slab (Fig. 4B). This complexity suggests strong stress heterogeneity in subducted slabs (17). However, supershear rupture during the M 6.7 earthquake brings its stress drop down to 32 MPa and its radiation efficiency to about 1.0 (Fig. 4B), which are much closer to values for the M 8.3 Okhotsk mainshock. Therefore, strong stress heterogeneity inside subducted slabs is not required to explain the 2013 Okhotsk mainshock and its M 6.7 aftershock. However, the difference in rupture speed (subshear versus supershear) indicates substantial spatial heterogeneity in the fracture strength or fracture energy within the slab.

Compared with shallow supershear events, this deep event has a relatively small rupture dimension and higher static stress drop (by a factor of ~10). Our estimate of high radiation efficiency (η ≈ 1.0) during the M 6.7 event is also consistent with theoretical predictions of low fracture energy during supershear ruptures (30). This constraint of low fracture energy bears on the question of deep earthquake faulting mechanisms, which is still enigmatic (15, 19). The 1994 Bolivia earthquake involved a large amount of fracture/thermal energy and radiated relatively little energy in seismic waves (16). In terms of energy partitioning, the supershear M 6.7 earthquake represents the opposite end member from the Bolivia earthquake, with almost all the available strain energy being radiated as seismic waves. This contrast is consistent with the idea of more than one rupture mechanism for deep earthquakes in slabs with different thermal states (18, 20, 21). The Okhotsk mainshock and aftershock in a cold slab ruptured with the transformational faulting mechanism, whereas the Bolivia earthquake in a warm slab was dominated by shear melting (18).

REFERENCES AND NOTES
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OCEAN MICROBES
Multispecies diel transcriptional oscillations in open ocean heterotrophic bacterial assemblages

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Oscillating diurnal rhythms of gene transcription, metabolic activity, and behavior are found in all three domains of life. However, diel cycles in naturally occurring heterotrophic bacteria and archaea have rarely been observed. Here, we report time-resolved whole-genome transcriptome profiles of multiple, naturally occurring oceanic bacterial populations sampled in situ over 3 days. As anticipated, the cyanobacterial transcriptome exhibited pronounced diel periodicity. Unexpectedly, several different heterotrophic bacterioplankton groups also displayed diel cycling in many of their gene transcripts. Furthermore, diel oscillations in different heterotrophic bacterial groups suggested population-specific timing of peak transcript expression in a variety of metabolic gene suites. These staggered multispecies waves of diel gene transcription may influence both the tempo and the mode of matter and energy transformation in the sea.

The coordination of biological activities into daily periodic cycles is a common feature of eukaryotes and is widespread among plants, fungi, and animals, including man (1). Among single celled noneukaryotic microbes, diel cycles have been well documented in cyanobacterial isolates (2–4), one halophilic archaeon (5), and bacterial symbionts of fish and squid (6, 7). Some evidence for diel cycling in microbial plankton has also been suggested on the basis of bulk community amino acid incorporation, viral production, or metabolite consumption (8–10). However, the existence of regular diel oscillations in free-living heterotrophic bacterial species has rarely been assessed.

Microbial community RNA sequencing techniques now allow simultaneous determination of whole-genome transcriptome profiles among multiple cooccurring species (11, 12), enabling high-frequency, time-resolved analyses of microbial community dynamics (12, 13). To better understand temporal transcriptional dynamics in oligotrophic bacterioplankton communities, we conducted a high-resolution multiday time series of bacterioplankton sampled from the North Pacific Subtropical Gyre (14).

To facilitate repeated sampling of the same planktonic microbial populations through time, automated Lagrangian sampling of bacterioplankton was performed every 2 hours over 3 days by using a free-drifting robotic Environmental Sample Processor (ESP) (13, 15) (fig. S1). After instrument recovery, planktonic microbial RNA was extracted, purified, converted to cDNA, and sequenced to assess whole-genome transcriptome dynamics of predominant planktonic microbial populations (tables S1 and S2). The recovered cDNAs were dominated by transcripts from
Prochlorococcus and several proteorhodopsin-containing or photoheterotrophic bacteria, including members of the Pelagibacter (SAR11), Roseobacter, SAR116, SAR86, and SAR324 clades (fig. S2).

