Metatranscriptome analyses indicate resource partitioning between diatoms in the field.

The MIT Faculty has made this article openly available. Please share how this access benefits you. Your story matters.

<table>
<thead>
<tr>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>As Published</td>
<td><a href="http://dx.doi.org/10.1073/pnas.1421993112">http://dx.doi.org/10.1073/pnas.1421993112</a></td>
</tr>
<tr>
<td>Publisher</td>
<td>National Academy of Sciences (U.S.)</td>
</tr>
<tr>
<td>Version</td>
<td>Final published version</td>
</tr>
<tr>
<td>Accessed</td>
<td>Tue Apr 18 17:31:50 EDT 2017</td>
</tr>
<tr>
<td>Citable Link</td>
<td><a href="http://hdl.handle.net/1721.1/99118">http://hdl.handle.net/1721.1/99118</a></td>
</tr>
<tr>
<td>Terms of Use</td>
<td>Article is made available in accordance with the publisher's policy and may be subject to US copyright law. Please refer to the publisher's site for terms of use.</td>
</tr>
<tr>
<td>Detailed Terms</td>
<td></td>
</tr>
</tbody>
</table>
Diverse communities of marine phytoplankton carry out half of global primary production. The vast diversity of the phytoplankton has long perplexed ecologists because these organisms coexist in an isotropic environment while competing for the same basic resources (e.g., inorganic nutrients). Differential niche partitioning of resources is one hypothesis to explain this “paradox of the plankton,” but it is difficult to quantify and track variation in phytoplankton metabolism in situ. Here, we use quantitative metatranscriptomie analyses to examine pathways of nitrogen (N) and phosphorus (P) metabolism in diatoms that cooccur regularly in an estuary on the east coast of the United States (Narragansett Bay). Expression of known N and P metabolic pathways varied between diatoms, indicating apparent differences in resource utilization capacity that may prevent direct competition. Nutrient amendment incubations skewed N/P ratios, elucidating nutrient-responsive patterns of expression and facilitating a quantitative comparison between diatoms. The resource-responsive (RR) gene sets deviated in composition from the metabolic profile of the organism, being enriched in genes associated with N and P metabolism. Expression of the RR gene set varied over time and differed significantly between diatoms, resulting in opposite transcriptional responses to the same environment. Apparent differences in metabolic capacity and the expression of that capacity in the environment suggest that diatom-specific resource partitioning was occurring in Narragansett Bay. This high-resolution approach highlights the molecular underpinnings of diatom resource utilization and how cooccurring diatoms adjust their cellular physiology to partition their niche space.

Significance

Nutrient availability plays a central role in driving the activities and large-scale distributions of phytoplankton, yet there are still fundamental gaps in understanding how phytoplankton metabolize nutrients, like nitrogen (N) and phosphorus (P), and how this metabolic potential is modulated in field populations. Here, we show that cooccurring diatoms in a dynamic coastal marine system have apparent differences in their metabolic capacity to use N and P. Further, bioinformatic approaches enabled the identification and species-specific comparison of resource-responsive (RR) genes. Variation of these RR gene sets highlights the disparate transcriptional responses these species have to the same environment, which likely reflects the role resource partitioning has in facilitating the vast diversity of the phytoplankton.
patterns in resource utilization potential through time by tracking the expression of species’ resource-responsive (RR) genes. When simultaneously applied to multiple species in a sample, this approach can resolve differences in the expressed gene compliment and how it is modulated, which may reflect resource partitioning of phytoplankton niche space (35). For example, this approach has uncovered species-specific expression of genes for the transport of organic compounds in the bacterioplankton (36–38), highlighting potential differences in resource partitioning. Although increasingly critical for identifying resource utilization in the bacterioplankton, metatranscriptome profiling has only recently been used to examine resource utilization in coastal eukaryotic phytoplankton populations (39), largely due to challenges in quantifying a transcriptional response in a mixed population and, until recently, the lack of reference genomes and transcriptomes for determining the origin of the transcriptional response. Cooccurring phytoplankton may possess different metabolic capabilities and responses to resource availability, which may then enable resource partitioning and the segregation of the fundamental niche or the realized niche. Knowledge of if and how these organisms modulate their niche space would allow predictive models to better resolve species distribution and ecosystem structure and function in the future ocean (26).

