Permanent draft genome sequence of Vibrio tubiashii strain NCIMB 1337 (ATCC19106)

Citation for published version:
Temperton, B, Thomas, S, Tait, K, Parry, H, Emery, M, Allen, M, Quinn, J, Macgrath, J & Gilbert, J 2011, 'Permanent draft genome sequence of Vibrio tubiashii strain NCIMB 1337 (ATCC19106)' Standards in genomic sciences, vol 4, no. 2, pp. 183-90. DOI: 10.4056/sigs.1654066

Digital Object Identifier (DOI):
10.4056/sigs.1654066

Link:
Link to publication record in Edinburgh Research Explorer

Document Version:
Publisher's PDF, also known as Version of record

Published In:
Standards in genomic sciences

Publisher Rights Statement:
Free via PMC.

General rights
Copyright for the publications made accessible via the Edinburgh Research Explorer is retained by the author(s) and / or other copyright owners and it is a condition of accessing these publications that users recognise and abide by the legal requirements associated with these rights.

Take down policy
The University of Edinburgh has made every reasonable effort to ensure that Edinburgh Research Explorer content complies with UK legislation. If you believe that the public display of this file breaches copyright please contact openaccess@ed.ac.uk providing details, and we will remove access to the work immediately and investigate your claim.
Permanent draft genome sequence of Vibrio tubiashii strain NCIMB 1337 (ATCC19106)

Ben Temperton1,2 and Simon Thomas1,2, Karen Tait1, Helen Parry1, Matt Emery3, Mike Allen1, John Quinn2, John MacGrath2, Jack Gilbert1,4,5

1 Plymouth Marine Laboratory, Prospect Place, The Hoe, Plymouth, UK
2 Queen’s University Belfast, School of Biological Sciences, Medical Biology Centre, Belfast, Northern Ireland
3 University of Plymouth, Department of Microbiology, Drakes Circus, Plymouth
4 Argonne National Laboratory, Argonne, IL, USA
5 Department of Ecology and Evolution, University of Chicago, Chicago, IL, USA

Vibrio tubiashii NCIMB 1337 is a major and increasingly prevalent pathogen of bivalve mollusks, and shares a close phylogenetic relationship with both V. orientalis and V. coralliilyticus. It is a Gram-negative, curved rod-shaped bacterium, originally isolated from a moribund juvenile oyster, and is both oxidase and catalase positive. It is capable of growth under both aerobic and anaerobic conditions. Here we describe the features of this organism, together with the draft genome and annotation. The genome is 5,353,266 bp long, consisting of two chromosomes, and contains 4,864 protein-coding and 86 RNA genes.

Introduction
The genus Vibrio is both numerous and ubiquitous within marine environments, with Vibrio species harbored within many diverse marine organisms, such as mollusks, shrimps, fishes, cephalopods and corals [1]. Comparative genome analysis has revealed a huge genetic diversity within this genus, which is driven by mutations, chromosomal rearrangements, loss of genes by decay or deletion, and gene acquisitions through duplication or horizontal transfer (e.g. the acquisition of bacteriophages, pathogenicity islands, and super-integrons), the combination of which presumably stimulates genetic and functional diversity and allows this group to colonize a wide variety of ecological niches and hosts [1,2].

Vibrio tubiashii was first described as three strains of Vibrio anguillarum by Tubiash et al [3] in 1965. The organisms were isolated from bivalve mollusks during an outbreak of bacillary necrosis in Milford, Connecticut, and deposited in the American Type Culture Collection as ATCC 19105, 19106 and 19109. These three strains were further elucidated and formally named as V. tubiashii by Had a et al [4] in 1984. Subsequently, several virulence factors have been identified [5,6] and the organism is increasingly implicated in major disease outbreaks in bivalve mollusks [1].

V. tubiashii is closely related to the proposed coral pathogen V. coralliilyticus, as well as V. orientalis, a bacterium associated with penaeid shrimps [7]. Indeed, V. coralliilyticus was initially designated as a V. tubiashii strain [8,9] due to their close similarity.