Phylogenetic analysis of gene transcripts in the most abundant taxa revealed the presence of some microdiversity (figs. S3 to S8). However, the most abundant transcripts sampled at any given time point were dominated by only a few genotypes within each population that persisted throughout the sampling period. An exception was Roseobacter, with transcripts for two different genes (groEL and dnaK) indicating the presence of genotypes that started at a very low

Table 1. Harmonic regression results. Sequence reads are the total number assigned to each taxon bin. Transcripts are the total number of unique ortholog clusters (see table S1) with at least one mapped sequence. Periodic is the total number of sequences identified as showing 24-hour periodicity using harmonic regression. The next row shows the proportion of variance explained by 24-hour periodicity in constrained principal components analysis (PCA). The fifth row indicates the Procrustes correlation between the first two principal components from Prochlorococcus and other taxa (unconstrained principal components analysis as shown in Fig. 3); P values based on 999 permutations. Bottom row is the correlation between pairwise transcriptome similarities at each time point from Prochlorococcus and heterotrophic taxa based on Mantel test on Euclidean distance matrices; P value based on 999 permutations.

<table>
<thead>
<tr>
<th>Taxon</th>
<th>Sequence reads</th>
<th>Transcripts</th>
<th>Periodic</th>
<th>Constrained PCA versus 24-hour clock</th>
<th>Procrustes test versus Prochlorococcus PCA</th>
<th>Mantel test versus Prochlorococcus</th>
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<td>3045</td>
<td>1491</td>
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<td>10</td>
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<td>SAR86</td>
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<td>4732</td>
<td>8</td>
<td>0.10 (P = 0.005)</td>
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Fig. 1. Laboratory versus field comparisons of periodic expression patterns in Prochlorococcus populations. (A) Scatter plot shows time of peak abundance for 973 transcripts identified as significantly periodic in both studies. Histograms show the total number of genes peaking in 1-hour intervals in this study (top) and the laboratory experiment (side). Black bars represent genes identified as significantly periodic in both studies, gray bars represent genes expressed in both studies but significantly periodic in only one, and white bars represent significantly periodic transcripts that were not detected in the other data set. For this comparison, we used published significance cutoffs from the laboratory study (4) but for consistency generated new peak times by using our harmonic regression approach and the published normalized mean expression levels for each time point. In general, the peak times generated using our approach closely matched published values for that data set. (B and C) Plots showing relative expression (normalized to mean expression level) over time for our metatranscriptome (top trace) and in microarray data (bottom trace) for selected transcripts. For comparison, experimental midnights (24 and 48 hours) from the microarray study are aligned with the 12:00 midnight samples from 9 and 10 September, respectively. All adenosine triphosphate (ATP) synthase subunits (B) and selected subunits from photosystem I (C) are shown.
abundance and increased in representation over the course of the time series. This variability could be due to an injection of a new population as water masses mixed during the latter portion of the time series or to an alteration in the relative transcriptional activities of two ecotypes that are responding to changes in the surrounding environment.

Transcriptional activity in *Prochlorococcus* was highly dependent on the time of day. Harmonic regression analyses indicated that nearly half (1491) of all *Prochlorococcus* population transcripts were significantly periodic (Table 1, Table S3, and Fig. 1). The expression patterns observed were similar to those of monocultures growing in controlled laboratory settings (4), but there were also notable differences (Fig. 1). For example, photosystem I gene expression exhibited a double peak in the wild *Prochlorococcus* transcriptome around noon (Fig. 1). In contrast, under laboratory conditions most photosystem I genes, for example, *psaL* and *psaF*, were found to peak just before noon, and *psaA* and *psaB* peaked shortly after noon (4).

The largest discrepancy between *Prochlorococcus* laboratory studies and our field observations was that a considerable number of *Prochlorococcus* transcripts in our field populations peaked around midday (Fig. 1). Some of these genes did exhibit periodicity in cultures but peaked at a different time of day than in field populations. A larger fraction of the field midday-peaking transcripts were either not periodically expressed or were not present in the culture experiments. In addition, 62% of the 10 a.m.–to–4 p.m. peaking transcripts in our field study lacked Kyoto Encyclopedia of Genes and Genomes (KEGG) orthology annotations, as opposed to those peaking in the evening or late at night.

