td>2</td>
<td>8.6</td>
<td>0.00</td>
<td>0.00</td>
<td>0.26</td>
<td>-0.17</td>
<td>0.94</td>
<td>0.00</td>
<td>0.65</td>
<td>0.00</td>
</tr>
<tr>
<td>LS 3</td>
<td>23</td>
<td>14.0</td>
<td>0.00</td>
<td>0.00</td>
<td>0.26</td>
<td>-0.17</td>
<td>0.94</td>
<td>0.00</td>
<td>0.65</td>
<td>0.00</td>
</tr>
<tr>
<td>LS 10</td>
<td>5</td>
<td>13.6</td>
<td>0.36</td>
<td>1.33</td>
<td>0.36</td>
<td>0.36</td>
<td>0.36</td>
<td>0.36</td>
<td>0.36</td>
<td>0.36</td>
</tr>
<tr>
<td>LS 29</td>
<td>81</td>
<td>90.2</td>
<td>0.50</td>
<td>0.63</td>
<td>0.63</td>
<td>0.63</td>
<td>0.63</td>
<td>0.63</td>
<td>0.63</td>
<td>0.63</td>
</tr>
<tr>
<td>LS 2</td>
<td>2</td>
<td>8.6</td>
<td>0.00</td>
<td>0.00</td>
<td>0.26</td>
<td>-0.17</td>
<td>0.94</td>
<td>0.00</td>
<td>0.65</td>
<td>0.00</td>
</tr>
</tbody>
</table>
Figure 2. The average flux (mean ± SE) of NH$_4^+$, NO$_3^-$ and DIN. The average flux is shown for each location normalized to sponge volume (A) and mass (B). Treatment groups with similar superscripts are not statistically different from each other. doi:10.1371/journal.pone.0072961.g002

Figure 3. Filtration of bacterioplankton from the water column by X. muta. Number of filtered cells (A), filtration efficiency (B), and particulate organic matter as carbon and nitrogen (C) available from filtered cells for X. muta from LSI (n = 4). Treatment groups (mean ± SE) with similar superscripts are not statistically different from each other. doi:10.1371/journal.pone.0072961.g003
Based on previous studies that have used stable isotope analysis to investigate the relationship between sponges and their symbionts [6, 7, 49], a cutoff for $\delta^{13}C$ of $-18\%$ or lower was used as an indication of photoautotrophic carbon fixation for X. muta. The bulk stable isotopic values measured for both C and N in X. muta tissue, comprising both the host tissue and prokaryotic biomass, were not significantly different between sites and similar to those documented in previous studies on X. muta [22, 23].

Using the cutoff values described above, there is no stable isotopic evidence that nitrogen fixation was occurring in X. muta (Fig. 1) although nifH genes have been sequenced from X. muta [Fiore and Lesser unpublished]. The $\delta^{13}C$ values of the sponge samples show evidence of photoautotrophy, although we cannot say definitively that there is transfer of C from symbionts to the sponge. Freeman and Thacker [49] demonstrated that high microbial abundance (HMA) sponges, such as X. muta, can obtain either C or N, or both, from their symbionts. Any interpretation of tissue stable isotopic signatures must, however, include heterotrophy on picoplankton for sponges. Based on the feeding study of LSI sponges, and despite the high abundance of resident bacteria, sponges are actively and non-selectively filtering most of the bacteria from the ambient water, which would supply significant amounts of POC and PON that could be potentially used by the host which is consistent with studies on other sponge species from the Caribbean [5, 37]. Taken together, the results of this study suggest some site related differences with sponges from the more open ocean sites of LSI and LC being more dependent on photoautotrophic sources of C and coastal FL sponges being more dependent on POM for their C requirements (Fig. 1).

The tracer experiments provide additional insight into what may be occurring in terms of the dynamics of C and N uptake by X. muta. $\delta^{15}N$ over time (A) for $^{15}N$ nitrate and $\delta^{13}C$ over time (B) for $^{13}C$ bicarbonate enriched sponge and bacterial fractions, and $\delta^{15}N$ over time for $^{15}N$ ammonium enriched sponge and bacterial fractions (C). Samples collected at 3 h were under low irradiances while samples collected at 6 h had been exposed to sunlight (Under neutral density screens representative of irradiances at the depth of collection) and samples from 12 h were collected at night (for both experiments). Treatment groups (mean ± SE) with similar superscripts are not statistically different from each other.

