DNA sequence of both chromosomes of the cholera pathogen *Vibrio cholerae*

John F. Heidelberg*, **Jonathan A. Eisen***, **William C. Nelson***, **Rebecca A. Clayton***, **Michelle L. Gwinn***, **Robert J. Dodson***, **Daniel H. Haft***, **Erin K. Hickey***, **Jeremy D. Peterson***, **Lowell Umayam***, **Steven R. Gill***, **Karen E. Nelson***, **Timothy D. Read***, **Hervé Tettelin***, **Delwood Richardson***, **Maria D. Ermolaeva***, **Jessica Vamathevan***, **Steven Bass***, **Haiying Qin***, **Ioana Dragoi***, **Patrick Sellers***, **Lisa McDonald***, **Teresa Utterback***, **Robert D. Fleishmann***, **William C. Nierman***, **Owen White** *, **Steven L. Salzberg***, **Hamilton O. Smith***†, **Rita R. Colwell**‡, **John J. Mekalanos**§, **J. Craig Venter***† & **Claire M. Fraser***

* *The Institute for Genomic Research, 9712 Medical Center Drive, Rockville, Maryland 20850, USA*
‡ *Center of Marine Biotechnology, University of Maryland Biotechnology Institute, 701 East Pratt Street, Baltimore, Maryland 21202, USA, and Department of Cell and Molecular Biology, University of Maryland, College Park, Maryland 20742, USA*
§ *Harvard Medical School, Department of Microbiology and Molecular Genetics, 200 Longwood Avenue, Boston, Massachusetts 02115, USA*

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Here we determine the complete genomic sequence of the Gram negative, γ-Proteobacterium *Vibrio cholerae* El Tor N16961 to be 4,033,460 base pairs (bp). The genome consists of two circular chromosomes of 2,961,146 bp and 1,072,314 bp that together encode 3,885 open reading frames. The vast majority of recognizable genes for essential cell functions (such as DNA replication, transcription, translation and cell-wall biosynthesis) and pathogenicity (for example, toxins, surface antigens and adhesins) are located on the large chromosome. In contrast, the small chromosome contains a larger fraction (59%) of hypothetical genes compared with the large chromosome (42%), and also contains many more genes that appear to have origins other than the γ-Proteobacteria. The small chromosome also carries a gene capture system (the integron island) and host 'addiction' genes that are typically found on plasmids; thus, the small chromosome may have originally been a megaplasmid that was captured by an ancestral *Vibrio* species. The *V. cholerae* genomic sequence provides a starting point for understanding how a free-living, environmental organism emerged to become a significant human bacterial pathogen.

Vibrio cholerae is the aetiological agent of cholera, a severe diarrhoeal disease that occurs most frequently in epidemic form[1]. Cholera has been epidemic in southern Asia for at least 1,000 years, but also spread worldwide to cause seven pandemics since 1817 (ref. 1). When untreated, cholera is a disease of extraordinarily rapid onset and potentially high lethality. Although clinical management of cholera has advanced over the past 40 years, cholera remains a serious threat in developing countries where sanitation is poor, health care limited, and drinking water unsafe.

Vibrio cholerae as a species includes both pathogenic and nonpathogenic strains that vary in their virulence gene content[2]. This bacterium contains a wide variety of strains and biotypes, receiving and transferring genes for toxins[3], colonization factors[4,5], antibiotic resistance[6], capsular polysaccharides that provide resistance to chlorine[7] and new surface antigens, such as the 0139 lipopolysaccharide and O antigen capsule[8,9]. The lateral or horizontal transfer of these virulence genes by phage[3], pathogenicity islands[10,11] and other accessory genetic elements[12] provides insights into how bacterial pathogens emerge and evolve to become new strains.

Vibrio species represent a significant portion of the culturable heterotrophic bacteria of oceans, coastal waters and estuaries[13,14]. Environmental studies show that these bacteria strongly influence nutrient cycling in the marine environment. Various species of this genus are also devastating pathogens for finfish, shellfish and mammals. There is still much to be learned about the aquatic ecology and natural history of *V. cholerae* including its autochthonous (native) existence in endemic locales during cholera-free, interepidemic periods, which environmental factors, such as climate[13,15], aided its re-emergence in Latin America, and which environmental factors are associated with its habitat in cholera endemic regions. For example, *V. cholerae*, during interepidemic periods, is an inhabitant of brackish and estuarine waters, and in these environments is associated with zooplankton and other aquatic flora and fauna[16]. The organism also enters a "viable but nonculturable"[17] state under certain conditions. Roles for these environmental interactions and this dormant physiological state in the emergence and persistence of pathogenic *V. cholerae* have been proposed[14,17].

Here we report the determination and analysis of the *Vibrio cholerae* genome sequence. This analysis represents an important step toward the complete molecular description of how this free-living environmental organism emerged to become a human pathogen by horizontal gene transfer.

