

Actinotalea subterranea sp. nov.

Submitted by Grouzdev, Denis

Species *Actinotalea subterranea*

Etymology

[sub.ter.ra.ne'a] **L. fem. adj.** *subterranea*, subterranean, referring to the site of isolation

Nomenclatural type

[NCBI Assembly: GCA_008364845.1](#) ^{Ts}

Reference Strain

[Strain sc|0040321](#): HO-Ch2 = [VKM Ac-2850](#) = [KCTC 49656](#)

Description

Description is based on two strains. Cells are Gram-stain-positive rods, motile at the early stage of incubation. Colonies formed after 5 days incubation on R2A medium at 28 °C are yellow, smooth, circular, convex, and non-transparent, with entire edges. Grows on PCA, nutrient agar, R2A, and LB media. Growth of the type strain occurs in the presence of 0–4% (*w/v*) NaCl (optimum, 1–2% NaCl), at pH 6.0–8.8 (optimum, pH 8.0–8.3) and at 10–40 °C (optimum, 28 °C). Growth of the reference strain occurs at 15–40 °C, 0–6% (*w/v*) NaCl and pH 6.2–8.5 with optimal conditions at 22–28 °C, 2–4% (*w/v*) NaCl and pH 6.6–7.5. Catalase-positive and oxidase-negative. Cells are positive for the following enzyme activities: esterase (C4), N-acetyl- β -glucosaminidase, α -galactosidase, β -galactosidase, α -glucosidase, β -glucosidase, β -glucuronidase, naphthol-AS-BI-phosphohydrolase, leucine arylamidase, esterase lipase (C8), lipase (C14), valine arylamidase, acid phosphatase, and cystine arylamidase, but negative for trypsin, α -chymotrypsin, alkaline phosphatase, α -mannosidase, urease, and α -fucosidase. Does not produce indole or H₂S, but NH₃ is produced from peptone. Negative for the methyl red test and Voges-Proskauer reaction. Reduces nitrate to nitrite in a medium with acetate, but does not reduce nitrate to N₂. Chemoorganoheterotrophic, facultatively anaerobic. In aerobic conditions utilizes peptone, yeast extract, acetate, and pyruvate; acid is produced from aesculin (Fe citrate), L-arabinose, arbutin, D-cellobiose, D-fructose, D-galactose, D-glucose, glycogen, D-mannose, salicin, D-sucrose, D-xylose, D-maltose, D-mannitol, starch, D-trehalose, D-turanose, and N-acetylglucosamine, but not from D-adonitol, amygdalin, D-arabinose, D-arabite, L-arabite, dulcitol, erythritol, D-fucose, L-fucose, inositol, inulin, D-lactose, D-lyxose, methyl- β D-xylopyranoside, methyl- α D-mannopyranoside, methyl- α D-glucopyranoside, D-melibiose, D-melicitose, gluconate K, 2-ketogluconate K, 5-ketogluconate K, D-ribose, L-rhamnose, D-sorbitol, L-sorbose, D-tagatose, xylitol, and L-xylose. In API 20E tests, positive for fermentation/oxidation of glucose, sucrose, and arabinose; and negative for arginine dihydrolase, lysine decarboxylase, ornithine decarboxylase, L-tryptophan deaminase, gelatinase, citrate utilization, indole production, and fermentation/oxidation of inositol, sorbitol, and rhamnose. Acetic, propionic, isobutyric, and iso-valeric acids and CO₂ are the major products of anaerobic glucose fermentation. Hydrogen is not produced. Sensitive to ampicillin (10 μ g), chloramphenicol (30 μ g), penicillin (10 μ g), ciprofloxacin (5 μ g), and erythromycin (15 μ g), but resistant to gentamicin (10 μ g) and kanamycin (30 μ g). The peptidoglycan type is A4 β , containing L-Orn (Lys)-D-Ser-D-Glu. The major cell-wall sugar is rhamnose; galactose, mannose, and glucose are also present. Major fatty acids (>5%) are *anteiso*-C15:0, C14:0, C16:0, and C15:0. The major menaquinone is MK-9(H4). The major polar lipids are diphosphatidylglycerol, unidentified glycolipids, and phosphoglycolipids.

The type strain is HO-Ch2T, isolated from the methanogenic enrichment obtained from a petroleum reservoir (Nurlat, Russia). The DNA G + C content of the genome of the type strain HO-Ch2T is 73.4% and the genome size is 4.0 Mb. The EMBL/GenBank accession numbers for the 16S rRNA gene sequence and genome sequence of strain HO-Ch2T are MT225794 and GCA_008364845.1, respectively.

Classification

Bacteria » *Actinomycetota* » *Actinomycetes* » *Micrococcales* » *Cellulomonadaceae* » *Actinotalea* » *Actinotalea subterranea*

References

Effective publication: Semenova et al., 2022 [2]

Registry URL

<https://seqco.de/i:39440>

References

1. Semenova et al. (2022). Correction: Semenova et al. Physiological and Genomic Characterization of *Actinotalea subterranea* sp. nov. from Oil-Degrading Methanogenic Enrichment and Reclassification of the Family Actinotaleaceae. *Microorganisms* 2022, 10, 378. *Microorganisms*. [DOI:10.3390/microorganisms10050862](https://doi.org/10.3390/microorganisms10050862)
2. Semenova et al. (2022). Physiological and Genomic Characterization of *Actinotalea subterranea* sp. nov. from Oil-Degrading Methanogenic Enrichment and Reclassification of the Family Actinotaleaceae. *Microorganisms*. [DOI:10.3390/microorganisms10020378](https://doi.org/10.3390/microorganisms10020378)

Register List Certificate of Validation

On behalf of the *Committee on the Systematics of Prokaryotes Described from Sequence Data* (SeqCode Committee), we hereby certify that the Register List **seqco.de/r:bk0hdqw0** submitted by **Grouzdev, Denis** and including 1 new name has been successfully validated.

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