Complete genome sequence of the marine planctomycete *Pirellula* sp. strain 1


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*Pirellula* sp. strain 1 ("Rhodopirellula baltica") is a marine representative of the globally distributed and environmentally important bacterial order Planctomycetales. Here we report the complete genome sequence of a member of this independent phylum. With 7,145 megabases, *Pirellula* sp. strain 1 has the largest circular bacterial genome sequenced so far. The presence of all genes required for heterolactic acid fermentation, key genes for the interconversion of C1 compounds, and 110 sulfatases were expected for this aerobic heterotrophic isolate. Although *Pirellula* sp. strain 1 has a proteinaceous cell wall, remnants of genes for peptidoglycan synthesis were found. Genes for lipid A biosynthesis and homologues to the flagellar L- and P-ring protein indicate a former Gram-negative type of cell wall. Phylogenetic analysis of all relevant markers clearly affiliates the Planctomycetales to the domain *Bacteria* as a distinct phylum, but a deepest branching is not supported by our analyses.

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**Methods**

**Sequencing Strategy.** Genome sequencing was performed by a combination of a clone-based and a whole-genome shotgun approach. Two plasmid libraries with 1.5- and 3.5-kb inserts and a cosmid library (Epicentre Technologies, Madison, WI) were built from *Pirellula* sp. strain 1 DNA. End sequences of inserts were determined by using Big Dye chemistry (ABI), M13 primers, and ABI 3,700 capillary sequencers (ABI) up to eightfold sequence coverage. All raw sequences were processed by PHRED (17) and controlled for vector or Escherichia coli contamination. Reads were assembled by PHRAP and manually finished by using GAPS (18). The quality of the sequence data was finished to reach a maximum of 1 error within 10,000 bases. Gap closure and finishing of the sequence were done by resequencing clones, primer-walking, and long-range PCR. Locations and sequence of repetitive sequence elements were additionally controlled by PCR.

**Open Reading Frame (ORF) Prediction.** Three different programs were used for ORF prediction, GLIMMER (19), CRITICA (20), and ORPHEUS (21). A nonredundant list of ORFs was generated by parsing the results with a self-written Perl-script. The script applied performs in the following way: For all ORFs that are predicted identically by all three gene finders, only one is kept. If the script recognizes identical stop positions but different starts and the difference is below 10% of the sequence length, only the longer ORF is kept. If the difference is more than 10%, both ORFs are kept.

**Annotation.** The software package PEDANT PRO (22) was used for annotation. All automatically generated results were evaluated manually for final annotation. Obviously overpredicted ORFs, e.g., overlapping ORFs without functional assignment, were marked for deletion and deleted after cross-checking by at least two independent annotators.

**Data Analysis.** For origin and terminus determination a combination of compositional indexes and oligomer distribution skew was used. The following compositional indexes were determined with self-written Perl-scripts: (i) cumulative GC skew [sum of (G − C)/(G + C) over adjacent windows of 10 kb]; (ii) keto excess [sum(GT) − sum(AC)]; (iii) purine excess [sum(AG) − sum(TC)] and the external program OLIGOSKEW6 (www.tigr.org/~salzberg/oligoskew6). Repeats were detected by the software REPETU (23). DNA flexibility such as curvature and bending...
was calculated with the BANANA program, and sequence twist was calculated with the program BTWISTED, both taken from the EMBOSS package (www.hgmp.mrc.ac.uk/Software/EMBOSS). Codon usage (codon adaptation index, CAI) was calculated with the CODONW-program (www.molbiol.ox.ac.uk). Highly expressed and alien genes according to Karlin and Mrazek (24) were identified with self-written Perl-scripts. For the phylogenetic distribution of the best BLAST hits the SEALS package and the taxonomy of the National Center for Biotechnology Information was used. Tat signals were found by extracting all proteins containing twin arginines plus two additional amino acids of the conserved Tat pattern (SRRXFLK).

For whole-genome visualization the software tool GENEWIZ (25) was used. Total gene numbers were calculated by searches against all publicly available genomes with Pfam profiles (http://pfam.wustl.edu) by using GENDB 1.1 (46). For phylogenetic reconstructions the preliminary sequence of Gemmata obscuriglobus UQM 2246 was obtained from The Institute for Genomic Research (www.tigr.org). The program package ARB was used for phylogenetic analysis (www.arb-home.de).