Genome analysis

The genome of *V. cholerae* was sequenced by the whole genome random sequencing method[18]. The genome consists of two circular chromosomes[19,20] of 2,961,146 (chromosome 1) and 1,072,314

Table 1 General features of the *Vibrio cholerae* genome

	Chromosome 1	Chromosome 2
Size (bp)	2,961,151	1,072,914
Total number of sequences	36,797	14,367
G+C percentage	47.7	46.9
Total number of ORFs	2,770	1,115
ORF size (bp)	952	918
Percentage coding	88.6	86.3
Number of rRNA operons (16S-23S-5S)	8	0
Number of tRNA	94	4
Number similar to known proteins	1,614 (58%)	465 (42%)
Number similar to proteins of unknown function*	163 (6%)	66 (6%)
Number of conserved hypothetical proteins†	478 (17%)	165 (15%)
Number of hypothetical proteins‡	515 (19%)	419 (38%)
Number of Rho-independent terminators	599	193

bp, base pairs. ORFs, open reading frames.
* Proteins of unknown function, significant sequence similarity (homology) to a named protein for which there is currently no known function.
† Conserved hypothetical protein, sequence similarity to a translation of another ORF, but there is currently no experimental evidence a protein is expressed.
‡ Hypothetical protein, no significant sequence similarity to another protein.

(chromosome 2) base pairs, with an average G+C content of 46.9% and 47.7%, respectively. There are a total of 3,885 predicted open reading frames (ORFs) and 792 predicted Rho-independent terminators; with 2,770 and 1,115 ORFs and 599 and 193 Rho-independent terminators on the individual chromosomes (Table 1, Figs 1 and 2). Most genes required for growth and viability are located on chromosome 1, although some genes found only on chromosome 2 are also thought to be essential for normal cell function (for example, *dsdA*, *thrS* and the genes encoding ribosomal proteins L20 and L35). Additionally, many intermediaries of metabolic pathways are encoded only on chromosome 2 (Fig. 3).

The replicative origin in chromosome 1 was identified by similarity to the *Vibrio harveyi* and *Escherichia coli* origins, co-localization of genes (*dnaA*, *dnaN*, *recF* and *gyrA*) often found near the origin in prokaryotic genomes, and GC nucleotide skew (G−C/G+C) analysis[21]. Based on these, we designated base-pair 1 in an intergenic region that is located in the putative origin of replication. Only the GC skew analysis was useful in identifying a putative origin on chromosome 2.

This genomic sequence of *V. cholerae* confirmed the presence of a large integron island (a gene capture system) located on chromosome 2 (125.3 kbp)[22,23]. The *V. cholerae* integron island contains all copies of the *V. cholerae* repeat (VCR) sequence and 216 ORFs (Fig. 1). However, most of these ORFs have no homology to other sequences. Among the recognizable integron island genes are three that encode gene products that may be involved in drug resistance (chloramphenicol acetyltransferase, fosfomycin resistance protein and glutathione transferase), several DNA metabolism enzymes (MutT, transposase, and an integrase), potential virulence genes (haemagglutinin and lipoproteins) and three genes which encode gene products similar to the 'host addiction' proteins (higA, higB and doc), which are used by plasmids to select for their maintenance by host cells.

Comparative genomics

The two-chromosome structure of *V. cholerae* allows for comparisons, both between the two chromosomes of this organism and between either of the *V. cholerae* chromosomes and the chromosomes of other microbial species. There is pronounced asymmetry in the distribution of genes known to be essential for growth and virulence between the two chromosomes. Significantly more genes encoding DNA replication and repair, transcription, translation, cell-wall biosynthesis and a variety of central catabolic and biosynthetic pathways are encoded by chromosome 1. Similarly, most genes known to be essential in bacterial pathogenicity (that is, those encoding the toxin co-regulated pilus, cholera toxin, lipopolysaccharide and the extracellular protein secretion machinery) are also located on chromosome 1. In contrast, chromosome 2 contains a larger fraction (59%) of hypothetical genes and genes of unknown function, compared with chromosome 1 (42%) (Fig. 4). This partitioning of hypothetical proteins on chromosome 2 is highly localized in the integron island (Fig. 2). Chromosome 2 also carries the 3-hydroxy-3-methylglutaryl CoA reductase, a gene apparently acquired from an archaea (Y. Boucher and W. F. Doolittle, personal communication).

The majority of the *V. cholerae* genes were very similar to *E. coli* genes (1,454 ORFs), but 499 (12.8%) of the *V. cholerae* ORFs showed highest similarity to other *V. cholerae* genes, suggesting recent duplications (Figs 5 and 6). Most of the duplicated ORFs encode products involved in regulatory functions (59), chemotaxis (50), transport and binding (42), transposition (18), pathogenicity (13) or unknown functions encoded by conserved hypothetical (62) and hypothetical proteins (113). There are 105 duplications with at least one of each ORF on each chromosome indicating there have been recent crossovers between chromosomes. The extensive duplication of genes involved in scavenging behaviour (chemotaxis and solute transport) suggests the importance of these gene products in

V. cholerae biology, notably its ability to inhabit diverse environments. These environments, in turn, may have selected the duplication and divergence of genes useful for specialized functions. Additionally, whereas El Tor strain N16961 carries only a single copy of the cholera toxin prophage, other *V. cholerae* strains carry several copies of this element[24,25], and strains of the classical biotype have a second copy of the prophage that is localized on chromosome 2 (ref. 20). Thus, virulence genes are presumed to be subject to selective pressure, affecting copy number and chromosomal location.