A number of factors may be responsible for differences in transcript dynamics between in laboratory cultures versus field *Prochlorococcus* populations. Maximal light levels at our study site at 23-m depth were frequently twofold higher (450 μmol Q m⁻² s⁻¹) than those used in laboratory microarray experiments (232 μmol Q m⁻² s⁻¹) (4). Fundamental genetic differences between our field populations and the *Prochlorococcus* strain used in laboratory culture experiments likely also contribute to the differences we observed. Other variables, including nutrient composition and organismal interactions, may also be factors in the observed differences. Although we could not identify obvious trends in the type or function of field population transcripts showing peak expression during the midday period, they did include a wide range of enzymatic functions that are more consistent with nutrient-responsive metabolic changes rather than a simple high-light stress response.

An abundant *Roseobacter* population also showed strong diel oscillations in its transcriptome profile, most notably in expressed genes involved in bacteriochlorophyll-associated aerobic anoxic photosynthesis. Overall, a large fraction of *Roseobacter* transcripts were periodically expressed (Table 1). Of these, the majority peaked during daylight hours, with only a few gene transcripts peaking at night (Fig. 2). Although this pattern contrasts with that observed in *Prochlorococcus*, where most normally expressed transcripts peaked at dawn or dusk, it was consistent with transcriptional regulation recently reported in *Dinoroseobacter shibae* (10).

Thirty-five of the 40 significantly periodic *Roseobacter* transcripts that peaked between 11 p.m. and 7 a.m. encoded genes belonging to a large photosynthetic “superoperon” (Fig. S9). Nightly expression of these genes, followed by immediate repression upon light onset, is consistent with the *D. shibae* study (10) and may be preparing cells for efficient solar energy harvest in the early morning hours. Functions that peaked during the daytime hours included ribosomal proteins, respiratory transcripts, genes involved in amino acid metabolism, and transporters (Fig. 2).
Protoehodopsin-containing photoheterotrophs including Pelagibacter (SAR11), SAR116, and SAR86 also showed evidence of diel periodicity in many of their gene transcripts (Fig. 2 and Table 1). All opsin-containing bacteria analyzed (SAR11, SAR116, SAR86, and SAR324) exhibited statistically significant diel oscillations in their protoehodopsin gene transcripts (table S3 and fig. S10). Peak expression of the opsin transcripts occurred near dawn in all these populations (fig. S10), potentially optimizing solar energy capture by the light-driven, proton-pumping rhodopsins.

Principal components analysis distinguished time series samples for each heterotroph by time of day (Fig. 3) and showed significant correlation with the light-driven behavior of Prochlorococcus (Table 1). Overall, these data are consistent with genome-wide transcriptional changes across the day-night cycle for each population. In addition, cooclustering of transcripts using GeneARMA (GA) (fig. S7) revealed suites of gene transcripts that exhibited similar expression patterns among different taxa (Fig. 4, figs. S11 to S14, and table S4). For example, a group of transcripts that fit highly similar GA expression models across multiple species included Pro GA5, Pro GA7, Pro GA9, Pro GA23, SAR11 GA6, SAR11 GA18, SAR116 GA2, and Roseobacter GA8 (Fig. 4, fig. S14, and table S4). These multispecies, day-peaking transcripts (fig. S14 and table S4) included gene products associated with respiration (Prochlorococcus, SAR11, SAR116, and Roseobacter), nitrogen metabolism (Prochlorococcus, SAR11, and SAR116), glycine metabolism (Prochlorococcus, SAR11, and Roseobacter), carbon monoxide metabolism (SAR116 and Roseobacter), and DNA synthesis (Prochlorococcus and Roseobacter). The cooclustering of gene transcripts revealed a complex pattern of expression through the day and across the time series and provides evidence for parallel trends in gene expression across multiple species (Fig. 4, fig. S11 to S14, and table S4).

Together, the transcriptional profiles of Roseobacter, SAR11, SAR116, and SAR86 indicate diel cycling of metabolic gene transcripts and suggest a multispecies wavelike progression of up-regulated gene suites across the day-night cycle (Fig. 4). Most conspicuously, a regular diel succession of translational, transcriptional, and respiratory gene transcripts was followed by peaks in transporter transcripts that possibly reflect a metabolic recovery phase (Fig. 2). Many of these metabolic pathway transcripts peaked earlier in the day in Roseobacter-field populations relative to other bacterial heterotrophs (Figs. 2 and 4 and table S4).

The overall transcriptional profile of SAR324 did not show as many transcript diel oscillations as other heterotrophic taxa. Instead, principal components analysis clustered SAR324 transcripts according to the specific day that they were collected (Fig. 3). In particular, the SAR324 group showed a strong separation between the first portion of the time series and the second in principal components analysis (Fig. 3). This split appears to be associated with the increases in temperature and salinity observed across the time series (fig. S1).