Herein, we examined pathways of resource metabolism between two cooccurring diatoms from the genera *Thalassiosira* and *Skeletonema*, sampled from a time-series site in Narragansett Bay. Narragansett Bay is a highly productive and dynamic estuarine environment on the east coast of the United States with an estimated bay-wide average net production of 269 g C m⁻² y⁻¹ (40). Quantitative metatranscriptomic techniques were developed and used to (i) assign taxonomic designation, (ii) assess and track changes in known metabolic capacity by means of the quantitative molecular fingerprint (OMF), (iii) statistically identify the RR gene set, and (iv) proportionalize the expression of RR genes to track species-specific responses through time using standardized transcriptional differentiation (STD) scores. This multifaceted computational approach enabled the unprecedented resolution of the unique strategies these two diatoms use for resource acquisition.

**Results and Discussion**

**Samples and Sequencing.** Narragansett Bay has seasonal blooms of diatoms that have been monitored through weekly cell counts for over 50 y at a long-term time-series station (41, 42). Five eukaryotic surface metatranscriptome samples were taken from surface seawater collected during May and June of 2012 at the time-series site, yielding over 358 million 100-bp, paired-end cDNA reads from the field [sample 1 (S1)–S5] (SI Appendix, Table 1). In conjunction with these field-based surveys, a nutrient amendment incubation experiment was performed with natural communities on May 30, 2012 (S3) to drive the community toward opposite extremes in the N/P ratio (Redfield ratio) (SI Appendix, Table 2). Eukaryotic metatranscriptomes from the five incubation treatments produced over 264 million 100-bp, paired-end cDNA reads (SI Appendix, Table 1).

To assign taxonomic designation, sequences from the time series were conservatively mapped (such that if a read mapped to more than one gene, it was discarded) to a sequence library containing all assembled sequences and annotations generated through the Marine Microbial Eukaryotic Transcriptome Sequencing Project (MMETSP) (43), which were made public as of March 17, 2014. The custom sequence library contained 401 transcriptomes across 209 species or cultured isolates. Between 62% and 71% of reads from the in situ samples mapped to the MMETSP database with diatoms dominating the libraries, representing 30–46% of the total mapped reads (Fig. 1). The peak in diatom representation coincided with a bloom of *Skeletonema* spp. detected in time-series cell counts (SI Appendix, Fig. 1) and a period of historical overlap between the *Skeletonema* and *Thalassiosira* genera. *Skeletonema* and *Thalassiosira* were well represented during the time period studied in both mapped RNA (Fig. 1) and cell counts (SI Appendix, Fig. 1). *Thalassiosira rotula* was present, but at low abundance, during the time series, whereas *Skeletonema* spp. was abundant, with sampling spanning a bloom of *Skeletonema* (>10 million cells per liter) with peak cell densities in S2 (May 21, 2012) (SI Appendix, Fig. 1). As such, subsequent analyses were focused on these two groups by remapping the data to representative transcriptomes: *T. rotula* and *Skeletonema costatum* (SI Appendix, Table 1). *S. costatum* was chosen because it was the transcriptome from the genus *Skeletonema* that recruited the most hits in the MMETSP data (35).

**Temporal Plasticity in Expressed Metabolic Capacity.** Metatranscriptome short reads were mapped to transcriptomes that had been annotated with Kyoto Encyclopedia of Genes and Genomes (KEGG) orthology (KO) (Dataset 1), allowing the expression of KO gene families within a KEGG module (higher level groupings of KO gene families into pathway or functional classifications) to be examined over time. Normalizing the expression of KEGG modules to the total KEGG annotated reads of each organism across time yielded the QMF, which highlighted differences between the two species and differences across time for each species (Fig. 2). A comparison of the total number of annotated genes falling into each of the KEGG modules revealed a close to one-to-one linear relationship (slope of 1.0948, $R^2 = 0.9123$) (SI Appendix, Fig. 2), indicating that the observed differences are not an artifact of gene distribution between organisms. The QMFs of the two organisms were distinct, and there were significant shifts in the QMF of each species over time reflecting considerable plasticity in the expressed metabolic capacity (SI Appendix, Fig. 3). Central carbohydrate metabolism, C fixation, and other carbohydrate metabolism were some of the most highly expressed KEGG modules in the field for both *Skeletonema* spp. and *T. rotula*, although higher for *Skeletonema* spp., where expression of these pathways peaked during S4, representing over 84% of mapped KEGG reads (Fig. 2). The
Species-Specific Resource Utilization Underpins Physiological Ecology.

KO gene families related to N and P metabolism were examined in the field samples to identify species-specific patterns in resource utilization. *Skeletonema* spp. and *T. rotula* both possess and express core pathways of N and P metabolism (e.g., the ornithine-urea cycle) (Fig. 3). Expression of these individual KO gene families was temporally variable, as was observed with the expression of KEGG modules, but related enzymes in a pathway exhibited a coordinated response (Fig. 3). For example, the nitrate transporter (K02575), nitrate reductase (K10534), and nitrite reductase (K00366) in *Skeletonema* spp. all had peak expression in S2 (Fig. 3). *Skeletonema* spp. and *T. rotula* share pathway homologs, including the same suite of N transporters (ammonium, nitrate, and amino acid), but these genes often had disparate patterns of expression between the two species (Fig. 3).