Several ORFs with apparently identical functions exist on both chromosomes which were probably acquired by lateral gene transfer. For example, glyA (encoding serine hydroxymethyl transferase) is found once on each chromosome but the phylogenetic analysis suggest the glyA copy on chromosome 1 branches with the α-Proteobacteria, whereas the copy on chromosome 2 branches with the γ-Proteobacteria (see Supplementary Information). The chromosome 2 glyA is flanked by genes encoding transposases, suggesting that this gene was acquired through a transposition event.

Figure 1 Linear representation of the *V. cholerae* chromosomes. The location of the predicted coding regions, colour-coded by biological role, RNA genes, tRNAs, other RNAs, Rho-independent terminators and *Vibrio cholerae* repeats (VCRs) are indicated. Arrows represent the direction of transcription for each predicted coding region. Numbers next to the tRNAs represent the number of tRNAs at a locus. Numbers next to GES represent the number of membrane-spanning domains predicted by the Goldman, Engleman and Steitz scale calculated by TopPred for that protein. Gene names are available at the TIGR web site (www.tigr.org) and as Supplementary Information.
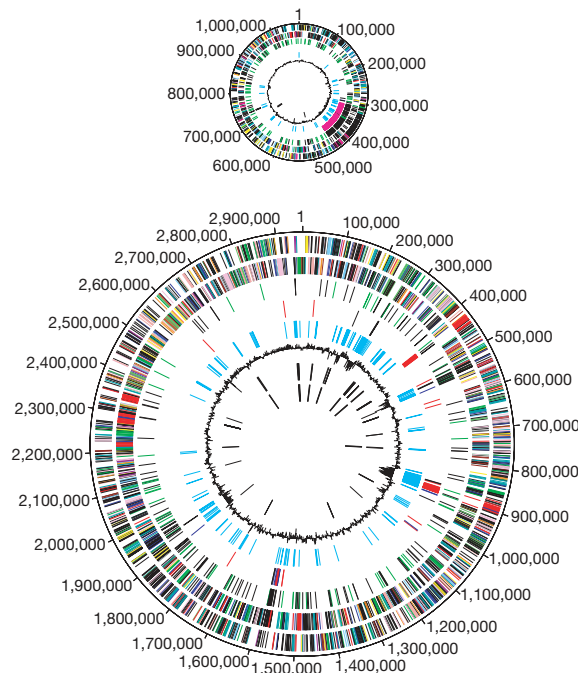

Figure 2 Circular representation of the *V. cholerae* genome. The two chromosomes, large and small, are depicted. From the outside inward: the first and second circles show predicted protein-coding regions on the plus and minus strand, by role, according to the colour code in Fig. 1 (unknown and hypothetical proteins are in black). The third circle shows recently duplicated genes on the same chromosome (black) and on different chromosomes (green). The fourth circle shows transposon-related (black), phage-related (blue), VCRs (pink) and pathogenesis genes (red). The fifth circle shows regions with significant χ^2 values for trinucleotide composition in a 2,000-bp window. The sixth circle shows percentage G+C in relation to mean G+C for the chromosome. The seventh and eighth circles are tRNAs and rRNAs, respectively.

Origin and function of the small chromosome of *V. cholerae*

Several lines of evidence suggest that chromosome 2 was originally a megaplasmid captured by an ancestral *Vibrio* species. The phylogenetic analysis of the ParA homologues located near the putative origin of replication of each chromosome shows chromosome 1 ParA tending to group with other chromosomal ParAs, and the ParA from chromosome 2 tending to group with plasmid, phage and megaplasmid ParAs (see Supplementary Information). In general, genes on chromosome 2, with an apparently identical functioning copy on chromosome 1, appear less similar to orthologues present in other γ-Proteobacteria species (see Supplementary Information). Also, chromosome 1 contains all the ribosomal RNA operons and at least one copy of all the transfer RNAs (four tRNAs are found on chromosome 2, but there are duplicates on chromosome 1). In addition, chromosome 2 carries the integron region, an element often found on plasmids[26]. Finally, the bias in the functional gene content is more easily explained, if chromosome 2

was originally a megaplasmid (Fig. 4). The megaplasmid presumably acquired genes from diverse bacterial species before its capture by the ancestral *Vibrio*. The relocation of several essential genes from chromosome 1 to the megaplasmid completed the stable capture of this smaller replicon. Apparently this capture of the megaplasmid occurred long enough ago that the trinucleotide composition and percentage G+C content between the two chromosomes has become similar (except for laterally moving elements such as the integron island, bacteriophage genomes, transposons, and so on). The two chromosome structure is found in other *Vibrio* species[19] suggesting that the gene content of the megaplasmid continues to provide *Vibrio* with an evolutionary advantage, perhaps within the aquatic ecosystem where *Vibrio* species are frequently the dominant microorganisms[14,16].

