Title
Genomic islands and the ecology and evolution of Prochlorococcus

Permalink
https://escholarship.org/uc/item/6506q5sk

Journal
Science, 311(5768)

ISSN
0036-8075

Authors
Coleman, ML
Sullivan, MB
Martiny, AC
et al.

Publication Date
2006-03-24

DOI
10.1126/science.1122050

License
CC BY 4.0

Peer reviewed
Genomic Islands and the Ecology and Evolution of Prochlorococcus

Maureen L. Coleman, Matthew B. Sullivan, Adam C. Martiny, Claudia Steglich, Kerrie Barry, Edward F. DeLong, Sallie W. Chisholm

Prochlorococcus ecotypes are a useful system for exploring the origin and function of diversity among closely related microbes. The genetic variability between phenotypically distinct strains that differ by less than 1% in 16S ribosomal RNA sequences occurs mostly in genomic islands. Island genes appear to have been acquired in part by phage-mediated lateral gene transfer, and some are differentially expressed under light and nutrient stress. Furthermore, genome fragments directly recovered from ocean ecosystems indicate that these islands are variable among co-occurring Prochlorococcus cells. Genomic islands in this free-living phototroph share features with pathogenicity islands of parasitic bacteria, suggesting a general mechanism for niche differentiation in microbial species.

Closely related bacterial isolates often contain remarkable genomic diversity (1, 2). Although its functional consequences have been described in a few model heterotrophic microbes (3), little is known about genomic microdiversity in the microbial phototrophs that dominate aquatic ecosystems. The marine cyanobacterium Prochlorococcus offers a useful system for studying this issue, because they are globally abundant, have very simple growth requirements, have a very compact genome (1.7 to 2.4 megabases [Mb]), and live in a well-mixed habitat. Although the latter appears to offer few opportunities for niche differentiation, Prochlorococcus populations consist of multiple coexisting ecotypes (4), whose relative abundances vary markedly along gradients of light, temperature, and nutrients (5–9). Even two high-light adapted (HL) ecotypes, whose type strains (MED4 and MIT9312) differ by only 0.8% in 16S ribosomal RNA (rRNA) sequence, have substantially different distributions in the wild (5–9).

Although whole-genome comparisons between the most distantly related Prochlorococcus isolates (97.9% 16S rRNA identity) have revealed the gross signatures of this niche differentiation (10), important insights into the evolution of diversity in this group likely lie in comparisons between very closely related strains, and between coexisting genomes from wild populations. Thus, we compared the complete genomes of the type strains, MED4 and MIT9312, that represent the two HL clades, and we analyzed genome fragments from wild cells belonging to these clades from the Atlantic and Pacific oceans.

The 1574 shared genes of MED4 and MIT9312 have conserved order and orientation, except for a large inversion around the replication terminus (Fig. 1). The average G + C content is similar in both genomes (31%), and the median sequence identity of the shared genes is 78%, surprisingly low for strains so similar at the rRNA locus (11). For most genes, synonymous sites are saturated and protein sequence identity is low (median 80%); this is likely a function of high mutation rates, given that HL Prochlorococcus lack several important DNA-repair enzymes (10, 12).

The strain-specific genes between MED4 and MIT9312 (236 in MIT9312 and 139 in MED4) occur primarily (80 and 74%, respectively) in five major islands (Fig. 1). Thus, these genomes have a mosaic structure similar to that of Escherichia coli genomes (1), though on a smaller scale. The islands are located in the same position in both genomes, implying that they are hotspots for recombination, and the length of island genes is similar to the whole-genome average, suggesting that they are not degraded. We hypothesize that these islands arose via lateral gene transfer and continually undergo rearrangement, on the basis of a number of characteristics. First, three islands are associated with tRNA genes (fig. S1), which are common integration sites for mobile elements (13). Second, the 3’ end of tRNA-proline, which flanks ISL3 in both genomes, is repeated 13 times in MIT9312-ISL3 (Fig. 2A) and three times in MED4-ISL3 (fig. S2), suggesting repeated remodeling of this island. Third, some of the genes found in a particular island in MED4 are found in a different island in MIT9312 (Fig. 1), a rearrangement that may have been mediated by a 48-base pair sequence element we call PRE1 (Prochlorococcus repeat element 1; fig. S3); portions of PRE1 are repeated, almost exclusively in islands, 13 times in MED4 (fig. S2), and 9 times in MIT9312 (Fig. 2A). Finally, up to 80% of the genes in any given MIT9312 island are most similar to the genes of noncyanobacterial organisms including phage, Eukarya, and Archaea, consistent with the recent observation that horizontally acquired genomic islands reflect a gene pool that differs from that of the core genome (14).