*Skeletonema* spp., the more abundant diatom, had high expression of KO gene families associated with the acquisition of nitrate and ammonia that were particularly amplified during the S2 bloom event. *T. rotula* had low expression of both of those transporters but high expression of a general amino acid transporter (Fig. 3). Amino acid transport (49) and nitrate transport (50) have previously been found to correlate inversely with intracellular nitrate concentration in the cell or in the presence of ammonia in the media. However, here, two closely related diatoms existing in the same parcel of water and the same nutrient environment are expressing genes to access different pools of dissolved N. Similar to nitrate transport, there was high expression of nitrate/nitrite reductase KO gene families in *Skeletonema* spp., whereas *T. rotula* appears to possess a different N reduction metabolism. This difference is observed in a KO gene family that is absent from the reference transcriptome of *Skeletonema* spp.: hydroxylamine reductase (Fig. 3). This gene has been found in the genomes of both *T. pseudonana* and *Phaeodactylum tricornutum*, and it is thought to have been acquired via lateral transfer from bacteria (51). The enzyme may potentially aid redox balancing and electron cycling from nitrate reduction (52).

Fig. 2. Quantitative metabolic fingerprint depicting the relative expression of KEGG modules for *Skeletonema* spp. and *T. rotula* in Narragansett Bay across the five sampling time points (S1–S5). Color indicates the proportion of total reads mapping to each KEGG module relative to all KEGG annotated reads.

The largest global shift in KEGG module expression was seen in *Skeletonema* spp. on S2 (SI Appendix, Fig. 3), when its density peaked at 11,520,000 cells per liter. The S2 time point for *Skeletonema* spp. had increased QMF signals in ATP synthesis, proteasome, and ubiquitin systems and decreased QMF signals in photosynthesis KEGG module in S2, an order of magnitude higher than other modules (Fig. 2). The temporal plasticity of transcript allocation to different aspects of metabolism for both species was striking and different between the two diatoms underscores the fundamental differences in expressed metabolic capacity that are present in these two cooccurring diatoms and highlights traits of a successful competitor (e.g., high expression of C metabolism).

**Skeletonema** spp., the dominant diatom during the study period (Fig. 1), had a higher proportion of transcripts related to growth relative to *T. rotula*, such as those transcripts encoding aspects of C metabolism, N metabolism, sulfur metabolism, and lipid metabolism (Fig. 2). Conversely, several KEGG modules were more highly expressed in *T. rotula* compared with *Skeletonema* spp., particularly those KEGG modules for glycan metabolism, phosphate, and amino acid transport systems, as well as repair system modules (Fig. 2). The majority of highly expressed KO modules (e.g., N metabolism) were based on moderate to high expression across several KO gene families, but the differences in expression at the module level were due to differences in the expression of a single KO gene family within the KEGG module in some cases. For example, the driver of the difference in the expression of glycan metabolism, which represented upward of 41% of all KEGG annotated reads for *T. rotula* compared with less than 0.6% for *Skeletonema* spp., was primarily associated with the high expression of a putative UDP–N-acetylglucosamine–dolichyl-phosphate N-acetylglucosamineprophosphotransferase (K01001). This gene was identified as a slatfin-like response gene associated with silica polymerization (48). Differences in silica metabolism may partially drive how the fundamental niche is segregated between these two diatoms. Taken together, the contrast in QMF between the two diatoms underscores the fundamental differences in expressed metabolic capacity that are present in these two cooccurring diatoms and highlights traits of a successful competitor (e.g., high expression of C metabolism).
Although the absence of this gene in Skeletonema spp. has not been definitively shown, the marked high expression of this gene in T. rotula suggests that this gene product represents a potential point of segregation in the metabolic capacity of these two species. Together, these data suggest that these species have disparate strategies for acquiring N and these differences...
may partially drive the relative success of *Skeletonema* spp. over the sample period.