Classification and features
Vibrio tubiashii 1337 belongs to the Gammaproteobacteria and are contained within the family, Vibrionaceae [Table 1]. Cells of Vibrio tubiashii are Gram-negative curved-rods of approximately 0.5 by 1.5 µm, which are motile in liquid media by means of a single sheathed, polar flagellum [3,4]. These cells are facultative anaerobes, [3,4,22]. It is catalase and oxidase positive, capable of splitting indole from tryptophan, and can use glucose, xylose, mannitol, rhamnose, sucrose, arabinose and acetate as sole carbon sources, and has β-galactosidase activity, despite an apparent inability to ferment lactose. V. tubiashii is capable of dissimilatory nitrate and nitrite reduction under anaerobic conditions, can use organic phosphorus during phosphate limitation, and can utilize 2-aminoethylphosphonate as a sole phosphorus source.
**Vibrio tubiashii** strain NCIMB 1337 (ATCC19106)

*V. tubiashii* has an absolute requirement for sodium and chloride ions, and is incapable of growth on media containing less than 0.5% W/V NaCl. The temperature optimum for growth is 25°C, but growth does occur in the range of 12-30°C. The organism is killed at 37°C. *V. tubiashii* has a biphasic pH response and grows optimally at both pH 8.0 and 6.5, but displays weakened growth at pH 7.0 and 7.5. The bacterium shows rapid growth on marine broth and produces buff colored, opaque, irregular, slightly convex colonies on marine agar, and yellow colonies, characteristic of the *Vibrionaceae*, on Thiosulfate-Citrate-Bile-Sucrose Agar (TCBS).

**Growth conditions and DNA isolation**

*Vibrio tubiashii* NCIMB 1337 (ATCC19106) was grown in marine broth (seawater + 1 gl⁻¹ yeast extract and 0.5 gl⁻¹ tryptone) at 25°C for 24 hours. DNA was extracted using the Qiagen DNAeasy blood and tissue kit, without modification of the manufacturer’s protocol.

**Genome sequencing and annotation**

**Genome sequencing**

The genome was sequenced using the Illumina sequencing platform. All general aspects of library construction and sequencing performed at the NERC Biomolecular analysis facility can be found on the NBAF website [23]. SOLEXA Illumina reads were assembled using VELVET Large Newbler contigs that were broken into 4,074 overlapping fragments of 1,000 bp and entered into the assembly as pseudo-reads. The sequences were assigned quality scores based on consensus q-scores with modifications to account for overlap redundancy and to adjust inflated q-scores. The error rate of the completed genome sequence is less than 1 in 100,000. Overall sequencing provided 131 × coverage of the genome.

**Genome annotation**

Genes were identified using the RAST server. The predicted CDSs were translated and used to search the National Center for Biotechnology Information (NCBI) nonredundant database, UniProt, TIGRFam, Pfam, PRIAM, KEGG, COG, and InterPro databases. The tRNAscanSE tool [24] was used to find tRNA genes, whereas ribosomal RNAs were found by using BLASTn against the ribosomal RNA databases. The RNA components of the protein secretion complex and the RNaseP were identified by searching the genome for the corresponding Rfam profiles using INFERNAL [25]. Additional gene prediction analysis and manual functional annotation was performed within the Integrated Microbial Genomes (IMG) platform developed by the Joint Genome Institute, Walnut Creek, CA, USA [26,27].