It is likely that phage, which often carry host genes (15, 16), mediate some of the island-associated lateral gene transfer, and the hli gene family in particular appears to have undergone repeated phage-host gene exchange (16). Of the 24 hli genes in MIT9312, 18 are found in the five major islands or their flanking regions. All 18 belong to the multicopy and sporadically distributed group that includes phage copies (Fig. 2A) and is well differentiated from widespread single-copy hli genes found in cyanobacteria (16). Other phagelike genes in islands include an integrase, DNA methylases, a second endonuclease (15), further supporting a link between phage and island dynamics.

Many island genes in the two strains appear to encode functions related to physiological stress and nutrient uptake and thus may be important in the high-light, low-nutrient surface waters dominated by HL Prochlorococcus. ISL2 and ISL5 in MIT9312, for example, encode 12 of the 24 hli genes, known to be important under a variety of stress conditions (17); they also encode two outer-membrane transporters.

Fig. 1. Whole-genome alignment showing the positions of orthologous genes in MED4 and MIT9312. Strain-specific genes appear on the axes. The locations of five major islands defined by whole-genome alignment (25) are shaded.
proteins; and a cyanophage-like homolog of phoH thought to be involved in the phosphate stress response (15). ISL3 in this strain contains a paralog of pshF, which encodes part of cytochrome b559, thought to protect against photoinhibition (18). Islands also contain genes involved in nutrient assimilation, including a cyanate transporter and lyase in MED4 and two transporters, for manganese/iron and amino acids, in MIT9312 (fig. S1).

In addition to genes involved in potentially growth-limiting processes, islands also contain genes that could play a role in selective mortality. ISL4 in both MED4 and MIT9312 encodes proteins involved in cell surface modification, including biosynthesis of lipopolysaccharide, a common phage receptor (19) (fig. S1). Phages are important agents of mortality in the oceans (20), and thus cell surface properties are likely under strong selection.

Clearly, for island genes to influence a cell’s fitness, they must be expressed. When MED4 cells are starved for phosphorus, nine ISL5 genes are differentially expressed, nearly all of unknown function (table S1). When cells are shifted to high light, 38 island genes are differentially expressed, including seven hli genes (table S1) that in Synechocystis encode proteins that accumulate when cells absorb excess excitation energy (e.g., under high light, nutrient limitation, and low temperatures) (17). Thus, 26% of all MED4 island genes are differentially expressed under P starvation or high-light stress; only one of these is differentially expressed under both conditions (concealed hypothetical gene PMM1416), suggesting that island genes contribute to specific stress responses.

The genome variation within the eMIT9312 clade [sensu (7)] was examined in wild populations of Prochlorococcus by aligning short genome fragments from the Sargasso Sea (21), where this clade dominates (7), against the MIT9312 genome (Fig. 2B). Nearly constant coverage was observed, confirming a stable core genome, except for notable gaps at ISL1, ISL3, and ISL4. This finding indicates that very few wild sequences match genes in these islands, and it supports the hypothesis that these regions are hypervariable in HL Prochlorococcus genomes. In contrast, genes belonging to ISL2 and ISL5 are relatively well represented in the Sargasso Sea data set (Fig. 2B, fig. S2). In MED4 and MIT9312, these islands contain about half of the hli genes, lack the tRNA genes implicated in integration of mobile elements, and contain a smaller fraction of noncyanobacterial genes than do the other islands. This finding suggests that the genes in these islands have become fixed in this wild population.