Supporting Information. All supporting information (Appendices I–8) is available on the PNAS web site, www.pnas.org. The complete annotation data and all supporting information are available on the home page of the REGX Project, www.regx.de. For fast searching a BLAST server is available for public use.

Results and Discussion

Genome Organization. With a size of 7,145,576 bases, Pirellula sp. strain 1 has the largest prokaryotic circular genome sequenced so far. Origin and terminus could be clearly identified by the change in cumulative GC and AT skews (Fig. 1). A single, unlinked rRNA operon was identified near the origin. Unlinked rrr operons have also been described for other planctomycetes (13) but the 460 kb region did not give any indications for lateral gene transfer. On the contrary, a regular codon adaptation index and the localization of housekeeping genes (e.g., several tRNA synthetases, ribosomal proteins, and flagella proteins) in this region indicate that most probably an internal chromosomal inversion has occurred. This conclusion is supported by five and four flanking transposases. Two of them are identical and have reverse orientation. In total, 16% (13) of all transposase genes are located within this region, supporting a hot spot for large genomic rearrangements.

Table 1. General features of the Pirellula sp. strain 1 genome

<table>
<thead>
<tr>
<th>Component of chromosome</th>
<th>Property</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total size, bases</td>
<td>7,145,576</td>
</tr>
<tr>
<td>G + C content, %</td>
<td>55.4</td>
</tr>
<tr>
<td>Coding sequences</td>
<td>7,325</td>
</tr>
<tr>
<td>Coding density, %</td>
<td>95</td>
</tr>
<tr>
<td>Average gene length, bases</td>
<td>939</td>
</tr>
<tr>
<td>Genes with similarities in databases*</td>
<td>3,380 (46%)</td>
</tr>
<tr>
<td>Genes with functional assignments</td>
<td>2,582 (35%)</td>
</tr>
<tr>
<td>tRNAs</td>
<td>17 (165) and (235–55)</td>
</tr>
<tr>
<td>rRNAs</td>
<td>70</td>
</tr>
<tr>
<td>Other stable RNAs</td>
<td>1 (ribozyme)</td>
</tr>
</tbody>
</table>

*Threshold for BLASTP E value ≤ 1 × 10–5, includes hits to hypothetical proteins.

Irregularity. A large inversion at position 87,500 to 431,000 of 343.5 kb is indicated by cumulative GC-skew and other structural parameters (Fig. 1 and Appendix 1, which is published as supporting information on the PNAS web site). Analysis of this anomalous region did not give any indications for lateral gene transfer. On the contrary, a regular codon adaptation index and the localization of housekeeping genes (e.g., several tRNA synthetases, ribosomal proteins, and flagella proteins) in this region indicate that most probably an internal chromosomal inversion has occurred. This conclusion is supported by five and four flanking transposases. Two of them are identical and have reverse orientation. In total, 16% (13) of all transposase genes are located within this region, supporting a hot spot for large genomic rearrangements.

Gene and Functional Prediction. An initial nonredundant list of 13,331 potential ORFs was generated. By manual annotation, this ORF set could be reduced to 7,325 ORFs, which equals a gene coverage of 95%. A BLASTX of all intergenic regions confirmed that a comprehensive ORF set was achieved. More than half (56%, 4,148) of the predicted ORFs had reliable functional predictions, which is about 20% less than the numbers found in general (26). This low percentage reflects the distinct phylogenetic position and the lack of molecular studies performed on Planctomycetes so far. An overview of the localization of the functional genes according to our functional classification (27) is given in Fig. 2. The complete annotation results are available at www.regx.de.

Genome Size. We found that 1,301 genes (i.e., 17.6% of all genes), with an average length of 464 aa, have more than one copy within the genome. In total, multicopy genes make up for about 25.4% of the genome. In total, multicopy genes make up for about 25.4% of the genome. Therefore extensive gene duplication is not the reason for the large genome size of Pirellula sp. strain 1. A large genome with an expanded genetic capability might be a prerequisite for environmental adaptability, as already discussed for the genome of Pseudomonas aeruginosa (29).