![Figure 1](image_url)

**Figure 1.** Phylogenetic tree highlighting the position of *V. tubiashii* NCIMB 1337 relative to other Vibrio strains. The tree was inferred from 1,159 aligned characters of the 16S rRNA gene sequence under the neighborhood joining criterion. Numbers above the branches are support values from 1,000 bootstrap replicates if greater than 60%.
Table 1. Classification and general features of *V. tubiashii* according to the MIGS recommendations

| MIGS ID | Property                  | Term                                | Evidence code |
|---------|---------------------------|-------------------------------------|---------------|
|         | Domain                    | Bacteria                            | TAS [10]      |
|         | Phylum                    | Proteobacteria                      | TAS [11]      |
|         | Class                     | Gammaproteobacteria                 | TAS [12,13]   |
| Current classification | Order                     | Vibrionales                         | TAS [14]      |
|         | Family                    | Vibrionaceae                        | TAS [15,16]   |
|         | Genus                     | Vibrio                              | TAS [15,17-19]|
|         | Species                   | *Vibrio tubiashii* NCIMB 1337       | TAS [4]       |
|         | Gram stain                | negative                            | IDA           |
|         | Cell shape                | Curved rods (vibroid)               | IDA           |
|         | Motility                  | motile via single polar flagellum   | IDA           |
|         | Sporulation               | Non-sporulating                     | IDA           |
|         | Temperature range          | Mesophile 12-30°C                   | IDA           |
|         | Optimum temperature       | 25°C                                | IDA           |
| MIGS 6.3 | Salinity               | Slightly halophytic, optimum 1-3% NaCl | IDA           |
| MIGS-22 | Oxygen requirement       | Aerobic/ facultative anaerobic      | IDA           |
|         | Carbon source             | Highly diverse                      | IDA           |
|         | Energy source             | Highly diverse                      | IDA           |
| MIGS-6  | Habitat                   | Marine invertebrates                | TAS [20]      |
| MIGS-16 | Biotic relationship       | Parassitico                         | TAS [3]       |
| MIGS-14 | Biosafety level           | 2                                   | TAS [4]       |
|         | Isolation                 | Moribund juvenile oyster (*Crassostrea virginica*) | TAS [3,4]  |
| MIGS-4  | Geographical location     | Milford, Connecticut, USA           | TAS [3]       |
| MIGS-5  | Sample collection time    | 01/02/1965                          | TAS [3]       |
| MIGS 4.1| latitude                  | 41.22 N                             | TAS [3]       |
| MIGS 4.2| longitude                 | -73.06 W                            | TAS [3]       |
| MIGS 4.3| Depth                    | Not reported                         |               |
| MIGS 4.4| Altitude                 | Marine                              | TAS [3]       |

Evidence codes - IDA: Inferred from Direct Assay (first time in publication); TAS: Traceable Author Statement (i.e., a direct report exists in the literature); NAS: Non-traceable Author Statement (i.e., not directly observed for the living, isolated sample, but based on a generally accepted property for the species, or anecdotal evidence). These evidence codes are from the Gene Ontology project [21]. If the evidence code is IDA, then the property was directly observed, for a live isolate by one of the authors, or an expert or reputable institution mentioned in the acknowledgements.

**Genome project information**

This organism was selected for sequencing on the basis of its increasing impact as a bivalve pathogen, and was funded by i-G Peninsula. The genome project is deposited in the IMG database and the complete genome sequence in GenBank (CP001643). Sequencing, finishing and annotation were performed by the GenePool Team at NERC Biomolecular Analysis Facility (NBAF) Edinburgh. A summary of the project information is shown in Table 2.

**Genomic properties**

The genome was assembled into 335 contigs and includes two circular chromosomes combining to give a total size of 5,353,266 bp (44.84% GC content). A total of 4,950 genes were predicted, 4,864 of which are protein-coding genes. 74.22% of protein coding genes were assigned to a putative function with the remaining annotated as hypothetical proteins. 658 protein coding genes belong to paralogous families in this genome corresponding to a gene content redundancy of 13.29%. The properties and the statistics of the genome are summarized in Tables 3-5.
Table 2. Project information

| MIGS ID   | Property               | Term                        |
|-----------|------------------------|-----------------------------|
| MIGS-31   | Finishing quality      | Draft                       |
| MIGS-28   | Libraries used         | Illumina                    |
| MIGS-29   | Sequencing platforms   | Illumina SOLEXA GAIIx       |
| MIGS-31.2 | Fold coverage          | 131×                        |
| MIGS-30   | Assemblers             | Velvet                      |
| MIGS-32   | Gene calling method    | RAST                        |
|           | Genome Database release| 181                         |
|           | Genbank ID             | 866909                      |
|           | Genbank Date of Release| December 12, 2010           |
|           | GOLD ID                | Gi07317                     |