Examination of 36 large genome fragments (1.1Mb total sequence; median size 34 kb) (table S2) from the Hawaii Ocean Time-Series Station (22) further confirms that a stable core genome surrounds islands of variability, because most fragments showed remarkable conservation of gene content and order with respect to the MED4 and MIT9312 genomes. Thirty-four of the 36 fragments were more similar to MIT9312 than to MED4; two contained tRNA operons, confirming their phylogenetic affiliation with the eMIT9312 clade (fig. S4). The eMIT9312 fragments have about 90% identity with the MIT9312 genome and about 80% with MED4 (Table 1). Collectively, these results suggest that the wild eMIT9312 population is a coherent group identifiable by sequence similarity in the absence of an rRNA operon (1). eMIT9312 genome fragments from this wild population are more similar to each other than to the genome of the type strain MIT9312 (isolated from the Atlantic Ocean), but still share only 93% average sequence identity (Table 1), indicating high coexisting diversity in core genes.

Five eMIT9312 genome fragments from the Hawaii sample border the major islands defined above. About 60% of the genes in these islands have no ortholog in either MED4 or MIT9312, and two fragments border ISL1, yet their gene content is largely different from each other and from the MIT9312 and MED4 genomes (fig. S5). Indeed, a third of the island genes in these two fragments are novel, i.e., have no detectable homologs, implying that cells have access to a large novel gene pool in the oceans (14). Like the islands in the MED4 and MIT9312 genomes, these two fragments contain signatures of mobility, including duplicated tRNA genes, copies of the repeat PRE1, and an integrase gene. This reveals that islands are dynamic even within a single ecotype clade as we have defined it.

One observation that stimulated this work is the dramatic difference in distribution and abundance of the two HL Prochlorococcus ecotype clusters (5–9), as defined by their rRNA internal transcribed spacer (ITS) sequence similarity. Although strains belonging to these two clusters have different island gene content, so do cells from field populations that belong to a single cluster. Therefore, other genomic features are likely to be important in explaining niche differentiation between eMED4 and eMIT9312 cells in the wild. Differential temperature adaptation, for example, which is thought to be an important determinant of ecotype distribu-

Fig. 2. Features of genomic islands (shaded) in the Prochlorococcus strain MIT9312 genome compared with wild sequences from the Atlantic and Pacific Oceans. (A) Locations of repetitive elements and hli genes in MIT9312, shown above or below the horizontal line for the forward or reverse strand, respectively. hli genes shown in pink belong to the single-copy conserved group and those shown in blue belong to the multicopy phage-encoded group (16). (B) Percent identity of Sargasso Sea shotgun database sequences (21) aligned to MIT9312 (top, left axis) and average coverage in the database of a given position in the MIT9312 genome (bottom, right axis). Log2(coverage) is set to —2 when coverage equals 0. (C) Genomic loci and percent identity of wild genome fragments (eMED4-like unless noted) aligned to MIT9312. Where the alignment is interrupted, a black line connects aligned segments of a single fragment. Fragments are projected down to 70% horizontal to visualize total coverage.

Table 1. Median pairwise percent identities, for all orthologous gene pairs and for large aligned regions >4 kb (23). Numbers in parentheses indicate the number of orthologous gene pairs from which the median was calculated.

