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Draft genome sequence of *Thermoactinomyces* sp. Gus2-1 isolated from the hot-spring Gusikha in Bargusín Valley (Baikal Rift Zone, Russia)



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ABSTRACT

The *Thermoactinomyces* sp. strain Gus2-1 was isolated from hot-spring sediments sample from the hot-spring Gusikha in Bargusín Valley (Baikal Rift Zone, Russia). The sequenced and annotated genome is 2,623,309 bp and encodes 2513 genes. The draft genome sequence of the *Thermoactinomyces* sp. strain Gus2-1 has been deposited at DDBJ/EMBL/GenBank under the accession JPZM01000000 and the sequences could be found at the site <https://www.ncbi.nlm.nih.gov/nucore/JPZM01000000>.

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Specifications

Organism/cell line/tissue	<i>Thermoactinomyces</i> sp. Gus2-1
Sex	–
Sequencer or array type	Ion PGM™ Template OT2 400 kits
Data format	Processed
Experimental factors	Bacteria
Experimental features	Whole genome sequence of <i>N. lepida</i> , assembly and annotation
Consent	Level of consent allowed for reuse if applicable
Sample source location	The hot-spring Gusikha (60 °C) in Bargusín Valley (Baikal Rift Zone, Russia)

1. Direct link to deposited data

<https://www.ncbi.nlm.nih.gov/nucore/JPZM01000000>.

2. Introduction

The genus *Thermoactinomyces* is one of the earliest known Actinomycete taxa, and the type species of this genus is *Thermoactinomyces vulgaris* [1]. The members of this genus are aerobic, endospore-forming, Gram-positive bacteria belonging to the order *Bacillales* [2]. They produce endospores that are formed endogenously inside the aerial and substrate hyphae of the bacteria [3]. Currently, researchers are finding new species belonging to this genus [4].

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3. Strain isolation

The strain Gus2-1 was isolated from sediments sample from the hot-spring Gusikha (60 °C) in Bargusín Valley (Baikal Rift Zone, Russia). *Thermoactinomyces* sp. Gus2-1 culture was cultivated in liquid medium containing 1% trypton, 0.5% yeast extract, and 3.5 M of NaCl. Eight ml of cell culture were pelleted by centrifugation and resuspended in 75 µl of H₂O by intense pipetting.

4. DNA isolation and sequencing

DNA was isolated using the DNA Purification Kit (Fermentas). Ion PGM™ Template OT2 400 and Ion PGM™ Template OT2 400 kits were used to create libraries for genome sequencing. Genome sequencing was performed on an IonTorrent platform (Applied Biosystems) using Ion PGM™ Sequencing 400 Kit in the SBRAS Sequencing Center.

5. Genome assembly and annotation

De novo assembly of short reads into contigs was performed using MIRA v. 4. Contigs shorter than 1000 bp were deleted. A total of 92 contigs yielded a genome sequence 2,623,309 bp long, and the G + C content is 48.01%. ORF prediction and automatic annotation was performed using NCBI PGAAP (http://www.ncbi.nlm.nih.gov/genome/annotation_prok). The complete genome sequence contained 2513 genes, 2315 CDS, 14 rRNAs (5S, 16S, 23S), 76 tRNAs, one ncRNA.

6. Phylogenetic analysis

Phylogenetic analysis was performed using 16S rRNA sequences with the UPGMA algorithm implemented in MEGA v.6. 16S rRNA

sequences of *Thermoactinomyces* type strains were found using the StrainInfo (www.straininfo.net) and GenBank (www.ncbi.nlm.nih.gov/nucleotide) databases. According to phylogenetic analysis, the *Thermoactinomyces* sp. strain Gus2-1 is most closely related to *Thermoactinomyces vulgaris*.

7. Nucleotide sequence accession numbers

The draft genome sequence for *Thermoactinomyces* sp. strain Gus2-1 has been deposited in DDBJ/EMBL/Genbank under the accession no. JPZM01000000. The 92 contigs have been deposited under accession no. JPZM01000001-JPZM01000092.

Conflict of interest

The authors declare that there is no conflict of interests with respect to the work published in this paper.

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