Potential Environmental Adaptations. Metabolism. The annotation process identified the standard pathways for heterotrophic bacteria such as glycolysis, the citrate cycle, and oxidative phosphorylation. Pirellula sp. strain 1 lacks the glyoxylate bypass and the Entner–Doudoroff pathway but exhibits the pentose-phosphate cycle. Fon-
thermore, it seems to be capable of synthesizing all amino acids (Appendix 2, which is published as supporting information on the PNAS web site).

Recent growth studies did not provide evidence that *Pirellula* sp. strain 1 can grow under nitrate-reducing or fermentative conditions. Interestingly, however, all genes required for heterolactic acid fermentation are present (Fig. 2 and Appendix 3, which is published as supporting information on the PNAS web site). Expression of the genes is likely, because the key enzyme lactate dehydrogenase has been predicted to be highly expressed on the basis of codon usage. Furthermore, both *Pirellula marina* and *Planctomyces limnophilus* have been described to be capable of carbohydrate fermentation (30). This capability could explain why planctomycetes were found in anoxic marine and freshwater sediments and anoxic terrestrial habitats (1–3).

**Motility.** The life cycle of *Pirellula* sp. strain 1 consists of an aggregate-forming sessile form and a motile swarmer cell. In the genome all genes for a functional flagellum could be determined, whereas except for cheY, essential genes for chemotaxis such as cheA, cheB, cheR, cheW, and cheZ could not be identified.

**Transporters.** As a free-living organism, *Pirellula* sp. strain 1 was expected to have a wide range of transporters (29). A comparative study with Pfam profiles for ABC-transporters against all 70 publicly available prokaryotic genomes revealed that the 55 ABC-transporters found in *Pirellula* sp. strain 1 is close to the calculated mean of 49 transporters. In comparison with other free-living bacteria this is only about one-third of the 148 ABC-transporters found with the same method in *Streptomyces coelicolor* A3 (2), but similar to the 45 transporters of *Caenorhabditis elegans* (Appendix 4, which is published as supporting information on the PNAS web site). Annotation revealed that ABC-transporters for ribose, oligopeptides, phosphate, manganese, nitrate, and sodium are present, but only one phosphotransferase system (PTS) specific for fructose could be identified. Exceptional is a set of ORFs for nitrate transport and nitrate/nitrite reduction that were predicted to be highly expressed (PHX) on the basis of codon usage (24). This set of ORFs could be essential in nitrogen-limited marine systems (31).

**Stress response.** The genome harbors homologues to superoxide dismutase and all four known types of catalases (Fig. 2). Methionine-sulfoxide reductases are present to repair oxidized methionine. By synthesis of a cytochrome *d* oxidase as an alternative to the regular cytochrome *aa*3 *Pirellula* sp. strain 1 should be able to cope with low oxygen concentrations. Many mechanisms are present to reduce the damaging effect of UV radiation. Besides the genes for SOS response (*recA, lexA, uvrA, uvrB, and uvrC*), *Pirellula* sp. strain 1 has a photolyase gene organized in an operon-like manner with genes encoding phytoene dehydrogenase and phytoene synthase. Probably, UV stress triggers the biosynthesis of a UV-protection carotenoid, which might be responsible for the pinkish color of *Pirellula* sp. strain 1. Regarding temperature stress, *Pirellula* sp. strain 1 has many homologues to heat and cold shock DNA-binding proteins. Detoxification seems to take place by means of unspecific export systems like cation efflux systems of the AcrB/AcrD/AcrF family or unspecific multidrug export systems for hydrophobic compounds. A cytochrome
P450 mono-oxygenase and an epoxide hydrolase are present for the detoxification of xenobiotics. Specific detoxification involves mercury reductase, arsenate reductase, and the ArsA arsenite-exporting ATPase. The harmful effect of d-tyrosine binding to tyrosyl-tRNA is minimized by D-tyrosyl-tRNA Tyr deacylase. In addition, Pirellula sp. strain 1 has a gene encoding a bacterial hemoglobin, which is believed to detoxify NO by oxidation. Finally, the genome has some homologues to carbon-starvation proteins, including DNA-protection proteins.