The diurnal patterns reported here for open-ocean heterotrophic bacterioplankton were different from those observed in a previous study of phylogenetically related coastal bacterioplankton using similar methods (12). For example, coastal versus open ocean SAR11 populations revealed differential expression levels among several orthologous transcript categories (fig. S15). Additionally, whereas the open-ocean SAR11 populations reported here exhibited statistically significant diel oscillations for many gene transcripts (Fig. 2), the coastal SAR11 populations did not.

Currently available data are insufficient to provide definitive mechanistic explanations for the diel behaviors we observed in different heterotrophic bacterioplankton species. It is possible that photoreceptors in these bacteria are involved in regulating light-dark cycles of transcriptional activity. Marine Roseobacter species have previously been shown to regulate their global transcriptional behavior in response to light (16), and laboratory cultures of Pelagibacter also exhibit light-responsive metabolic behaviors (18). However, differences between the behaviors of SAR11 coastal versus open ocean field populations (fig. S15), as well as comparisons of several taxa in our field study versus laboratory experiments on related cultivated isolates (Fig. 1), suggest that other factors may be at play in regulating diel behavior among these different bacterioplankton populations.

Previous studies have proposed that tight metabolic coupling between primary producers and consumers in microbial plankton might elicit conspicuous diel cycling in heterotrophic bacterial activities (9). The diel cycling we observed among different bacterioplankton species is consistent with this hypothesis, with multiple coexisting heterotroph populations exhibiting diurnal oscillations resembling those of their photoautotrophic neighbors. We postulate that the tightly coupled multispecies temporal expression patterns observed may elicit corresponding waves of species-specific metabolic responses at regular time intervals, potentially coordinating diverse biogeochemical activities in these complex microbial communities. Such temporal coordination of biogeochemical activities among multiple species may be important regulators of both the tempo and the mode of microbial matter and energy transformation in the sea.
Fig. 4. Timing of expression of functional gene clusters among different taxa, clustered by the similarities of their temporal expression patterns.

Heat map shows cluster models for all GA clusters, colored by mean-centered relative expression (red, high; blue, low). Black and yellow bars show the daily photoperiod. Each box represents a single sampling event; for sample times, see table S1. Dendrograms show cluster model similarity (Pearson correlations, average linkage clustering, scale bar at upper right represents a correlation of 0.5). The total number of genes (column A), significantly periodic genes (B), and genes associated with photosynthesis (C), ribosome (D), oxidative phosphorylation (E), amino acid metabolism (F), and transport (G) (defined as for Fig. 2), are listed for each cluster. See table S4 for the identities of genes found within any specific GeneARMA (GA) cluster.
**MICROECONOMICS**

Harnessing naturally occurring data to measure the response of spending to income

Michael Gelman, 1 Shachar Kariv, 2 Matthew D. Shapiro, 1,3* Dan Silverman, 3,4 Steven Tedes 3, 5

This paper presents a new data infrastructure for measuring economic activity. The infrastructure records transactions and account balances, yielding measurements with scope and accuracy that have little precedent in economics. The data are drawn from a diverse population that overrepresented males and younger adults but contains large numbers of underrepresented groups. The data infrastructure permits evaluation of a diverse population that overrepresents males and younger adults but contains large numbers of underrepresented groups. The data infrastructure permits evaluation of a benchmark theory in economics that predicts that individuals should use a combination of saving, borrowing, and making the timing of income irrelevant for the timing of spending. As in previous studies and in contrast to the predictions of the theory, there is a response of spending to the arrival of anticipated income. The data also show, however, that this apparent excess sensitivity of spending results largely from the coincident timing of regular income and regular spending. The remaining excess sensitivity is concentrated among individuals with less liquidity.

Economic researchers and policy-makers have long sought high-quality measures of individual income, spending, and assets from large and heterogeneous samples. For example, when policy-makers consider whether and how to stimulate the economy, they need to know how individuals will react to changes in their income. Will individuals spend differently? Will they save at a different rate or reduce their debt, and when? There are many obstacles to obtaining reliable answers to these important questions. One obstacle is that existing data sources on individual income and spending have substantial limits in terms of accuracy, scope, and frequency.

