



Draft Genome Sequences of *Methanobrevibacter curvatus* DSM11111, *Methanobrevibacter cuticularis* DSM11139, *Methanobrevibacter filiformis* DSM11501, and *Methanobrevibacter oralis* DSM7256

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Here, the draft genome sequences of four different *Methanobrevibacter* species are presented. Three of the *Methanobrevibacter* species (*M. curvatus*, *M. cuticularis*, and *M. filiformis*) have been isolated from the termite hindgut, while *M. oralis* was isolated from human subgingival plaque.

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Methanobacteriales, in particular members of the genera *Methanobrevibacter*, have been shown to be some of the most abundant methanogenic archaea in various intestinal environments, including the human gut (1), the termite hindgut (2), and the ovine and bovine rumen (3). Recent studies have indicated that *Methanobrevibacter* species may not only be contributing to greenhouse gas emissions from farm animals (4, 5), but may also have effects on human physiology and health (6, 7). It is therefore of great interest to gain a better understanding of how different *Methanobrevibacter* species have adapted to specific host environments at the molecular level. Genome sequences of *Methanobrevibacter* species have been obtained so far for strains from the human intestinal tract and the rumen (8–12), but not from any insect guts or the human oral cavity. *Methanobrevibacter oralis* DSM7256, isolated from the human subgingival plaque (13), is also the first sequenced representative of all human oral methanogens.

Genomic DNA was ordered by the DSMZ (Braunschweig) or was isolated using the MasterPure complete DNA purification kit (Epicentre, Madison, WI, USA). The extracted DNA was used to generate Illumina-shotgun libraries according to the manufacturer's protocol (Illumina, San Diego, CA, USA). Sequencing was conducted using a MiSeq and MiSeq reagent kit v3 (2 × 300 bp paired end) as recommended by the manufacturer (Illumina). Sequencing resulted in 1,934,710 (*M. filiformis*), 1,983,778 (*M. ora-*

lis), 3,298,762 (*M. curvatus*), and 3,533,158 paired end reads (*M. cuticularis*), respectively. Trimmomatic 0.32 (14) was used to filter low-quality reads and for clipping of adapter contaminations. The assembly was performed with the SPAdes genome assembler software 3.6.2 (15). Coverages were determined using QualiMap version 2.1 (15, 16) and automatic annotation was performed using the software tool PROKKA (17). General genome features are listed in Table 1.

Sequencing the genomes of the four different *Methanobrevibacter* genomes provides reference sequences for comparative analyses with other *Methanobrevibacter* genomes and may reveal adaptive traits of *Methanobrevibacter* species to different environments. Some characteristic features and differences between *Methanobrevibacter* species are already apparent from formal description of the type strains, e.g., presence of catalase activity in the three *Methanobrevibacter* species from the termite hindgut (18, 19). The genome sequences allow the identification of the potential molecular basis of this enzyme activity: A monofunctional heme-dependent catalase similar to the enzyme purified from *M. arboriphilus* (20). The gene encoding this enzyme is present in each of the genomes of the three *Methanobrevibacter* species isolated from the termite hindgut, but appears to be absent from the genome of *M. oralis*.

Nucleotide sequence accession numbers. These whole-genome shotgun projects have been deposited at DDBJ/EMBL/

TABLE 1 Genome features and GenBank accession numbers of sequenced strains

Strain	Genome size (bp)	G+C content (%)	No. of scaffolds (>500 bp)	No. of CDSs ^a	No. of rRNAs	No. of tRNAs	Accession no.
<i>M. filiformis</i> DSM 11501	2,606,143	26.99	295	1,933	3	29	LWMT00000000
<i>M. oralis</i> DSM 7256	2,140,433	27.71	136	1,994	5	24	LWMU00000000
<i>M. curvatus</i> DSM 11111	2,414,608	25.72	232	1,969	4	31	LWMV00000000
<i>M. cuticularis</i> DSM 11139	2,608,702	26.79	169	2,061	3	30	LWMW00000000

^a CDSs, coding sequences.

GenBank under the accession numbers listed in Table 1. The versions described here are the first versions.

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