It is unclear why chromosome 2 has not been integrated into chromosome 1. Perhaps chromosome 2 plays an important specialized function that provides the evolutionary selective pressure to
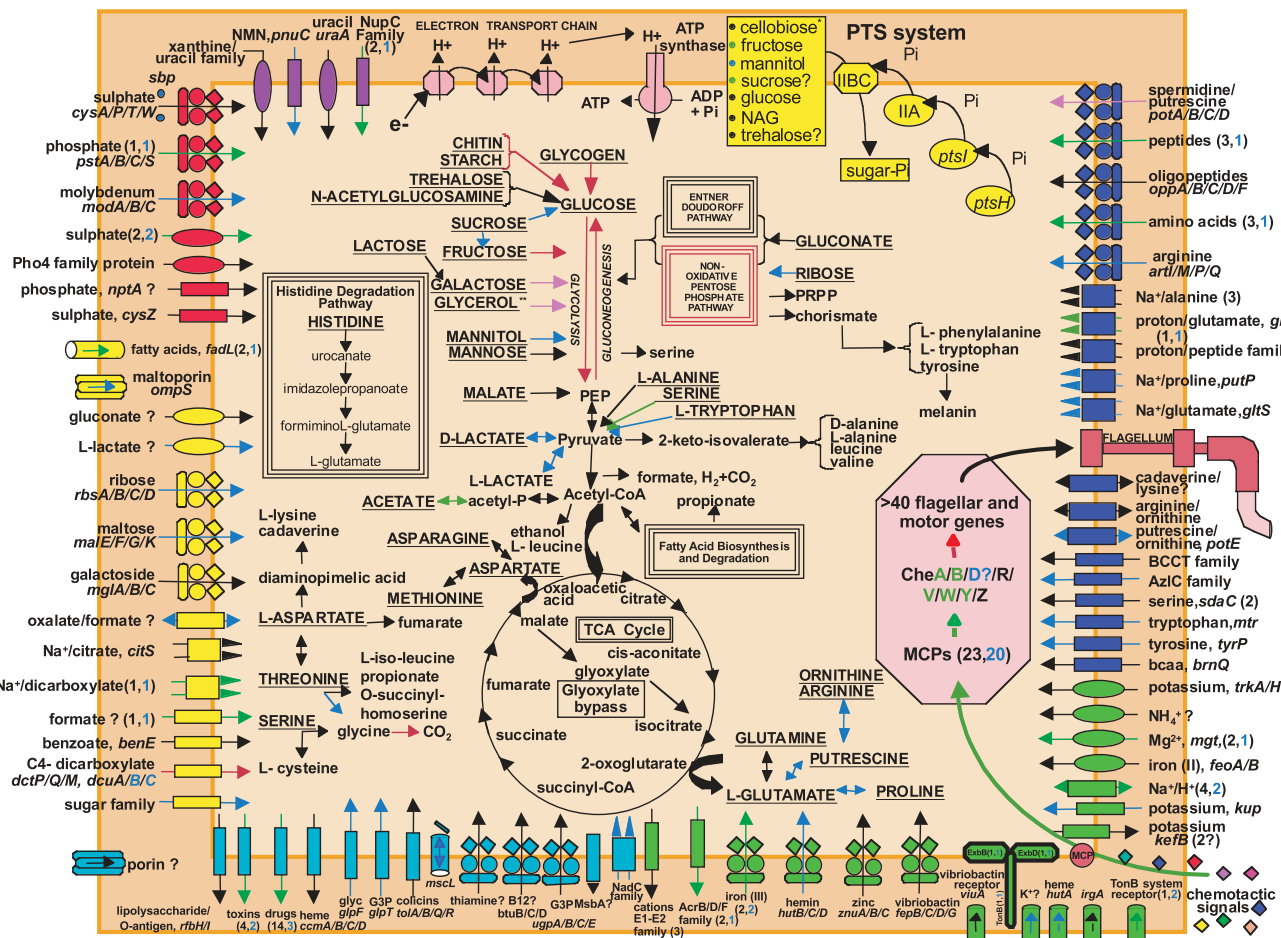

Figure 3 Overview of metabolism and transport in *V. cholerae*. Pathways for energy production and the metabolism of organic compounds, acids and aldehydes are shown. Transporters are grouped by substrate specificity: cations (green), anions (red), carbohydrates (yellow), nucleosides, purines and pyrimidines (purple), amino acids/peptides/amines (dark blue) and other (light blue). Question marks associated with transporters indicate a putative gene, uncertainty in substrate specificity, or direction of transport. Permeases are represented as ovals; ABC transporters are shown as composite figures of ovals, diamonds and circles; porins are represented as three ovals; the large-conductance mechanosensitive channel is shown as a gated cylinder; other cylinders represent outer membrane transporters or receptors; and all other transporters are drawn as rectangles. Export or import of solutes is designated by the direction of the arrow through the transporter. If a precise substrate could not be determined for a transporter, no gene name was assigned and a more general common name reflecting the type of substrate being transported was used. Gene location on the two chromosomes, for both

transporters and metabolic steps, is indicated by arrow colour: all genes located on the large chromosome (black); all genes located on the small chromosome (blue); all genes needed for the complete pathway on one chromosome, but a duplicate copy of one or more genes on the other chromosome (purple); required genes on both chromosomes (red); complete pathway on both chromosomes (green). (Complete pathways, except for glycerol, are found on the large chromosome.) Gene numbers on the two chromosomes are in parentheses and follow the colour scheme for gene location. Substrates underlined and capitalized can be used as energy sources. PRPP, phosphoribosyl-pyrophosphate; PEP, phosphoenolpyruvate; PTS, phosphoenolpyruvate-dependant phosphotransferase system; ATP, adenosine triphosphate; ADP, adenosine diphosphate; MCP, methyl-accepting chaemotaxis protein; NAG, *N*-acetylglucosamine; G3P, glycerol-3-phosphate; glyc, glycerol; NMN, nicotinamide mononucleotide. Asterisk, because *V. cholerae* does not use cellobiose, we expect this PTS system to be involved in chitobiose transport.