Table 3. Summary of genome*

| Label          | Size (Mb) |
|----------------|-----------|
| Chromosome 1   | 3.4       |
| Chromosome 2   | 1.9       |

*Two chromosomes with no plasmids. Approximate chromosome size estimated by Pulse field gel electrophoresis

Table 4. Nucleotide content and gene count levels of the genome

| Attribute                  | Value     | % of total\(^a\) |
|----------------------------|-----------|------------------|
| Size (bp)                  | 5,353,266 | 100%             |
| G+C content (bp)           | 2,400,750 | 44.87%           |
| Coding region (bp)         | 4,627,782 | 86.45%           |
| Total genes\(^b\)          | 4950      | 100%             |
| RNA genes                  | 86        | 1.74%            |
| Protein-coding genes       | 4864      | 98.26%           |
| Genes in paralog clusters  | 658       | 13.29%           |
| Genes assigned to COGs     | 3674      | 74.22%           |
| Genes with signal peptides | 1655      | 33.43%           |
| Genes with transmembrane helices | 1167 | 23.58% |
| Paralogous groups          | 658       | 13.29%           |

\(^a\)The total is based on either the size of the genome in base pairs or the total number of protein coding genes in the annotated genome.

\(^b\)Also includes 54 pseudogenes and 5 other genes.
### Table 5. Number of genes associated with the 25 general COG functional categories

| Code | Value | % Value | Description |
|------|-------|---------|-------------|
| J    | 200   | 4.86    | Translation |
| A    | 1     | 0.02    | RNA processing and modification |
| K    | 369   | 8.96    | Transcription |
| L    | 154   | 3.74    | Replication, recombination and repair |
| B    | 1     | 0.02    | Chromatin structure and dynamics |
| D    | 37    | 0.9     | Cell cycle control, mitosis and chromosome partitioning |
| Y    |       |         | Nuclear structure |
| V    | 75    | 1.82    | Defense mechanisms |
| T    | 432   | 8.31    | Signal transduction mechanisms |
| M    | 227   | 5.51    | Cell wall/membrane biogenesis |
| N    | 148   | 3.59    | Cell motility |
| U    | 146   | 3.55    | Intracellular trafficking and secretion |
| O    | 173   | 4.2     | Posttranslational modification, protein turnover, chaperones |
| C    | 203   | 4.93    | Energy production and conversion |
| G    | 248   | 6.02    | Carbohydrate transport and metabolism |
| E    | 348   | 8.45    | Amino acid transport and metabolism |
| F    | 105   | 2.55    | Nucleotide transport and metabolism |
| H    | 159   | 3.86    | Coenzyme transport and metabolism |
| I    | 119   | 2.89    | Lipid transport and metabolism |
| P    | 188   | 4.57    | Inorganic ion transport and metabolism |
| Q    | 77    | 1.77    | Secondary metabolites biosynthesis, transport and catabolism |
| R    | 445   | 10.81   | General function prediction only |
| S    | 356   | 8.65    | Function unknown |
| -    | 1276  | 25.78   | Not in COGs |

a) The total is based on the total number of protein coding genes in the annotated genome.

### Genomic comparison

Based on COG I.D the *Vibrio tubiashii* genome shows most similarity to the genome of *V. coralliilyticus* (R² = 0.96) and to *V. orientalis* (R² = 0.94), while showing less similarity to *V. shilonii* (R²= 0.86) [Table 6]. This is in contrast to the 16S-based analysis shown in Figure 1. However, it should be noted that 16S rRNA analysis often poorly discriminates vibrios due to low sequence heterogeneity in the 16S gene [28].