Although N has been observed to be a primary nutritional driver in Narragansett Bay (47, 53, 54), P may also drive the dynamics of these two diatoms. *Skeletonema* spp. shows elevated expression of a sodium phosphate cotransporter (NPT), again with peak expression during S2 (bloom). *T. rotula* does not express the NPT as highly but, by comparison, has a much higher transcript count for a putative P transporter that is not detected in *Skeletonema* spp. (Fig. 3). These transporters may have different kinetic properties that allow the two species to diverge in their PO₄ uptake strategies. Genes associated with the scavenging of P from organic molecules, such as glycerophosphoryl diester phosphodiesterase (GDP), also suggest differences in expressed metabolic capacity between the two species. GDP may be associated with exogenous metabolism of dissolved organic P (DOP) or internally in the cleaving of P from lipids (55, 56). The expression of GDP by *Skeletonema* spp., with a peak around S2, and the apparent absence of this transcript in *T. rotula* suggest *Skeletonema* spp. may be accessing a pool of DOP that is not being used by *T. rotula*. In *T. pseudonana*, related transcripts are tightly linked to concomitant changes in the proteome and biochemical activities (56). If these transcriptional patterns are linked to similar changes in activities, then these insights suggest that there is a fundamental difference in the metabolic capacity being expressed in the same environment by the two diatoms. *Skeletonema* spp. is both actively taking up PO₄ and hydrolysing organic sources, whereas *T. rotula* is not hydrolysing DOP and is taking up inorganic PO₄ by a different mechanism. In summary, these data suggest that these two diatoms have a unique metabolic capacity for the utilization of specific forms of N and P. Such disparate resource utilization potential could be a niche-defining feature that underpins diatom diversity as well as the “winner-loser” dynamic observed here with the differences in cell abundance between the species.

**Identification and Modulation of RR Genes in Situ Highlight Species-Specific Differences.** To identify and quantitatively track RR genes in situ, incubation experiments were used to examine species-specific transcriptional responses to shifts in N/P ratios. Comparing the expression patterns between like nutrient treatments (+N vs. −N and +P vs. −P) for each of the organisms enabled the statistical identification of a suite of RR genes (57) and stable reference genes (58). RR gene counts were normalized to the stable reference genes (*SI Appendix, Fig. 4*), resulting in stable gene normalized counts (SGNCs). Calculation of an SGNC is similar in concept to the reference gene normalization done in quantitative RT-PCR (qRT-PCR) (59) or metatranscriptome studies of prokaryotes (60), with the added value of not having to rely on reference genes from model diatoms.

Of the transcripts expressed at greater than two tags per million (TPM) for at least one treatment, 24.5% and 17.9% were identified as RR by being significantly up- or down-regulated in the N or P treatments for *Skeletonema* spp. and *T. rotula*, respectively (*SI Appendix, Table 3 and Dataset 2*). As is common with eukaryote studies (32), the majority of the RR genes do not have a KEGG annotation (Fig. 4A and Dataset 2). The portion of the RR gene set annotated with KEGG ontology for *Skeletonema* spp. and *T. rotula* revealed that, relative to the full KEGG profile, genes comprising genetic information processing associated with replication (encompassing ribosomes, nucleotide replication, and processing) were underrepresented for both organisms in the RR set (*SI Appendix, Fig. 5*). By contrast, the RR sets were enriched for energy metabolism, carbohydrate metabolism, and lipid metabolism, which encompass pathways known to be associated with the metabolism of N and P (Fig. 4A and *SI Appendix, Fig. 5*). Specific genes in this set included, but were not limited to, those genes associated with N assimilation (e.g., glutamate dehydrogenase, glutamine synthase, nitrate reductase), dissolved organic N utilization (e.g., urease, aminopeptidase, amino acid transport system), P scavenging (e.g., phosphate transporter, NPT), and DOP utilization (e.g., phosphatases) (*Dataset 2*). A number of these genes have been shown to be N- or P-responsive in transcriptional studies with cultures of the diatom *T. pseudonana* (56, 61), and transporters and enzymes for the processing of organic N or P, as observed here, are well known to be RR in many phytoplankton (56, 62–64). Overall, these genes demonstrated patterns of regulation in situ (Fig. 4B and *SI Appendix, Fig. 6*) similar to what has been observed in culture (56,
In the incubations, the NPT was significantly up-regulated in the −P treatment for both species (Fig. 4B), which is consistent with P regulation of a T. pseudonana NPT homolog (Thaps_24435) observed in culture experiments (56). Nitrate reductase, which has been shown to be regulated by N in T. pseudonana (Thaps_25299) (61), was up-regulated in −N for T. rotula, but not for Skeletonema spp. (SI Appendix, Fig. 6). In fact, members of this large gene family (SI Appendix, Fig. 7) showed disparate regulation in both species (SI Appendix, Fig. 6). These data demonstrate that the use of nutrient amendments is robust for normalizing and identifying N- and P-responsive genes in the field that are consistent with known signals, but they also point to the value of in situ analyses, because application of a priori knowledge about regulation from model diatoms could lead to misinterpretations.