suppress integration events when they do occur. For example, if under some environmental condition there is a difference in copy number between the chromosomes, then chromosome 2 may have accumulated genes that are better expressed at higher or lower copy number than genes on chromosome 1. A second possibility is that, in response to environmental cues, one chromosome may partition to daughter cells in the absence of the other chromosome (aberrant segregation). Such single-chromosome-containing cells would be replication-defective but still maintain metabolic activity ('drone' cells), and, therefore, be a potential source of "viable, but non-culturable (VBNC)" cells observed to occur in *V. cholerae*[17]. Such 'drone' cells may also play a role in *V. cholerae* biofilms[7,27,28] by, for example, producing extracellular chitinase, protease and other degradative enzymes that enhance survival of cells in a biofilm, retaining two chromosomes without directly competing with these cells for nutrients.

Transport and energy metabolism

Vibrio cholerae has a diverse natural habitat that includes association with zooplankton in a sessile stage, a planktonic state in the water column, and the capacity to act as a pathogen within the human gastrointestinal tract. It is, therefore, no surprise that this organism maintains a large repertoire of transport proteins with broad substrate specificity and the corresponding catabolic pathways to enable it to respond efficiently to these different and constantly changing ecosystems (Fig. 4). Many of the sugar transporter systems and their corresponding catabolic pathway enzymes are localized on a single chromosome (that is, ribose and lactate transport and degradation enzymes are contained on chromosome 2, whereas the trehalose systems reside on chromosome 1). However, many of the other energy metabolism pathways are split (that is, chitin, glyco-lysis, and so on) between the chromosomes (Fig. 3).

In aquatic environments, chitin often represents a source of both carbon and nitrogen. This energy source is important for *V. cholerae* as it is associated with zooplankton, which have a chitinous exoskeleton[13,15,29]. *Vibrio cholerae* degrades chitin by a pathway that is very similar to that of *Vibrio furnissii*[30]. Sequence analysis suggests a phosphenolpyruvate phosphotransferase system (PTS)

for cellobiose transport, but as *V. cholerae* does not use cellobiose, it is more likely that this PTS is involved in transport of the structurally similar compound, chitobiose, analogous to the situation proposed for *Bourrelia burgdorferi*[18].

The three anions that are transported by ABC transport systems in *V. cholerae* are molybdenum, phosphate and sulphate. Molybdenum transport genes (*modA/B/C*) are all located on chromosome 2, and most of the sulphate transport genes are on the large molecule. However, copies of the genes for phosphate transporters are found in both chromosomes. The genes in these two phosphate transport operons are different from each other and do not represent a recent duplication; instead, this suggests that one may be an acquired operon.

Interchromosomal regulation

Several of the regulatory pathways, both for regulation in response to environmental and pathogenic signals, are divided between the two chromosomes. These included pathways for starvation survival, 'quorum sensing' and expression of the entertoxigenic haemolysin, HlyA.

During periods of nutrient starvation, *V. cholerae*, and other Gram-negative bacteria, enter the stationary phase and, later, the viable but nonculturable (VBNC) state[14,17]. The alternative sigma factor σ^{38} (*rpoS*) is required for survival of *V. cholerae* in the environment but not for pathogenicity[31], and therefore probably plays an important role in the initiation of the VBNC state. There is one copy of *rpoS*, located on chromosome 1, near the oriC. The RpoS regulates expression of several other proteins, including catalase, cyclopropane-fatty-acyl-phospholipid synthase and HA/protease, which are found on both chromosomes[31].

Genes involved in 'quorum sensing', or cell-density-dependent regulation, also exist on both chromosomes of *V. cholerae*. In bioluminescent *Vibrio* species (notably *Vibrio fischeri* and *V. harveyi*), quorum sensing is used to control light production. Although this strain of *V. cholerae* lacks the genes for bioluminescence, it does have the genes required for the autoinducer-2 (AI-2) quorum-sensing mechanism[32] (*luxOPQSU*) but this pathway is split between the chromosomes with *luxOSU* on chromosome 1 and *luxPQ* on chromosome 2. Similarly, another transcriptional regulatory gene,
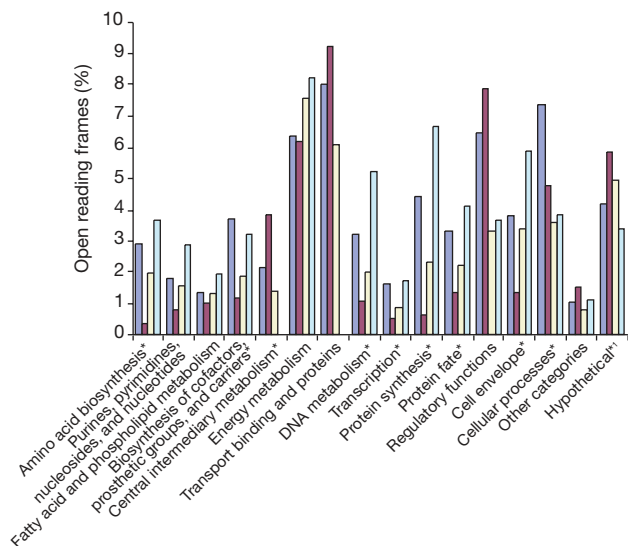