### Regulatory systems

The *Vibrio tubiashii* NCIMB 1337 genome contains multiple quorum sensing systems, most notably a luxM/N system which has two adjacent copies of the luxN gene. In addition, there is a luxS/PQ system, with the lux P and Q gene appearing consecutively. There is also a cqsA/S system. It is probable that these three systems converge on the phospho-relay transfer system encoded by the luxO/luxU/hapR genes. There are two additional lux genes (LuxT and LuxZ). The genome also contains the rpoN gene encoding for the sigma-54 factor, which may indicate the presence of the two-component phosphorylation-dephosphorylation cascade described in *V. harveyi* [29] (note: *Vibrio harveyi* is also known as *Lucibacterium harveyi* and *Beneckea harveyi*).
**Vibrio tubiashii** strain NCIMB 1337 (ATCC19106)

Table 6. Comparison of the genome of **Vibrio tubiashii** NCIMB 1337 with other sequenced Vibrios

| Genome Name | **Vibrio coralliilyticus** ATCC BAA-450 | **Vibrio orientalis** CIP 102891 | **Vibrio shilonii** AK1 | **Vibrio tubiashii** NCIMB 1337 |
|-------------|-----------------------------------------|---------------------------------|----------------------|-----------------------------|
| Genes       | 5,144                                   | 4,297                           | 5,438                | 4,950                       |
| RNA         | 122                                     | 128                             | 78                   | 86                          |
| w/ Func Pred| 3,687                                   | 3,185                           | 3,517                | 4,062                       |
| w/ Func Pred %| 71.68%                                  | 74.12%                          | 64.67%               | 82.06%                      |
| Enzymes     | 1,143                                   | 1,058                           | 1,258                | 1,116                       |
| Enzymes %   | 22.22%                                  | 24.62%                          | 23.13%               | 22.55%                      |
| KEGG        | 1397                                    | 1,257                           | 1,511                | 1,354                       |
| KEGG %      | 27.16%                                  | 29.25%                          | 27.79%               | 27.35%                      |
| COG         | 3815                                    | 3,302                           | 4,093                | 3,674                       |
| COG %       | 74.16%                                  | 76.84%                          | 75.27%               | 74.22%                      |
| Pfam        | 4127                                    | 3,520                           | 4,379                | 3,976                       |
| Pfam %      | 80.23%                                  | 81.92%                          | 80.53%               | 80.32%                      |
| TIGRfam     | 1,643                                   | 1,515                           | 1,708                | 1,651                       |
| TIGRfam %   | 31.94%                                  | 35.26%                          | 31.41%               | 33.35%                      |
| Signal peptide | 1,733                                    | 1,408                           | 1,214                | 1,655                       |
| Signal peptide %| 33.69%                                  | 32.77%                          | 22.32%               | 33.43%                      |
| TransMb     | 1,227                                   | 1,018                           | 1,326                | 1,167                       |
| TransMb Perc | 23.85%                                  | 23.69%                          | 24.38%               | 23.58%                      |
| Pfam Clusters | 2,183                                    | 2,091                           | 2,163                | 2,186                       |
| COG Clusters | 2,030                                    | 1,943                           | 2,087                | 2,041                       |
| TIGRfam Clusters | 1,310                                    | 1,246                           | 1,300                | 1,323                       |
| GC Perc     | 0.46                                    | 0.45                            | 0.44                 | 0.45                        |
| Bases       | 5,680,628                               | 4698244                         | 5,701,826            | 5,353,266                   |

**Antibiotic resistance**

There are six separate genes encoding for putative β-lactamases within the genome, but only two have homology at the protein levels with any known *Vibrio* β-lactamases. There is also a multiantibiotic resistance protein MarC, associated with an operon containing a variety of multidrug resistance proteins. This operon is controlled by a MerR type transcriptional regulator, which is often associated with antibiotic resistance [30], and may account for the kanamycin resistance observed in this strain by the authors.

**Acknowledgements**

*We wish to thank* i-G Peninsula (Prospect Place, the Hoe, Plymouth, Devon, UK) for providing funding for this project, and NBAF Edinburgh for performing the sequencing.