Of the RR gene sets for Skeletonema spp. and T. rotula, only 17.7% and 12.7% of the genes, respectively, could be annotated with KEGG ontology (Fig. 4A). Identifying differentially regulated genes in situ through experimental manipulations allowed the expression patterns of genes to be tracked even when their function was unknown. As an example, two RR gene families were identified with homologs in Skeletonema spp. and T. rotula (Fig. 4B and SI Appendix, Fig. 7). RR gene family 1 (RR1) was up-regulated in −P compared with +P for both species (Fig. 4B). Homologs from RR1 were also identified in other diatom genomes (Fray_268075, Phatr_19661, Pseemu_319824, and Thaps_32459) (SI Appendix, Fig. 7). Annotations for these genes were limited, although Fray_268075 was identified as possibly involved in intracellular trafficking, secretion, or vesicular transport, suggesting these proteins may be involved in poly-P metabolism (65). RR2 demonstrated significantly different patterns of regulation in the two species: up-regulated in −N compared with +N for T. rotula but down-regulated in −N compared with +N for Skeletonema spp. (Fig. 4B). A homolog from RR2 was identified in T. pseudonana (Thaps_22648) (SI Appendix, Fig. 7) and was poorly characterized, with the best BLAST hit to a human dentin sialophosphoprotein. This finding suggests RR2 could be associated with biomineralization.

To enable cross-comparison of the RR genes between species, their expression was put into a greater metabolic context by proportionalizing the expression in the field to the transcriptional range observed in the incubations with extremes in the N/P ratio. This became particularly important when targeted assessment by qRT-PCR to compare expression patterns between species in culture (66). Briefly, the SGNC of a gene in the field was bounded by the SGNC from each of the nutrient treatments to yield the STD score for both N (STDN) and P (STDp) (Fig. 4C). The STD score was used to compare expression directly relative to its maximum and minimum capacity, where values of STD ≥ 1 indicate signals were similar to the deplete condition and values of STD ≤ 0 indicate similarity to the replete condition. The STDN and STDp were plotted for genes from the NPT and the two highlighted RR gene families over the time series (Fig. 4C). The NPT for both Skeletonema spp. and T. rotula showed elevated expression during S2. RR1, which was also identified as significantly expressed in −P compared with +P, also showed elevation during S2 (the bloom). The expression of RR1, however, was also elevated on S4 for both diatoms, which was not seen for the NPT. However, the STDp was >1 for Skeletonema spp., indicating a far more P-deficient response in Skeletonema spp. compared with T. rotula, which never demonstrated P-sensitive expression in the field comparable to the P-sensitive expression observed in the −P incubations (Fig. 4C). RR2 showed different patterns of expression across time for both species. Most interesting perhaps was the low STDN score for Skeletonema spp. during S2 (the bloom), indicating that RR2 expression was more similar to the +N treatment, whereas the STDN for T. rotula was greater than 1, suggesting that RR2 expression was more similar to the −N treatment (Fig. 4C). These three targeted examples suggest that during the large bloom of Skeletonema spp., Skeletonema spp. was expressing genes in a pattern more similar to the −P and +N treatments, whereas T. rotula was expressing genes similar only to the −N treatment. Notably, these orthogonal patterns were associated with the same environment.

The STDN and STDp for all of the RR genes were calculated (Dataset 2) to expand upon the single-gene analyses above. The RR genes were plotted based on the STDN/STDp ratio (SI Appendix, Fig. 8) to examine how similar the pattern was to the incubation N/P ratio (Fig. 5A and SI Appendix, Fig. 9). Redfield regimes have historically been used to characterize different aquatic environments based on the ratio of nutrient resources required for growth. For example, a Redfield ratio of N/P = 16, here called “Redfield,” would predict neither P nor N limitation. As expected, RR genes identified as N-regulated genes fall primarily into the N/P < Redfield quadrant and P-regulated genes fall primarily into the N/P > Redfield quadrant for both Skeletonema spp. and T. rotula (Fig. 5A). Observing patterns in the distribution of these genes across time, S2 stands out among the time points, where a significant (88%) proportion of the P-regulated genes from Skeletonema spp. move far into the N/P > Redfield quadrant (Fig. 5A). This N/P > Redfield physiology is consistent with the single-gene analyses (Fig. 4C) and suggests P availability may have constrained Skeletonema spp. populations during the bloom sample (S2). Conversely, a large proportion (59%) of the N-regulated genes in T. rotula move into the N/P < Redfield quadrant (Fig. 5A) consistent with the divergent responsiveness of RR2 observed for T. rotula compared with Skeletonema spp. (Fig. 4C). In fact, with the exception of S4 and S5, where T. rotula had even distribution between the N/P > Redfield and N/P < Redfield quadrants, the two species always showed statistically significant (Tukey’s honest significant difference analysis p < 0.05) orthogonal responses in the distribution of the RR gene set across the two quadrants (Fig. 5B and SI Appendix, Fig. 10). These patterns, combined with the temporal variability in gene expression patterns, indicate a finely tuned response to the environment, which is distinctive for each diatom species. Although there are many potential controls on diatom dynamics in Narragansett Bay, including top-down processes like predation (67, 68), these patterns of RR gene expression suggest the presence of bottom-up nutrient control on diatom population dynamics in Narragansett Bay.