Figure 4 Percentage of total *Vibrio cholerae* open reading frames (ORFs) in biological roles compared with other γ-Proteobacteria. These were *V. cholerae*, chromosome 1 (blue); *V. cholerae*, chromosome 2 (red); *Escherichia coli* (yellow); *Haemophilus influenzae* (pale blue). Significant partitioning ($P < 0.01$) of biological roles between *V. cholerae* chromosomes is indicated with an asterisk, as determined with a χ^2 analysis. 1, Hypothetical contains both conserved hypothetical proteins and hypothetical proteins, and is at 1/10 scale compared with other roles.
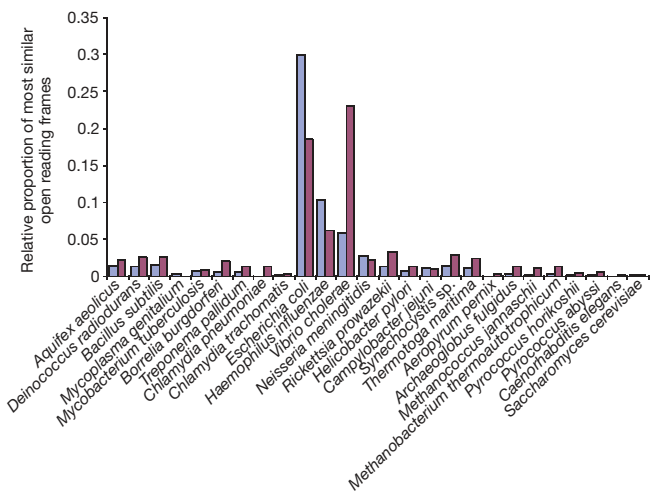
Figure 5 Comparison of the *V. cholerae* ORFs with those of other completely sequenced genomes. The sequence of all proteins from each completed genome were retrieved from NCBI, TIGR and the *Caenorhabditis elegans* (wormpep16) databases. All *V. cholerae* ORFs (large chromosome, blue; small chromosome, red) were searched against all other genomes with FASTA3. The number of *V. cholerae* ORFs with greatest similarity ($E \le 10^{-5}$) are shown in proportion to the total number of ORFs in that genome. There were no ORFs that were most similar to a *Mycoplasma pneumoniae* ORF.
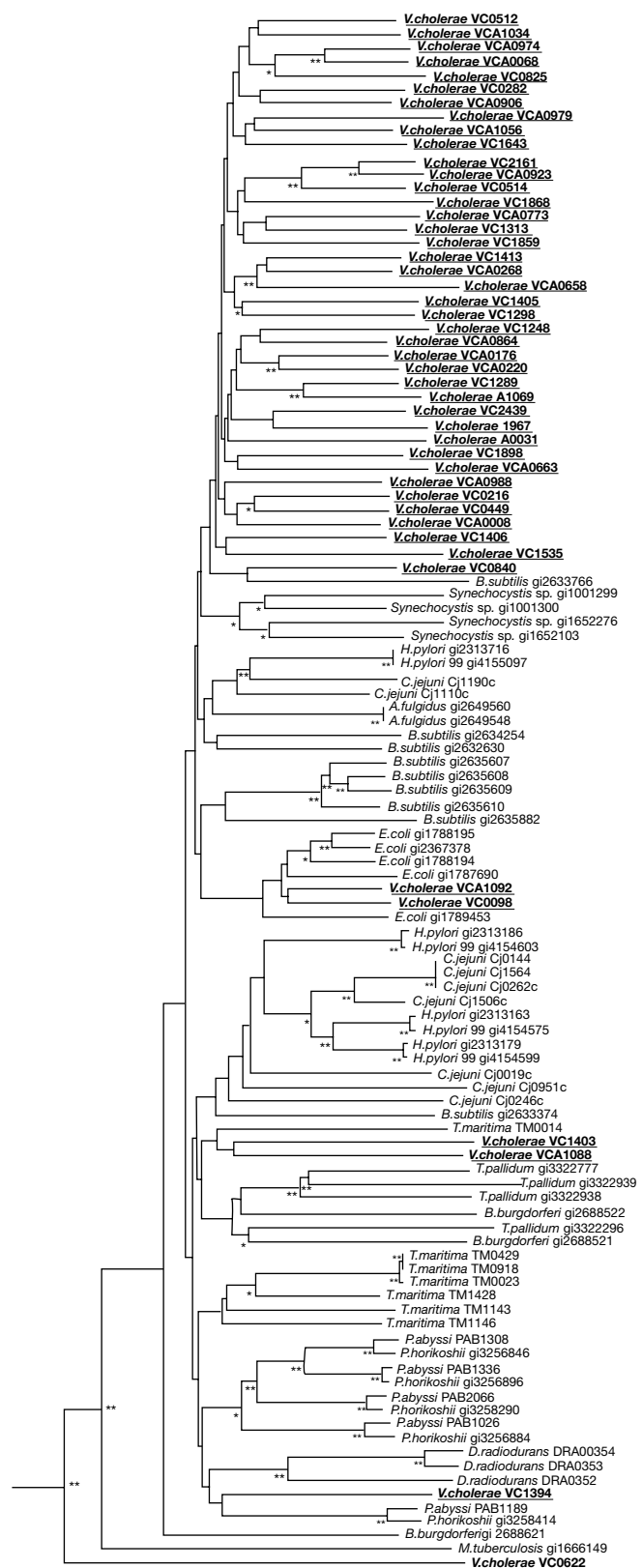
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Figure 6 Phylogenetic tree of methyl-accepting chemotactic proteins (MCP) homologues in completed genomes. Homologues of MCP were identified by FASTA3 searches of all available complete genomes. Amino-acid sequences of the proteins were aligned using CLUSTALW, and a neighbour-joining phylogenetic tree was generated from the alignment using the PAUP* program (using a PAM-based distance calculation). Hypervariable regions of the alignment and positions with gaps in many of the sequences were excluded from the analysis. Nodes with significant bootstrap values are indicated: two asterisks, >70%; asterisk, 40–70%.