**References**

1. Thompson FL, Iida T, Swings J. “Biodiversity of vibrios,” *Microbiology and molecular biology reviews*. [Table of contents]. *Microbiol Mol Biol Rev* 2004; **68**:403-431. PubMed doi:10.1128/MMBR.68.3.403-431.2004

2. Colwell RR, Huq A. Environmental Reservoir of *Vibrio cholerae* The Causative Agent of Cholera. *Ann N Y Acad Sci* 1994; **740**:44-54. PubMed doi:10.1111/j.1749-6632.1994.tb19852.x

3. Tubiash HS, Chanley PE, Leifson E. *Bacillary necrosis*, a disease of larval and juvenile bivalve mollusks. I. Etiology and epizootiology. *J Bacteriol* 1965; **90**:1036-1044. PubMed doi:10.1128/JB.90.3.1036-1044.1965

4. Hada HS, West PA, Lee JV, Stemmler J, Colwell RR. *Vibrio tubiashii* sp. nov., a Pathogen of Bivalve Mollusks. *Int J Syst Bacteriol* 1984; **34**:1-4. doi:10.1099/00207713-34-1-1

5. Beaubrun JLG, Kothary MH, Curtis SK, Flores NC, Eribo BE, Tall BD. Isolation and characterization of *Vibrio tubiashii* outer membrane proteins and determination of a toxR homolog. *Appl EnvironMicrobiol* 2008; **74**:907-911. PubMed doi:10.1128/AEM.02052-07
6. Kothary MH, Delston RB, Curtis SK, McCardell BA, Tall BD. Purification and characterization of a vulnificolin-like cytolytin produced by Vibrio tubiashii. Appl Environ Microbiol 2001; 67:3707-3711. PubMed doi:10.1128/AEM.67.8.3707-3711.2001

7. Abraham T. Distribution of luminous bacteria in semi-intensive penaeid shrimp hatcheries of Tamil Nadu, India. Aquaculture 2004; 232:81-90. doi:10.1016/S0044-8486(03)00485-X

8. Beaubrun JJG, Kothary MH, Curtis SK, Flores NC, Eribo BE, Tall BD. Isolation and characterization of Vibrio tubiashii outer membrane proteins and determination of a toxR homolog. Appl Environ Microbiol 2008; 74:907-911. PubMed doi:10.1128/AEM.02052-07

9. Ben-Haim Y, Zicherman-Keren M, Rosenberg E. Temperature-Regulated Bleaching and Lysis of the Coral Pocillopora damicornis by the Novel Pathogen Vibrio corallilyticus. Appl Environ Microbiol 2003; 69:4236-4242. PubMed doi:10.1128/AEM.69.7.4236-4242.2003

10. Woese CR, Kandler O, Wheelis ML. Towards a natural system of organisms: proposal for the domains Archaea, Bacteria, and Eucarya. Proc Natl Acad Sci USA 1990; 87:4576-4579. PubMed doi:10.1073/pnas.87.12.4576

11. Garrity GM, Holt JG. The Road Map to the Manual. In: Garrity GM, Boone DR, Castenholz RW (eds), Bergey’s Manual of Systematic Bacteriology, Second Edition, Volume 1, Springer, New York, 2001, p. 119-169.

12. List Editor. Validation of publication of new names and new combinations previously effectively published outside the IJSEM. List no. 106. Int J Syst Evol Microbiol 2005; 55:2235-2238. doi:10.1099/ijs.0.64108-0

13. Garrity GM, Bell JA, Liburn T. Class III. Gammaproteobacteria class. nov. In: Garrity GM, Brenner DJ, Krieg NR, Staley JT (eds), Bergey’s Manual of Systematic Bacteriology, Second Edition, Volume 2, Part B, Springer, New York, 2005, p. 1.