This work addresses fundamental knowledge gaps in how phytoplankton species are able to cooccur while they compete for the same basic resources. Cooccurring diatoms appear to have different functional capabilities in N and P metabolism, and this metabolic potential is modulated in field populations in a distinctive way for each diatom. These findings suggest that differential resource partitioning is occurring between these two diatoms in situ. Such resource partitioning could facilitate the vast diversity of the phytoplankton and the structure, function, and productivity of aquatic ecosystems. In culture studies, resource-related transcriptional changes have been shown to be tightly choreographed with changes in protein activities, and biochemical pools (56, 62, 69). If further work is similarly able to link the transcriptional patterns observed here with changes in enzymatic activities or uptake rates, then shifts in the RR gene sets might reflect aspects of the realized niche and how it differs between these two species. These detailed in situ transcriptional comparisons would not have been possible without proportionalization to metabolic capacity (STD), which provides a quantitative means to compare transcriptional patterns directly between species. This approach could be applied to other systems, organisms, or environmental parameters to identify responsive genes and proportionalize their expression, with the aim of answering similar questions about how cooccurring species adjust their cellular physiology to partition their niche space.
Surface seawater was collected for Skeletonema spp. (yellow) andSkeletonema spp. (cyan) and T. rotula (dark blue) were proportionalized relative to the expression of those genes in nutrient incubations, yielding the $S_{D}$ and $S_{B}$ for each gene. These data are plotted for S1–S5. (B) Proportion of identified RR genes falling into the N/P > Redfield and N/P < Redfield quadrants for T. rotula (T) and Skeletonema spp. (S). FC, fold change.

Materials and Methods
Experimental Setup and Sample Collection. Surface seawater was collected and sampled for total community RNA at the long-term sampling site in Narragansett Bay (41°34′12″ ‘N, 71°23′24″ ‘W) during 2012 (May 16, May 21, May 30, June 4, and June 8, here called S1 through S5) in conjunction with the weekly time-series sampling effort. To diminish the influence of diel signals, samples were collected and processed between 0830 and 0900 local time. Near-surface water was collected in an acid-washed carboy and then filtered on polycarbonate filters (5.0-μm pore size, 47 mm) using a peristaltic pump. Filters were then placed in cryovials and stored in liquid N until RNA extraction. In this manner, all samples were preserved within 15 min of collection. In addition to sampling for total community RNA, phytoplankton abundance was measured as part of the long-term weekly survey (70, 71).

A nutrient amendment incubation experiment was performed on May 30, 2012, with S3 representing the t = 0 of the experiment. Water collected in conjunction with S3 was prefiltered through 200-μm mesh to remove large zooplankton grazers and placed into acid-washed 2.5-L bottles. Triplicate bottles were then amended with nutrients to create five treatments: +N, +P, −N, −P, and ambient control. The +N and +P treatments were designed to eliminate the N and P stress signals, respectively, whereas the −N and −P treatments were supplemented with everything except the nutrient in question (e.g., the −N treatment was amended with P, Si, Fe, and vitamins) to force the drawdown of N and P, respectively (SI Appendix, Table 2). N and P amendment concentrations were selected to be 10-fold the seasonal average ambient N and P concentrations in the surface waters of Narragansett Bay measured at station II. The Si, Fe, and f/5 vitamin amendments were made in proportion to the f/5 media ratios (72). Bottles were placed in a flow-through incubator at ambient temperatures and photosynthetically active radiation (PAR) to mimic the collection depth. The incubation was run for 48 h, at which point all treatments were sampled for total community RNA as described above by filtering and snap-freezing 2 L of biomass from each replicate bottle.