hlyU (ref. 33), is located on chromosome 1, while the gene it regulates, hlyA, is located on chromosome 2.

DNA repair

Vibrio cholerae has genes encoding several DNA-repair and DNA-damage-response pathways, including nucleotide-excision repair, mismatch-excision repair, base-excision repair, AP endonuclease, alkylation transfer, photoreactivation, DNA ligation, and all the major components of recombination and recombinational repair, including initiation, recombination and resolution[34]. In addition, homologues of many of the genes involved in the SOS response in *E. coli* are found. The presence of three photolyase homologues, more than have been found in other bacterial species, probably allows for the ability to photoreactivate the two major forms of ultraviolet-induced DNA damage (cyclobutane pyrimidine dimers and 6-4 photoproducts), and may also allow use of a range of wavelengths of light used for the energy required for photoreactivation. It is also of interest that many of the repair genes are on chromosome 2 (*alkA, ada1, ada2, phr3, mutK, sbcD, dcm, mutT3*), indicating that this chromosome is probably required for full DNA repair capability.

Pathogenicity

Toxins. The genome sequence of *V. cholerae* El Tor N16961 revealed a single copy of the cholera toxin (CT) genes, *ctxAB*, located on chromosome 1 within the integrated genome of CTXϕ, a temperate filamentous phage[3]. The receptor for entry of CTXϕ into the cell is the toxin-coregulated pilus (TCP)[3], and the TCP gene cluster (see below) also resides on chromosome 1. Like the structural genes for CT and TCP, the regulatory gene, *toxR*, which controls their expression *in vivo*[35], is also located on chromosome 1.

On the other side of CTXϕ prophage is a region encoding an RTX toxin (*rtxA*), and its activator (*rtxC*) and transporters (*rtxBD*)[36]. A third transporter gene has been identified that is a paralogue of *rtxB*, and is transcribed in the same direction as *rtxBD*. Downstream of this gene are two genes encoding a sensor histidine kinase and response regulator. Trinucleotide composition analysis suggests that the RTX region was horizontally acquired along with the sensor histidine kinase/response regulator, suggesting these regulators effect expression of the closely linked RTX transcriptional units.

Also present are genes encoding numerous potential toxins, including several haemolysins, proteases and lipases. These include *hap*, the haemagluttinin protease, a secreted metalloprotease that seems to attack proteins involved in maintaining the integrity of epithelial cell tight junctions[37], and *hlyA*, encoding a secreted haemolysin that displays enterotoxic activity[38]. In contrast to CTX, RTX and all known intestinal colonization factors, the *hap* and *hlyA* genes virulence factors reside on chromosome 2.

Vibrio cholerae has been reported to produce shiga-like toxins[39]; however, the sequence did not reveal genes encoding specific homologues of the A or B subunits of shiga toxin. Also not detected were genes encoding homologues of *E. coli* heat stable toxin (ST), which have been detected in other pathogenic strains of *V. cholerae*[40].

Colonization factors. The critical intestinal colonization factor of *V. cholerae* is the TCP, a type IV pilus[5,41]. The genome sequence confirmed that the genes involved in TCP assembly (*tcpABCDEFGHIJNQRST*) reside on chromosome 1 (ref. 20) as part of a proposed 'pathogenicity island' (also referred to as VPI) composed of recently acquired DNA that encodes not only TCP, but also other genes associated with the ToxR regulatory cascade, such as *acfABCD, toxT, aldA* and *tagAB*[10,11]. Trinucleotide composition analysis suggest that this 45.3-kb segment begins at a 20-bp site upstream of *aldA*, and encompasses a helicase-related protein and a transcriptional activator which both share homology with bacteriophage proteins. At the other end of the segment of atypical trinucleotide composition is a phage family integrase and the

other copy of the 20-bp site, which is presumably the target for integration of the island onto the chromosome[10,11]. It has been proposed that the TCP/ACF island corresponds to the genome of a filamentous phage that uses TCP pilin as a coat protein[4]. However, other than the three genes encoding phage-related proteins (that is, the helicase, transcriptional activator and integrase) we could find no other genes on the island that encoded products with significant homology to the conserved gene products of other filamentous phages or the structural proteins of nonfilamentous phages.