**Antibiotics.** Several ORFs potentially coding for polyketide antibiotics and nonribosomal polypeptide antibiotics or a mixture of both have been determined in the genome of **Pirellula** sp. strain 1. In general the ORFs are unusually long, coding for proteins from 916 up to 3,665 aa.

**Sulfatases.** The **Pirellula** sp. strain 1 genome harbors 110 genes encoding proteins with significant similarity to prokaryotic (82 genes; 75%) and eukaryotic (28 genes; 25%) sulfatases. For instance, similarity was found to alkylsulfatase of **Pseudomonas aeruginosa**, to arylsulfatases of **Pseudomonas sp.**, to mucin-desulfatating sulfatase of **Prevotella sp.**, and to archaeal arylsulfatase, as well as to mammalian iduronate-2-sulfatase and arylsulfatases A and B. In comparison, the analysis of 70 published prokaryotic genomes with a specific Pfam profile revealed a maximum of only 6 sulfatases found in the **Pseudomonas aeruginosa** PAO1 genome. In **Pirellula** sp. strain 1, the sulfatase genes are distributed across the genome in 22 clusters containing two to five genes (Fig. 2).

In **Pirellula** sp. strain 1, all detected sulfatase gene products, except for the three alkylsulfatases, are of the cysteine type; 85 (79%) of them show the canonical CXXXR motif and are hence considered as potentially functional (32). In contrast to the known bacterial cysteine-type sulfatases, which are cytosolic enzymes (e.g., arylsulfatase AtsA of **Pseudomonas aeruginosa**), for 26 (31%) of the 85 potentially functional sulfatases in **Pirellula** sp. strain 1 a signal peptide is predicted with high probability, suggesting an extracellular localization of the proteins.

The fact that the sulfatase genes in **Pirellula** sp. strain 1 outnumber those present in all other known prokaryotic genomes by two orders of magnitude raises the question about their physiological role. Bacterial sulfatases seem to be primarily used in sulfur scavenging, and their expression is known to be tightly regulated and dependent on the sensing of sulfur deprivation (32). As marine systems are characterized by high inorganic sulfate concentrations, sulfur limitations should not occur. Therefore, **Pirellula** sp. strain 1 might use its sulfatases to access more effectively the carbon skeleton of sulfated compounds as an energy source rather than to meet its sulfate requirements. Cleavage of sulfate esters in sulfated high molecular weight glycoproteins (mucins) to increase the efficiency of polymer degradation by other enzymes has been described for **Prevotella** sp. RS2. Seven of the 110 sulfatase genes in **Pirellula** sp. strain 1 encode proteins with high similarity to mucin-desulfatating sulfatase of **Prevotella** sp. RS2, an enzyme that seems to be specific for the cleavage of sulfate from N-acetylglucosamine 6-sulfate in mucin side chains (33). Remarkably, one of the seven genes in **Pirellula** sp. strain 1 is located next to a gene encoding a protein with some similarity to N-acetylglucosamine-6-phosphate deacetylase, a protein involved in metabolism of N-acetylglucosamine. This compound is known to support growth of **Pirellula** sp. strain 1 (M.B., unpublished results) resembles very closely a pathway of formaldehyde oxidation/detoxification in methylotrophic proteobacteria. These tetrahydromethanopterin (H₄MPT)-dependent enzymes were previously thought to be unique for anaerobic methanogenic and sulfate-reducing **Archaea**. Recently, however, they have been shown to play an essential physiological role in methylotrophic proteobacteria (34). **Pirellula** sp. strain 1, to our knowledge, the first bacterial organism outside the proteobacterial division found to contain genes encoding H₄MPT-dependent enzymes. In context with the fact that planctomycetes constitute an independent phylum, our finding revives the discussion on the evolutionary processes leading to the distribution of these archaeal genes.