<table>
<thead>
<tr>
<th>Orthologs (nucleotides)</th>
<th>MED4–MIT9312</th>
<th>MED4-eMIT9312 fragments</th>
<th>MIT9312-eMIT9312 fragments</th>
<th>Overlapping fragments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Orthologs (amino acids)</td>
<td>78.4 (1574)</td>
<td>79.5 (1063)</td>
<td>90.6 (1092)</td>
<td>93.2 (434)</td>
</tr>
<tr>
<td>Large aligned regions</td>
<td>79.0</td>
<td>79.9</td>
<td>90.7</td>
<td>92.6</td>
</tr>
</tbody>
</table>
Toll-Like Receptor Triggering of a Vitamin D–Mediated Human Antimicrobial Response

Philip T. Liu,1,2* Steffen Stenger,4a Huiying Li,3 Linda Wenzel,4 Belinda H. Tan,1,2 Stephan R. Kruitzik,2 Maria Teresa Ochoa,4 Jürgen Schaubert,5 Kent Wu,6 Christoph Meinken,4 Diane L. Kamen,6 Manfred Wagner,4 Robert Bals,8 Andreas Steinmeyer,9 Ulrich Zügäl,10 Richard L. Gallo,3 David Eisenberg,3 Martin Hewison,11 Bruce W. Hollis,12 John S. Adams,11 Barry R. Bloom,13 Robert L. Modlin1,2†

In innate immune responses, activation of Toll-like receptors (TLRs) triggers direct antimicrobial activity against intracellular bacteria, which in murine, but not human, monocytes and macrophages is mediated principally by nitric oxide. We report here that TLR activation of human macrophages up-regulated expression of the vitamin D receptor and the vitamin D–1–hydroxylase genes associated with known antimicrobial functions (5), and the mechanism of human microbial activity remains unresolved.

In studies of resistance to M. tuberculosis, we observed that activation of TLR2/1 reduced the viability of intracellular M. tuberculosis in human monocytes and macrophages but not in monococyte-derived dendritic cells (DCs) (Fig. 1A and (5, 6)). Consequently, we used DNA microarrays to examine gene expression profiles of monocytes and DCs stimulated with a synthetic 19-KD M. tuberculosis–derived lipo-peptide (TLR2/1L) or treated with medium (6). A two-way ANOVA was applied to the array data to identify genes differentially expressed in the two cell types after TLR2/1L treatment (6).

The innate immune system provides a rapid host mechanism for defense against microbial pathogens in Drosophila, innate immunity is mediated in part by the Toll family of pattern-recognition receptors, whose activation induces expression of a series of antimicrobial peptides (1). The mammalian TLR homologs, including the TLR2 and TLR1 heterodimer (2), similarly recognize a variety of microbial-derived ligands, including bacterial lipopeptides. Activation of TLRs results in a direct antimicrobial response in monocytes and macrophages in vitro. In mice, this activity is mediated principally through generation of nitric oxide (5, 4). However, we found that TLR2/1–induced antimicrobial activity in human macrophages is not affected by inhibitors of nitric oxide or reactive oxygen intermediates (5), and the mechanism of human microbial activity remains unresolved.

In studies of resistance to M. tuberculosis, we observed that activation of TLR2/1 reduced the viability of intracellular M. tuberculosis in human monocytes and macrophages but not in monococyte-derived dendritic cells (DCs) (Fig. 1A and (5, 6)). Consequently, we used DNA microarrays to examine gene expression profiles of monocytes and DCs stimulated with a synthetic 19-KD M. tuberculosis–derived lipo-peptide (TLR2/1L) or treated with medium (6). A two-way ANOVA was applied to the array data to identify genes differentially expressed in the two cell types after TLR2/1L treatment (6).

The innate immune system provides a rapid host mechanism for defense against microbial pathogens in Drosophila, innate immunity is mediated in part by the Toll family of pattern-recognition receptors, whose activation induces expression of a series of antimicrobial peptides (1). The mammalian TLR homologs, including the TLR2 and TLR1 heterodimer (2), similarly recognize a variety of microbial-derived ligands, including bacterial lipopeptides. Activation of TLRs results in a direct antimicrobial response in monocytes and macrophages in vitro. In mice, this activity is mediated principally through generation of nitric oxide (5, 4). However, we found that TLR2/1–induced antimicrobial activity in human macrophages is not affected by inhibitors of nitric oxide or reactive oxygen intermediates (5), and the mechanism of human microbial activity remains unresolved.