The maltose-sensitive haemagglutinin (MSHA) is unique to the El Tor biotype of *V. cholerae*. Initially characterized as a haemagglutinin, it was later found to be a type IV pilus[42,43]. The MSHA biogenesis (MshHIJKLMNEGF) and structural (MshBACD) proteins are all clustered on chromosome 1. There are no apparent integrases or transposases that might define this region as a pathogenicity island or suggest an origin for it other than *V. cholerae*. In support of this conclusion, trinucleotide composition analysis shows that this region has similar composition to the rest of the chromosome, suggesting that if these genes were acquired it was very early in the *Vibrio* phylogenetic history. Recently, several investigators have reported that MSHA is not required for intestinal colonization, nor does it seem to appreciably affect the efficiency of colonization[44–46], but instead plays a role in biofilm formation[27,28]. Accordingly, this pilus may be important for the environmental fitness of *Vibrio* species rather than for pathogenic potential.

The *pilA* region of *V. cholerae* genome apparently encodes a third type IV pilus, although it has not been visualized[47]. This gene cluster includes a gene encoding a prepilin peptidase (PilD) that is required for the efficient processing of protein complexes with type IV prepilin-like signal sequences including TCP, MSHA and EPS[47,48]. The EPS system of *V. cholerae* encodes a type II secretion system involved in extracellular export of CT and other proteins. The EPS system is encoded by chromosome 1 but, like the MSHA genes, trinucleotide analysis suggests that the EPS genes of *V. cholerae* have not been recently acquired. In contrast, trinucleotide composition analysis suggests that the *pilA* gene cluster was acquired by horizontal transfer. Thus, analysis of the *V. cholerae* genome sequence provides some evidence that older gene clusters, like MSHA and EPS, have become dependent on newly acquired genes such as PilD.

Conclusions

The *Vibrio cholerae* genome sequence provides a new starting point for the study of this organism's environmental and pathobiological characteristics. It will be interesting to determine the gene expression patterns that are unique to its survival and replication during human infection[35] as well as in the environment[13,14,16]. Additionally, the genomic sequence of *V. cholerae* should facilitate the study of this model multi-chromosomal prokaryotic organism. Comparative genomics between several species in the genus *Vibrio* will provide a better understanding of the origin of the new small chromosome and the role that it plays in *Vibrio* biology. The genome sequence may also provide important clues to understanding the metabolic and regulatory networks that link genes on the two chromosomes. Finally, *V. cholerae* clearly represents a promising genetic system for studying how several horizontally acquired loci located on separate chromosomes can still efficiently interact at the regulatory, cell biology and biochemical levels. □

Methods

Whole-genome random sequencing procedure.

Vibrio cholerae N16961 was grown from a single isolated colony. Cloning, sequencing and assembly were as described for genomes sequenced by TIGR[18]. One small-insert plasmid library (2–3 kb) was generated by random mechanical shearing of genomic DNA. One large insert library was ligated into λ-DASHII/EcoRI vector (Stratagene). In the initial sequence phase, approximately sevenfold sequence coverage was achieved with 49,633 sequences from plasmid clones. Sequences from both ends of 383 λ-clones served as a genome scaffold, verifying the orientation, order and integrity of the contigs. The plasmid and λ sequences were jointly assembled using TIGR Assembler. Sequence gaps were closed by editing the end sequences and/or primer walking on plasmid clones. Physical gaps were closed by direct sequencing of genomic DNA, or combinatorial polymerase chain reaction (PCR) followed by sequencing the PCR product. The final genome sequence is based on 51,164 sequences.

ORF prediction and gene family identification. An initial set of ORFs, likely to encode proteins, was identified with GLIMMER[49], and those shorter than 30 codons were eliminated. ORFs that overlapped were visually inspected, and in some cases removed. ORFs were searched against a non-redundant protein database[18]. Frameshifts and point mutations were detected and corrected where appropriate. Remaining frameshifts and point mutations are considered to be authentic and were annotated as 'authentic frameshift' or 'authentic point mutation'. ORFs were also analysed with two sets of hidden Markov models (HMMs) constructed for a number of conserved protein families (1,313 from Pfam v3.1 (ref. 50) and 476 from the TIGRFAM) by use of the HMMER package. TopPred was used to identify membrane-spanning domains in proteins.

Paralogous gene families were constructed by searching the ORFs against themselves using BLASTX, identifying matches with $E \leq 10^{-5}$ over 60% of the query search length, and subsequently clustering these matches into multigene families. Multiple alignments for these protein families were generated with the CLUSTALW program and the alignments scrutinized.

Distribution of all 64 trinucleotides (3-mers) for each chromosome was determined, and the 3-mer distribution in 2,000-bp windows that overlapped by half their length (1,000 bp) across the genome was computed. For each window, we computed the χ^2 statistic on the difference between its 3-mer content and that of the whole chromosome. A large value of this statistic indicates that the 3-mer composition in this window is different from the rest of the chromosome. Probability values for this analysis are based on the assumption that the DNA composition is relatively uniform throughout the genome. Because this assumption may be incorrect, we prefer to interpret high χ^2 values merely as indicators of regions on the chromosome that appear unusual and demand further scrutiny.

Homologues of the genes of interest were identified using the BLASTP and FASTA3 search programs. All homologues were then aligned to each other using the CLUSTALW program with default settings. Phylogenetic trees were generated from the alignments using the neighbour-joining algorithm as implemented by the PAUP* program (with a PAM matrix based distance calculation). Regions of the alignment that were hypervariable or were of low confidence were excluded from the phylogenetic analysis. All alignments are available upon request.

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