**Cell Biology. Cell wall.** Planctomycetes are the only group of free-living members of the domain **Bacteria** known so far that have no peptidoglycan in their cell walls. Instead, they are stabilized by a protein sacculus with disulfide bonds (35). A systematic investigation for genes involved in peptidoglycan biosynthesis revealed that murB, murE, murG, dddA, and upk (bacA) are present. Furthermore, **Pirellula** sp. strain 1 possesses the gene glmS, which is involved in the formation of N-acetyl-α-glucosamine, a precursor for peptidoglycan biosynthesis. Other key enzymes, such as MurA, MurC, MurD, MurF, and DdaA for the final cross-linking of peptidoglycan, are notably absent from the **Pirellula** sp. strain 1 genome. The preservation at least some of the genes of the peptidoglycan synthesis pathway suggests that **Pirellula** sp. strain 1 is not a descendant from a bacterium evolving before the invention of peptidoglycan, as proposed earlier (36). It rather seems that after the development of a proteinaceous cell envelope in planctomycetes, genes for peptidoglycan biosynthesis were successively lost.

**Membrane.** It is noteworthy that the **Pirellula** sp. strain 1 genome harbors all genes required for biosynthesis of lipid A, the major constituent of the lipopolysaccharide (LPS) layer in Gram-negative bacteria. The presence of these genes is in line with earlier reports of presence of lipid A with unusual portions of long-chain 3-OH fatty acids in members of the **Pirellula/Planctomycetes** group (12). Nevertheless, the key enzymes necessary for the biosynthesis of an O-specific side chain (O-antigen ligase; O-antigen polymerase) are absent from the **Pirellula** sp. strain 1 genome. The presence of lipid A and homologues to the flagellar L- and P-ring protein suggests that the cell envelope of planctomycetes was converted from a Gram-negative type of cell wall. Furthermore, **Pirellula** sp. strain 1 lacks the signature sequences in the ribosomal protein S12 and SecF typical for low and high G+C Gram-positive bacteria, respectively (37).

**Compartmentalization.** One of the most striking properties of the **Planctomycetes** is their complex internal structures (1). Ribosomes are located only within the riboplasm, therefore proteins targeted to the paryphoplasm have to overcome the intracytoplasmic membrane. This requires effective protein targeting. A com-
Regulation is based to a greater extent on altering the promoter tors. Currently, with 65 sigma factors, only factors, including 16 ECF (extracytoplasmic function) sigma factors, are important. Peptides showed that a comparison for Tat (twin arginine translocation) signal site). A comparison for Tat (twin arginine translocation) signal peptides showed that *Pirellula* sp. strain 1 has the highest number of all genomes investigated (135; 18.9 per megabase). Effective protein targeting might be the basis for the polar organization of *Pirellula* sp. strain 1 and for distinct features (e.g., stalks, holdfast substance, crateriform structures) present only in certain regions of the cells.

**Cell division.** Cell division involves a plethora of genes. The most important are *fsz*, *fsA*, *fsi*, *fsL*, *fsQ*, *fsN*, *zipA*, and *fsW* (38). Surprisingly, with the exception of *fsK*, all genes are absent from the genome of *Pirellula* sp. strain 1. A lack of the key enzyme FtsZ, the major constituent of the septal replication ring, has so far been reported only for chlamydiae, the *Crenarchaeota*, and *Ureaplasma urealyticum* (38). Not much is known about replication in planctomycetes, especially how the cell compartments are distributed to the daughter cells. Altogether, cell division in *Pirellula* sp. strain 1 must follow a different pathway than reported for the model organisms *E. coli* and *B. subtilis*.

**Life cycle.** *Pirellula* sp. strain 1 exhibits a life cycle similar to *C. crescentus* (39). Surprisingly, no homologue to the master response regulator protein CtrA has been found within the genome of *Pirellula* sp. strain 1. However, the origin in *Pirellula* sp. strain 1 contains some patterns similar to the CtrA binding site pattern TTAAN-TTAA upstream of *dnaN* (e.g., TTAAN-AAAAC), which might indicate a similar control mechanism.

**Regulation.** Analysis of the *Pirellula* sp. strain 1 genome with 116 relevant Pfam 7.2 family models shows only 135 genes with motifs for predicted transcriptional regulators. No evidence for eukaryotic-like transcriptional regulators could be found. There are 68 response regulators, which allow microorganisms to respond to changes in their environment, but common bacterial regulators such as LysR are absent or underrepresented. A comparative analysis in all currently available bacterial genomes was performed. The results confirm earlier findings that the proportion of the genome encoding transcriptional regulators increases with genome size (29, 40) (Appendix 6, which is published as supporting information on the PNAS web site). Nevertheless with a genome size of more than 7 megabases and only 2% predicted regulatory genes, *Pirellula* sp. strain 1 clearly contradicts this trend. It remains to be determined whether these results reflect a lack of knowledge in the diversity of regulatory proteins or even unknown gene regulation mechanisms. For example, a unique family of predicted DNA-binding proteins has been reported in the genome of *S. coelicolor* A3(2) (40), which might constitute a family of *Streptomyces*-specific transcriptional regulators. Regulation of metabolic capacities of *Pirellula* sp. strain 1 were recently addressed by a proteomic approach, which revealed differential protein patterns in response to carbohydrate substrates used for growth (41).

