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Microbial methane oxidation is one of the key drivers of oxygen consumption in marine sediments and the overlying water column (1). Methanotrophic bacteria are the primary producers of many cold and hot seep ecosystems (2, 3). Here, we report three genome sequences of gammaproteobacterial methanotrophs isolated from three marine ecosystems. Methylobacter marinus A45 (a methanol-adapted strain, formerly Methylomonas methanica A4, ACM 4717) was isolated from sewage outfall sediment near Los Angeles, CA (4). Methylobacter sp. strain BBA5.1 was isolated from the surface layer of estuary sediment collected at low tide near Newport, Bay Estuary (CA) (5). Methylomonas vadi IT-4 (= JCM 13665T = DSM 18976T) was isolated from a

<table>
<thead>
<tr>
<th>Species</th>
<th>Sequencing platform(s)</th>
<th>Genome assembly and annotation</th>
<th>Genome coverage (×)</th>
<th>Genome size (Mb)</th>
<th>No. of scaffolds (no. of contigs)</th>
<th>Core (accessory) metabolic pathways*</th>
<th>NCBI accession no.</th>
</tr>
</thead>
<tbody>
<tr>
<td>M. marinus A45</td>
<td>Illumina</td>
<td>Velvet 1.1.05, AllPaths, Phrap 4.24, Prodigal 2.5</td>
<td>1,237</td>
<td>4.99</td>
<td>9 (49)</td>
<td>pMMO, pXmo, Mxa, XoxF1, XoxF2, H,F,H,MPT, FDH, RuMP, EMP, EDD, dPPP, PPP, pSC, TCA</td>
<td>ARVS000000000</td>
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<tr>
<td>Methylobacter sp. BBA5.1</td>
<td>Illumina, PacBio RS</td>
<td>AllPaths, Prodigal 2.5</td>
<td>290</td>
<td>5.07</td>
<td>87 (91)</td>
<td>pMMO, pXmo, Mxa, XoxF1, XoxF2, H,F,H,MPT, FDH, RuMP, EMP, EDD, dPPP, PPP, pSC, TCA</td>
<td>JQKS000000000</td>
</tr>
<tr>
<td>M. vadi IT-4</td>
<td>Illumina, PacBio RS</td>
<td>Prodigal 2.5</td>
<td>272</td>
<td>4.33</td>
<td>1 (1)</td>
<td>pMMO, Mxa, XoxF, H,F,H,MPT, FDH, RuMP, EMP, EDD, dPPP, PPP, pSC, TCA</td>
<td>JPON000000000</td>
</tr>
</tbody>
</table>

* dPPP, dissipatory pentose-phosphate pathway; EDD, Entner-Doudoroff pathway; EMP, Embden-Meyerhof-Parnas pathway; FDH, formate dehydrogenases; H,F, folate-linked C1 transfer; H,MPT, methanopterin-linked C1 transfer; Mxa, PQQ-linked methanol dehydrogenases; pMMO, membrane-bound methane monooxygenase; pSC, partial serine cycle; pXmo, methane/ammonia monooxygenase-related proteins of unknown function; PPP, pentose-phosphate pathway; RuMP, assimilatory ribulose monophosphate pathway; Xox, PQQ-linked methanol and formaldehyde dehydrogenases (i.e., no evidence for the glyoxylate regeneration pathway was found); TCA, tricarboxylic acid cycle.

This is contribution 12 from OMeGA.
microbial mat of a shallow submarine hydrothermal system near Taketomi Island, Okinawa, Japan (6).

DNA samples from the three strains were prepared using the standard phenol-chloroform method (7). DNA sequence data were obtained at the Joint Genome Institute using a combination of PacBio (8) and Illumina (9) technologies, and draft genome sequences were assembled. The computational tools used for genome sequencing and assembly are listed in Table 1.

All three sequenced marine methanotrophs are obligate methane and methanol utilizers. All three genomes harbor genes typical for type I methanotrophs, including genes encoding particulate methane monooxygenase (pmoCA), the PQQ-dependent methanol dehydrogenases (mxaF and multiple copies of oxxF), genes for tetrahydromethanopterin (H₂MPT)- and tetrahydrofolate (H₄F)-dependent C₄ transfer pathways, genes of the ribulose monophosphate pathway, including its phosphoketolase variant (10), and genes encoding a complete tricarboxylic acid (TCA) cycle and a partial serine cycle (10) (Table 1). The gpmABC gene clusters (11) linked to a distant homologue of the nitrate-nitrite transporter (narK) were found in the Methylobacter sp. strain BB5.1 and M. marinus A45 genomes. A phosphoenolpyruvate carboxylase gene (pcg) was found encoded in M. vadi IT-4 only. Genes encoding soluble methane monooxygenase, known glyoxylate regeneration pathways, and RuBisCO were not detected. Genes involved in ammonium and nitrate assimilation are present in all three genomes. The genomes of strains A45 and BBA5.1 contain all genes necessary to provide for urea hydrolysis and nitrogen fixation. M. vadi IT-4 has the potential for dissimilatory nitrite reduction to nitric oxide, as suggested by the presence of nir genes. The NADH:ubiquinone reductase (H⁺)-translocating genes (nuoABCDEFGHJKLMN) were identified in M. marinus A45 only. All strains possess genes encoding Na⁺-transporting NADH:ubiquinone oxidoreductase (napABCD), ubiquinol-cytochrome bc complex, cytochrome b, cytochrome c oxidase, cytochrome P450 and P460, and cytochrome d ubiquinol oxidase. Cytochrome bo, quinol oxidase was found in M. vadi IT-4 only. Both Methylobacter species possess genes encoding the Na⁺-translocating ferredoxin:NAD⁺ oxidoreductase complex (rfABCDGE). All genomes contain genes encoding pyruvate-ferredoxin, flavodoxin oxidoreductases, and all three strains possess ectoine biosynthesis genes.

The genome of M. marinus A45 includes a chromosomally integrated complete copy of a bacteriophage genome (predicted size, 65 kb) integrated in the chromosome, indicating the possibility of lysogenic infection in methanotrophic bacteria. These genomes provide a valuable resource to obtain new insights into environmental controls of fitness and diversity in methanotrophs, mechanisms of genetic exchange within methanotrophic communities, and the potential for the development of new genetic tools for methanotrophs.

Nucleotide sequence accession numbers. The genome sequences have been deposited in GenBank under the accession numbers listed in Table 1.

ACKNOWLEDGMENTS

We thank all members of the Organization for Methanotroph Genome Analysis for collaboration (OmEGa) and Genoscope (France) for access to its MicroScope platform for comparative genome analysis (http://www.genoscope.cns.fr/age/microscope/home/).

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REFERENCES