14. Garrity GM, Holt JG. Taxonomic Outline of the Archaea and Bacteria. In: Garrity GM, Boone DR, Castenholz RW (eds), Bergey’s Manual of Systematic Bacteriology, Second Edition, Volume 1, Springer, New York, 2001, p. 155-166.

15. Skerman VBD, McGowan V, Sneath PHA. Approved Lists of Bacterial Names. Int J Syst Bacteriol 1980; 30:225-420. doi:10.1099/00207713-30-1-225

16. Véron M. La position taxonomique des Vibrio et de certaines bactéries comparables. C R Acad Sci Hebd Seances Acad Sci 1965; 261:5243-5246.

17. Pacini F. Osservazione microscopiche e deduzioni patologiche sul cholera asiatico. Gazette Medica de Italiana Toscana Firenze 1854; 6:405-412.

18. Shewan J, Veron M. Genus I. Vibrio Pacini 1854, 411. In: Buchanan RE, Gibbons NE (eds), Bergey’s Manual of Determinative Bacteriology, Eighth Edition, The Williams and Wilkins Co., Baltimore, 1974, p. 340-345.

19. Judicial Commission. Opinion 31. Conservation of Vibrio Pacini 1854 as a Bacterial Generic Name, Conservation of Vibrio cholerae Pacini 1854 as the Nomenclatural Type Species of the Bacterial Genus Vibrio, and Designation of Nontype Strain of Vibrio cholerae Pacini. Int Bull Bacteriol Nomencl Taxon 1965; 15:185-186. doi:10.1099/00207713-15-3-185

20. Hada HS, West PA, Lee JV, Stemmler J, Colwell RR. Vibrio tubiashii sp. nov., a Pathogen of Bi-valve Mollusks. Int J Syst Bacteriol 1984; 34:1-4. doi:10.1099/00207713-34-1-1

21. Ashburner M, Ball CA, Blake JA, Botstein D, Butler H, Cherry JM, Davis AP, Dolinski K, Dwight SS, Eppig JT, et al. Gene Ontology: tool for the unification of biology. Nat Genet 2000; 25:25-29. PubMed doi:10.1038/75556

22. Pillidge CJ, Colwell RR. Nucleotide sequence of the 5S rRNA from Listonella (Vibrio) ordalii ATCC 33509 and Listonella (Vibrio) tubiashii ATCC 19105. Nucleic Acids Res 1988; 16:3111. PubMed doi:10.1093/nar/16.7.3111

23. NBAF website. http://nbaf.nerc.ac.uk

24. Lowe TM, Eddy SR. tRNAscan-SE: a program for improved detection of transfer RNA genes in genomic sequence. Nucleic Acids Res 1997; 25:955-964. PubMed doi:10.1093/nar/25.5.955

25. INFERNAL http://infernal.janelia.org

26. Markowitz VM, Korzeniewski F, Palaniappan K, Szeto E, Werner G, Padki A, Zhao X, Dubchak I, Hugenholtz P, Anderson I, et al. The Integrated Microbial Genomes (IMG) system. Nucleic Acids Res 2006; 34:D344-D348. PubMed doi:10.1093/nar/gkj024

27. The DOE Joint Genome Institute. http://img.jgi.doe.gov

28. Thompson FL, Gevers D, Thompson CC, Dayndt P, Naser S, Hoste B, Munn CB, Swings J.
Phylogeny and molecular identification of vibrios on the basis of multilocus sequence analysis. *Appl Environ Microbiol* 2005; 71:5107-5115. PubMed doi:10.1128/AEM.71.9.5107-5115.2005

29. Lilley BN, Bassler BL. Regulation of quorum sensing in *Vibrio harveyi* by LuxO and sigma-54. *Mol Microbiol* 2000; 36:940-954. PubMed doi:10.1046/j.1365-2958.2000.01913.x

30. Brown NL, Stoyanov JV, Kidd SP, Hobman JL. The MerR family of transcriptional regulators. *FEMS Microbiol Rev* 2003; 27:145-163. PubMed doi:10.1016/S0168-6445(03)00051-2