**Sigma factors.** *Pirellula* sp. strain 1 encodes for a total of 51 sigma factors, including 16 ECF (extracytoplasmic function) sigma factors. Currently, with 65 sigma factors, only *S. coelicolor* has a higher number. Nevertheless, it seems that in *Pirellula* sp. strain 1 gene regulation is based to a greater extent on altering the promoter specificity of the RNA polymerase. Further support for this hypothesis comes from the observation that genes for essential pathways such as purine or biotin and amino acid biosynthesis are not organized in operons, as is known for most of the prokaryotes. This scattering, together with the split rRNA operon, requires a different way of regulation.

**Evolution.** Phylogeny. The currently accepted bacterial systematics based on 16S rRNA assigns the Planctomycetes with the genera *Pirellula*, *Planctomyces*, *Isosphaera*, and *Gemmatia* as an independent monophyletic phylum (9). For evaluation, phylogenetic trees for commonly used alternative markers such as the 23S rRNA, elongation factors Tu and G, ATP-synthase subunits, *RecA*, heat shock proteins HSP60 and HSP70, RNA polymerase, and DNA gyrase subunits as well as the aminoacyl-tRNA synthetases of *Pirellula* sp. strain 1 were constructed. A common *Pirellula–Gemmatia* (G. *obscuriglobus* UQM.2246) cluster separated from all other phyla or major subgroups is seen in trees derived from 23S rRNA, elongation factors, ATP-synthase subunits α and β, DNA gyrase subunits A and B, the heat shock proteins HSP60 and HSP70, the *recA* protein, and most of the tRNA synthetases. Multiple copies are present in the case of ATP-synthase α and β as well as the DNA gyrase A and B subunits. The sequence divergence of the duplicates corresponds to the phylum level. Thus the individual monophyletic *Pirellula–Gemmatia* pairs are separated from other phyla in the respective trees. However, the multiple copies of heat shock proteins HSP60 and HSP70 cluster in common groups with the *Pirellula* and *Gemmatia* proteins phylogenetically intermixed. Keeping in mind the possible pitfalls and the differences in resolving power when functional genes are used for tree reconstruction (42), the status as an independent phylum affiliated to the domain *Bacteria* is clearly supported in the majority of analyses. A deepest branching of planctomycetes within the bacterial subtree as reported recently (10) is not convincingly supported by any of the markers, as evaluated by applying different tree-building methods, parameters, and significance tests.

**ATP-synthases.** *Pirellula* sp. strain 1 is the only bacterium described so far that contains two F,F′,F′′ ATP-synthases (Fig. 2). By gene organization (α, c, b, δ, α, γ, β, e) and similarity one ATP-synthase resembles the “standard” ATP-synthase that is common in all *Bacteria* (43). Nevertheless, the J gene found in all *Bacteria* except for *Thermotoga maritima* is missing. The second set of F,F′,F′′-ATP-synthase genes has a different operon structure with β and e genes followed by the J, X, α, c, and b genes and the α, γ genes (Appendix 7, which is published as supporting information on the PNAS web site). The strong conservation and the high similarity of this ATP-synthase operon to *Methanosarcina barkeri* indicates a lateral gene transfer event. It remains unclear whether the genes were transferred from the archael to the bacterial domain or vice versa.