RNA Extraction and Sequencing. Filters from triplicate bottles, representing ~6 L of water, were pooled by treatment and extracted for each of the in situ and incubation experiment samples. RNA was extracted from individual filters with the RNeasy Mini Kit (Qiagen), following a modified version of the yeast protocol. Briefly, lysif buffer and RNA-clean zinc beads were added to the filter, and samples were vortexed for 1 min, placed on ice for 30 s, and then vortexed again for 1 min. Samples were processed following the yeast protocol. The resulting RNA was eluted in water and then treated for possible DNA contamination using a TURBO DNA-free Kit (Ambion) following the Rigorous Dnase protocol. RNA from each triplicate was then pooled by sample or treatment, using the RNA Cleanup Protocol from the RNeasy Mini Kit. The total RNA (>1,000 ng for each sample) was then enriched for euukaryotic mRNA through a poly-A pull-down onto oligo-dT beads. The resulting enriched RNA sample then went through library preparation with the TruSeq RNA Prep Kit (Illumina). Libraries were sequenced at the Columbia University Genome Center with an Illumina HiSeq2000. Each sample was sequenced to produce ~60 million, 100-bp, paired-end reads (SI Appendix, Table 1). Raw sequence data quality was visualized using FastQC (73) and then cleaned and trimmed using Trimmomatic version 0.27 (paired-end mode; 6-bp-wide sliding window for quality below 20; minimum length of 25 bp) (74). All project sequence reads are available at the National Center for Biotechnology Information (NCBI) under accession number SRP055134.

Transcriptome and Genome Mapping. To assign a taxonomic identification to the reads, a database was created from transcriptomes made publicly available through the MMETSP as of March 17, 2014. In total, 401 transcriptomes from 209 species or cultured isolates were collected. Like-species transcriptomes were combined (regardless of strain or condition) using CD-HIT-EST (98% identity, word size of 9). The resulting clustered set of transcripts was considered to be the representative transcriptome for the species or cultured isolate. The 209 transcriptomes created in this manner were concatenated to form a comprehensive species-level transcriptome database from the MMETSP library. Due to the large size of the resulting MMETSP database, trimmed reads were mapped to the MMETSP using the Burrows-Wheeler aligner (75) and then counted using the HTSeq 0.6.1 package (76).

Transcriptomes from two ecologically relevant diatom species in Narragansett Bay were selected: S. costatum RCC1716 (MMETSP0013, accessed from the publicly available transcriptome databases of the Moore Foundation Marine Microbiology Initiative-supported MMETSP, National Center for Genome Resources) and T. rotula CCMP3096 (a custom assembly available at NCBI under BioSample SAMN03349676). These transcriptomes were individually clustered using CD-HIT-EST (parameters: -c 0.98, −n 9) (77). The resulting clustered set of transcripts was then concatenated to form a reference transcriptome database. Trimmed reads from the field and incubation samples were mapped to this transcriptome database using
Bowtie2 version 2.2.1 (parameters: −a, −sensitive) (78). As a point of comparison, reads were also mapped using Bowtie2 version 2.2.1 under the same parameters to the genome of the model centric diatom species, T. pseudonana CCMP1335 (version 3.0), an organism not known to be abundant in Narragansett Bay. Mapped reads were then counted by transcript using the HTSeq 0.6.1 python package (parameters: −m union, −n 0) (76). Reads aligning to more than one full transcript were not counted. KEGG pathways were assigned to the assembled sequences with the online KEGG Automatic Annotation Server, using the bidirectional best-hit method to obtain KO annotations. In this study, only genes with a normalized count (NC) (raw count)total number of genes mapped to an organism of at least two TPM in at least one of the field or incubation samples were included, thus limiting the sample set to 4,318 genes for T. rotula (19.3% of the transcriptome) and 20,921 genes for Skeletonema spp. (75.6% of the transcriptome). This difference in coverage is directly related to their relative abundance in the population.

Transcriptome Clustering. To assess relatedness of genes within Skeletonema spp. and T. rotula, the transcriptomes were translated using ORF predictor (proteomics.ysu.edu/tools/ORFPredictor.html) using a reference BLASTx alignment against the NCBI database with an e-5 cutoff (79). These translated peptide sequences were then combined with the translated proteins from the diatom genomes Frigilariaopsis cylindrus CCMP1102 version 1.0, P. tricornutum CCMP632 version 2.0, Pseudonitzschia multiseries CLN-47 version 1.0, and T. pseudonana CCMP1335 version 3.0, which were collected from the Joint Genome Institute database (genome.jgi-psf.org). A protein similarity network was then created using EGN, a software program that automates the reconstruction of gene networks from protein sequences through reciprocal BLASTp analysis (e-value <1e-5, 20% hit identity threshold, 5% best reciprocal threshold of best e-value, 90% minimal match coverage threshold) (80, 81). Networks were then visualized and manipulated using Cytoscape 3.0, where the layout of the network was produced using an edge-weighted, spring-embedded model based on e-value, meaning that genes that are closer together are more similar (82, 83). Known RR genes from previous transcriptome studies of the diatom species T. pseudonana were selected for analysis: (i) the P-responsive gene, Thaps_24435, which is an NPT (56) and (ii) the N-responsive gene, Thaps_25299, which is an assimilatory nitrate reductase (61).