doi:10.1371/journal.pone.0072961.g004

Figure 4. $\delta^{15}N$ and $\delta^{13}C$ isotopic values from uptake experiments with X. muta. $\delta^{15}N$ over time (A) for $^{15}N$ nitrate and $\delta^{13}C$ over time (B) for $^{13}C$ bicarbonate enriched sponge and bacterial fractions, and $\delta^{15}N$ over time for $^{15}N$ ammonium enriched sponge and bacterial fractions (C). Samples collected at 3 h were under low irradiances while samples collected at 6 h had been exposed to sunlight (Under neutral density screens representative of irradiances at the depth of collection) and samples from 12 h were collected at night (for both experiments). Treatment groups (mean ± SE) with similar superscripts are not statistically different from each other.

doi:10.1371/journal.pone.0072961.g004
the prokaryotic community. Results of the HCO₃⁻ tracer experiment indicate that there is uptake by prokaryotes and then immediate equilibrium of C with the sponge tissue. These results show that there is an active autotrophic community present in X. muta and provides evidence for transfer of C from symbiont to host. Uptake of HCO₃⁻ is not limited to autotrophs, and heterotrophic bacteria may have also taken up HCO₃⁻ for use in anaerobic pathways [63].

The NH₄⁺ tracer experiment indicates that the symbiotic prokaryotic community of X. muta actively assimilated the NH₄⁺ and the host sponge is the likely source of NH₄⁺. The small increase in sponge host δ¹⁵N during the NH₄⁺ experiment may also indicate direct uptake of NH₄⁺ by the sponge, as has been demonstrated for corals [64]. Alternatively, as in the NO₃⁻ tracer experiment, the accumulation of tracer in the sponge fraction may also be explained by transfer of N from the bacteria to the host.

NO₃⁻ can also be utilized by the prokaryotic community, as demonstrated by the increased δ¹⁵N of the bacterial fractions incubated with ¹⁵NO₃⁻ (Fig 4 A). Photosynthetically driven NO₃⁻ uptake has been demonstrated in planktonic communities [65], and may explain the increase in NO₃⁻ uptake when the sponges were exposed to natural solar radiation. However, it should be noted that heterotrophic bacteria could take up NO₃⁻ as well [66]. As the symbiotic nitrifiers in the sponge produce NO₃⁻ it would provide a source of NO₃⁻ for uptake by the photosynthetic community as well as a substrate for denitrification. Similarly, NH₄⁺ is also a likely substrate for aerobic ammonia oxidation by the crenarcheote community, which is supported by the recovery of crenarchaeal amoA genes in X. muta [34], expression of amoA genes [34] and tracer studies [8]. If assimilatory and dissimilatory processes are competing for NO₃⁻, NO₂⁻, and NH₄⁺, which have been documented in other communities [67], then this may have a significant role in nitrogen cycling in the sponge holobiont. Lastly, the anaerobic oxidation of NH₄⁺ (anammox) is another process that may utilize both NH₄⁺ and the NO₂⁻ generated from nitrification. The rates, however, of anammox had been previously demonstrated to occur in X. muta, and we show here that other nitrogen transformations including denitrification and/or anammox may occur in these sponges as well as the importance of active suspension feeding on the nitrogen rich pool of picoplankton. Further work is needed to better characterize the flux of DIN from X. muta and other sponges on Caribbean coral reefs including whether sponges can fix N₂. This will require additional investigations on the functional activity of the symbiotic prokaryotic community of sponges using a combination of experimental and molecular approaches (i.e., transcriptomics) that will yield insight into the taxonomy and function of this community, and how this impacts nutrient fluxes and biogeochemical cycling on Caribbean coral reefs.

**Acknowledgments**

We thank Marc Slattery, Deborah Gochfeld, Erica Henkin, Cole Eason, Sylvester Lee, Julia Stevens, Christopher Freeman, Julie Olson, Mauritius Bell, and the Aquarius team for help in the field. Marshall Otter at the stable isotope laboratory at the Marine Biological Laboratory performed the stable isotope analyses. Marilyn Fogel and Roxane Bowden at the Carnegie Institution of Washington for additional support and assistance with stable isotope analyses. Paul Henderson at the Nutrient Analytical Facility at Woods Hole Oceanographic Institute conducted the nutrient analyses and the flow cytometry analyses were performed at the J.J. MacIsaac Facility for Aquatic Cytometry at the Bigelow Laboratory for Ocean Sciences.

**Author Contributions**

Conceived and designed the experiments: CLF MPL. Performed the experiments: CLF MPL. Analyzed the data: CLF DBM MPL. Contributed reagents/materials/analysis tools: CLF DBM MPL. Wrote the paper: CLF DBM MPL.
Figure 5. Potential dissimilatory N transformations occurring in X. muta based on the observed net flux of NH₄⁺ and NO₃⁻ from the sponge. Font size for a given process indicates the relative importance of that process. Locations may appear more than once due to differences in individual sponges at each location.

doi:10.1371/journal.pone.0072961.g005

References