In studies of resistance to M. tuberculosis, we observed that activation of TLR2/1 reduced the viability of intracellular M. tuberculosis in human monocytes and macrophages but not in monococyte-derived dendritic cells (DCs) (Fig. 1A and (5, 6)). Consequently, we used DNA microarrays to examine gene expression profiles of monocytes and DCs stimulated with a synthetic 19-KD M. tuberculosis–derived lipo-peptide (TLR2/1L) or treated with medium (6). A two-way ANOVA was applied to the array data to identify genes differentially expressed in the two cell types after TLR2/1L treatment (6).

The innate immune system provides a rapid host mechanism for defense against microbial pathogens in Drosophila, innate immunity is mediated in part by the Toll family of pattern-recognition receptors, whose activation induces expression of a series of antimicrobial peptides (1). The mammalian TLR homologs, including the TLR2 and TLR1 heterodimer (2), similarly recognize a variety of microbial-derived ligands, including bacterial lipopeptides. Activation of TLRs results in a direct antimicrobial response in monocytes and macrophages in vitro. In mice, this activity is mediated principally through generation of nitric oxide (5, 4). However, we found that TLR2/1–induced antimicrobial activity in human macrophages is not affected by inhibitors of nitric oxide or reactive oxygen intermediates (5), and the mechanism of human microbial activity remains unresolved.

In studies of resistance to M. tuberculosis, we observed that activation of TLR2/1 reduced the viability of intracellular M. tuberculosis in human monocytes and macrophages but not in monococyte-derived dendritic cells (DCs) (Fig. 1A and (5, 6)). Consequently, we used DNA microarrays to examine gene expression profiles of monocytes and DCs stimulated with a synthetic 19-KD M. tuberculosis–derived lipo-peptide (TLR2/1L) or treated with medium (6). A two-way ANOVA was applied to the array data to identify genes differentially expressed in the two cell types after TLR2/1L treatment (6).

The innate immune system provides a rapid host mechanism for defense against microbial pathogens in Drosophila, innate immunity is mediated in part by the Toll family of pattern-recognition receptors, whose activation induces expression of a series of antimicrobial peptides (1). The mammalian TLR homologs, including the TLR2 and TLR1 heterodimer (2), similarly recognize a variety of microbial-derived ligands, including bacterial lipopeptides. Activation of TLRs results in a direct antimicrobial response in monocytes and macrophages in vitro. In mice, this activity is mediated principally through generation of nitric oxide (5, 4). However, we found that TLR2/1–induced antimicrobial activity in human macrophages is not affected by inhibitors of nitric oxide or reactive oxygen intermediates (5), and the mechanism of human microbial activity remains unresolved.

In studies of resistance to M. tuberculosis, we observed that activation of TLR2/1 reduced the viability of intracellular M. tuberculosis in human monocytes and macrophages but not in monococyte-derived dendritic cells (DCs) (Fig. 1A and (5, 6)). Consequently, we used DNA microarrays to examine gene expression profiles of monocytes and DCs stimulated with a synthetic 19-KD M. tuberculosis–derived lipo-peptide (TLR2/1L) or treated with medium (6). A two-way ANOVA was applied to the array data to identify genes differentially expressed in the two cell types after TLR2/1L treatment (6). Genes up-regulated in monocytes, but not in DCs, with significant P values (P < 0.05; the false discovery rate (FDR), which is the expected proportion of false rejections among all rejections, was 0.09) were cross-referenced against a list of genes associated with known antimicrobial function, yielding two candidates: vitamin D receptor (VDR) and S100A12, a calcium-binding pro-inflammatory molecule (7) (Fig. 1B). Although TLR2/1 stimulation of DCs up-regulated specific genes characteristic of activation (Fig. 1B), the selective up-regulation of the VDR gene in monocytes prompted us to examine further selected VDR-related genes. From these analyses,