Identification of Stable and Nutrient-Responsive Genes. Intercomparison of nutrient-incubation experiments enabled the identification of both nutrient-responsive genes and stably expressed reference genes for T. rotula and Skeletonema spp. For each organism, RR genes were identified by comparing the counts for that organism in +N to the −N incubation and the +P to the −P incubation, respectively, using analysis of sequence counts (ASC), an empirical Bayes method, which estimates the prior distribution from the data itself (57). ASC analyses were run using raw count data from each species separately. Genes were considered to be differentially regulated between treatments if for a fold change of 2.0, the posterior probability (post hoc Tukey test show the divergence of species across time (P < 0.05).

Normalization of Metatranscriptome Data. Counts from the field were first normalized to the sequences belonging to the species in the library (Eq. 1). For a particular species, c, the number of reads mapping to a gene g, c_{i,g}, was normalized to the sum of all of the counts across all genes for that organism, yielding the NC, similar to normalization techniques used for metatranscriptome data (32, 84):

\[ NC_{i,g} = \frac{c_{i,g}}{\sum_{g \in G} c_{i,g}} \]  

(Henceforth, only genes for which NC > 2 TPM in at least one sample (incubation or field) were considered. To facilitate interspecies comparisons, the NC was normalized to the geometric mean of the set of stable reference genes, R, yielding an SGNC. The calculation of an SGNC (Eq. 2) for metatranscriptome data was designed to emulate the normalization used in qRT-PCR studies (85).

\[ SGNC_{i,g} = \frac{NC_{i,g}}{\left( \prod_{c} NC_{i,c} \right)^{1/R}} \]  

The nutrient-responsive genes identified as differentially expressed in the nutrient incubations (SI Appendix, Table 2) were then selected for investigation in the field metatranscriptomes (S1–S5). The SGNCs from the field (SGNC_{field}) for these nutrient-related genes were bounded by the SGNCs from like nutrient incubations to calculate the STD_{N} and STD_{P} (Eqs. 3 and 4).

\[ STD_{N} = \frac{SGNC_{field} - SGNC_{N}}{SGNC_{N} - SGNC_{N}} \]  

\[ STD_{P} = \frac{SGNC_{field} - SGNC_{P}}{SGNC_{P} - SGNC_{P}} \]  

For example, in calculating STD_{N}, the SGNC_{field} is put in the range of the SGNC_{N} and SGNC_{N}. In consequence, if the STD_{N} for a gene in the field equals 1, it is more similar in expression to the +N treatment, and if it equals 0, it is more similar in expression to the −N treatment. As such, a plot of STD_{N} against STD_{P} can divide the space into two main theoretical quadrants N/P > Redfield (STD_{N} > 1 and STD_{P} < 0) and N/P < Redfield (STD_{N} > 1 and STD_{P} < 0) (SI Appendix, Fig. 8). The total number of genes falling into each of the quadrants was counted by varying the bounds considered: the N/P > Redfield ratio quadrant (STD_{N} > C and STD_{P} < C) for 0.25 < C < 0.75 and the N/P < Redfield ratio quadrant (STD_{N} < C and STD_{P} > C) for 0.25 < C < 0.75. To approximate variation conservatively, the value of C was varied over 10 different values, and the average and SD for the percentages of genes falling into each of the quadrants were quantified. Similarity of data between species by quadrant was assessed using an ANOVA with a generalized linear model. The results from a post hoc Tukey test show the divergence of species across time (P < 0.05).

ACKNOWLEDGMENTS. We thank L. Whitney and M. Mercier for their efforts to grow T. rotula for the assembly. The MMETSP samples used in this study were sequenced, annotated, and assembled with the Assembly by Short Sequences (AbYSS) pipeline at the National Center for Genome Resources, H.A. was supported by the Department of Defense through the National Defense Science and Engineering Graduate Fellowship Program. This research was supported by funds from the National Science Foundation (NSF) Environmental Genomics and NSF Biological Oceanography Programs through Grant OCE-0723667 (to S.T.D., B.D.J., and T.A.R.) and Grant OCE-0962208 (to B.D.J.), and by the Joint Genome Institute/Department of Energy Community Sequencing Program through Grant CSP/795793 (to B.D.J., S.T.D., and T.A.R.). Support was also provided by the Gordon and Betty Moore Foundation through Grant GBF2637 to the National Center for Genome Resources for the MMETSP.