**Phylogenetic Distribution of Best BLAST Hits.** All 7,325 potential proteins (ORFs) in the *Pirellula* sp. strain 1 genome were searched against the National Center for Biotechnology Information non-redundant database. By setting the cut-off for the BLASTP expectation value ≤1 × 10−3, significant hits could be obtained for 3,380 genes. Of these genes, 83% were assigned to the domain *Bacteria*, 9% and 8% to the domains *Archaea* and *Eukarya*, respectively (Fig. 2). The large number of hits to eukaryotes is exceptional. *T. maritima*, for example, shows 24% hits to *Archaea* but only 2% to *Eukarya*, and in *E. coli* only 0.4% of the best hits assign to *Archaea* or *Eukarya*. Nevertheless, among the 270 genes found in *Pirellula* sp. strain 1 no trend for a certain organism of origin or a distinct functional category could be detected (Appendix 8, which is published as supporting information on the PNAS web site). Furthermore, the two genes for the integrin α-V and inter-α-trypsin inhibitor found in *Pirellula marina* and *G. obscuriglobus*, considered to be typical for *Eukarya* (44), could not be detected in the *Pirellula* sp. strain 1 genome.
**Relationship to Chlamydia.** The phylogenetic analysis of a comprehensive set of markers with different tree-building methods and confidence tests revealed that only the trees for DNA gyrase, RNA polymerase C, and lysyl- and valyl-tRNA synthetase supported a moderate relationship to *Chlamydia*. Using only distance matrix methods, we also found a remote relationship to *Chlamydia* in some ribosomal proteins, DNA A, Hsp60, Rho, the protein component of RNase P, and CTP synthase. According to the indel method of Gupta and Griffiths (37), *Pirellula* sp. strain 1 branches off between spirochetes and *Chlamydia*.

As already mentioned, *Pirellula* sp. strain 1 and *Chlamydia* share some noteworthy features: both lost their peptidoglycan cell walls in favor of a proteinaceous cell envelope. Furthermore, *Chlamydia* and *Pirellula* sp. strain 1 have two dnaA copies and lack ftsZ, indicating an unknown mode of cell division. In addition, the ribosomal protein L30 is missing from the sp copron in *Chlamydia* as well as in *Pirellula* sp. strain 1. Although the complete genome sequence does not confirm a close relationship of *Chlamydia* in some cases but also gives hints for the specific adaptations to the different habitats and the origin of the unique combination of morphological and ultrastructural properties.

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**Conclusions.** The genome sequence of *Pirellula* sp. strain 1 has revealed insights into adaptations of free-living marine bacteria. Environmental versatility demands an enhanced genetic complexity harbored by larger genomes (29, 40). In case of *Pirellula* sp. strain 1 no indication for a recent expansion of the genome could be detected. In contrast, except for some irregularities (Fig. 1), the plot of the cumulative GC-skew of *Pirellula* sp. strain 1 is very “smooth” with clear maxima and minima (Appendix I). From the genome features it is now possible to propose certain lifestyle of *Pirellula* sp. strain 1. In the water column *Pirellula* sp. strain 1 gains energy from the aerobic oxidation of mono- or disaccharides derived from the cleavage of sulfated polymers produced by algae. Protection systems for UV and the expression of carotenoids protect *Pirellula* sp. strain 1 from irradiation at the water surface. Nitratre transporters support growth even under limited-nitrogen conditions common in continental shelf areas. The holdfast substance enables *Pirellula* sp. strain 1 to attach to nutrient-rich marine snow particles slowly sedimenting to the sea floor. When the bacterium reaches the sediment the expression of cytochrome *d* oxidase allows survival under low oxygen conditions. Anoxic conditions force *Pirellula* sp. strain 1 to switch to heterolactic acid fermentation or pathways involving formaldehyde conversion if not to support growth, then at least to allow basic maintenance metabolism. With the expression of genes for carbon starvation *Pirellula* sp. strain 1 can even outlast periods of nutrient depletion. The formation of swarmer cells helps *Pirellula* sp. strain 1 to reach for new resources. The high number of sigma factors allows the tight control of gene expression under changing environmental conditions.

Exceptionally interesting will be the genome comparison of *Pirellula* sp. strain 1 with the freshwater isolate *G. obscuriglobus* UOM.2246 (45), currently being sequenced by The Institute for Genomic Research, and *Gemmata* sp. Wil-1, being sequenced by Integrated Genomics. It will not only reveal common traits of the *Planctomycetes* but also give hints for the specific adaptations to the different habitats and the origin of the unique combination of morphological and ultrastructural properties.