This paper advances the measurement of income and spending with new high-frequency data derived from the actual transactions and account balances of individuals. It uses these measures to evaluate the predictions of a benchmark economic theory that states that the timing of anticipated income should not matter for spending. Like previous research, it finds that there is a response of spending to the arrival of anticipated income. The data show that, on average, an individual’s total spending rises substantially above average daily spending on the day that a paycheck or Social Security check arrives, and remains high for at least the next 4 days. The data also allow the construction of variables that show, however, that this apparent excess sensitivity of spending results in large part from the coincident timing of regular income and regular spending. The remaining excess sensitivity is concentrated among individuals who are likely to be liquidity-constrained.

Traditionally, researchers have used surveys such as the Consumer Expenditure Survey (CEX) to measure individual economic activity. Such surveys are expensive to implement and require considerable effort from participants and are therefore fielded infrequently, with modest-sized samples. Researchers have recently turned to administrative records, which are accurate and can be frequently refreshed, to augment survey research. So far, however, the administrative records have typically represented just a slice of the population.

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**Table 1. Check versus ACS demographics (percent).** The sample size for Check is 59,072, 35,417, 28,057, and 63,745 for gender, age, education, and region, respectively. The sample size for ACS is 2,441,532 for gender, age, and region and 2,158,014 for education.

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<tr>
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**Census Bureau region**

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PERSPECTIVES

Taking the pulse of ocean microbes

Gene expression patterns of open-ocean microbes are tightly coupled

By E. Virginia Armbrust

Far from land, relatively few types of very small microbes thrive in the warm, nutrient-poor surface waters of the subtropical gyres. Here, the cyanobacterium Prochlorococcus dominates photosynthesis and generates a sizable fraction of the organic matter that fuels the growth of abundant bacteria, including Pelagibacter and a number of uncultured strains (1, 2). On page 207 of this issue, Ottesen et al. (3) use high-resolution sampling to document an exquisitely tight coupling between light-driven gene expression patterns of Prochlorococcus and gene expression patterns of other members of the bacterial community in surface waters at Station Aloha in the North Pacific Subtropical Gyre (see the photo and figure). The study illustrates the power of combining at-sea observation technology with DNA sequencing to understand marine microbial processes.

To obtain microbial samples at sea, oceanographers commonly follow “drifters” that track currents, collecting samples from as close to the drifter as possible with the goal of repeatedly sampling the same parcel of water. The robotic environmental sample processor used by Ottesen et al. represents a crucial advance in sampling technology: It both tracks a given parcel of water and automatically filters and preserves microbial communities for community gene transcription (transcriptome) analysis. Such innovations in autonomous sampling are essential for understanding how marine organisms respond to the rapid chemical and physical changes in today’s oceans.

One motivation for sampling microbial communities at sea is the premise that the...
Exquisite synchrony. Surface waters of subtropical gyres contain relatively few types of abundant microbes, as indicated by the different colors. Photosynthesis in these regions is dominated by the cyanobacterium Prochlorococcus. Ottesen et al. have used autonomous sampling technology to obtain in situ transcription profiles of microbial communities over the day/night cycle. Profiles for a subset of genes from two idealized microbes are shown—one photosynthetic, such as Prochlorococcus, and the other containing the light-driven proton pump proteorhodopsin. The bar at the bottom indicates night (dark bar) and day (yellow bar). The observed periodicity in transcription profiles may translate into a periodicity of metabolic processes in these open-ocean communities.

The observed synchrony of gene expression may rely on at least two factors: The pulse of organic carbon from Prochlorococcus may harmonize the overall transcriptional responses of the bacteria, and proteorhodopsin may further synchronize the transcriptional patterns of particular suites of genes to the light/dark cycle. These microbial interactions could be particularly important in nutrient-poor open-ocean environments, in which genomic streamlining (7) may leave organisms dependent on one another for lost metabolic functions (the black queen hypothesis) (11). Additional tiers of microbial interactions could also be at play if specific organic molecules serve as signals that manipulate the behavior of microorganisms in manners similar to that seen in terrestrial ecosystems (12). The enormous data set generated by Ottesen et al. provides a fantastic resource with which to address these questions.

The ability to simultaneously monitor transcriptional profiles of multiple organisms in situ over time provides the opportunity to answer questions about how microorganisms interact with their environment and with each other, and how these interactions influence ecosystem stability. The large proportion of genes with unknown functions will continue to present challenges. The greatest challenge will be to understand how far-reaching the genetic choreography observed at Station Aloha is over time and space, and how these interactions expand across the entire microbial community.

REFERENCES