



# Draft Genome Sequence of *Desemzia* sp. Strain C1, Producing Hydrogen Peroxide, Isolated from Oil-Contaminated Soil

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**ABSTRACT** Here, we report the draft genome sequence of *Desemzia* sp. strain C1, which was isolated from oil-contaminated soil in South Korea and produces hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>). The genome of *Desemzia* sp. strain C1 contains genes encoding various oxidases involved in H<sub>2</sub>O<sub>2</sub> production and resistance to oxidative stress.

The genus *Desemzia* includes Gram-positive, non-spore-forming, microaerophilic bacteria (1). The information on the *Desemzia* genus is limited because further novel species have not been reported since the first report by Steinhaus in 1941 (2).

Oil-contaminated soil was collected after removal with a spatula of surface soil (depth of 3 cm) from an auto repair shop in Gwangju, South Korea (35°12'22.3"N, 126°54'01.6"E). *Desemzia* sp. strain C1, producing hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>), was isolated based on the Prussian blue zone-forming reaction of ferric cyanide and H<sub>2</sub>O<sub>2</sub> produced by bacteria (3). Bacteria producing H<sub>2</sub>O<sub>2</sub> were grown in brain heart infusion (BHI) broth (BD BBL, Sparks, MD, USA) and Trypticase soy broth containing 3 g yeast extract (TSBY) (BD Difco) at 30°C under static conditions, and the H<sub>2</sub>O<sub>2</sub> produced was quantified using the Amplex Red hydrogen peroxide/peroxidase assay kit (Invitrogen, Waltham, MA, USA). *Desemzia* sp. strain C1 produced a maximum of 0.23 mM H<sub>2</sub>O<sub>2</sub> in BHI broth, which was 4 times higher than that of *Streptococcus oralis* KACC 13048<sup>T</sup>, a well-known H<sub>2</sub>O<sub>2</sub> producer, in TSBY (Fig. 1). Therefore, we sequenced the whole genome of *Desemzia* sp. strain C1 to identify genes related to H<sub>2</sub>O<sub>2</sub> production.

Genomic DNA (gDNA) was extracted using the MagAttract high-molecular-weight (HMW) DNA kit (Qiagen, Hilden, Germany) after culture in BHI broth for 36 h at 30°C under static conditions. Extracted gDNA was sheared using the Megaruptor 3 (Diagenode SA, Liège, Belgium), and small fragments of less than 3 kb were removed using AMPure XP beads (Beckman Coulter, Pasadena, CA, USA). The DNA library was constructed by using the SMRTbell Express template preparation kit v2.0 (Pacific Biosciences [PacBio], Menlo Park, CA, USA) (4). The SMRTbell library was sequenced using the Sequel Sequencing kit v3.0 (PacBio) and a SMRT Cell 1M v2 (PacBio), resulting in 344,884 reads (*N*<sub>50</sub>, 8,836 bp). The draft genome of *Desemzia* sp. strain C1 was constructed based on PacBio sequencing data (5). Sequencing analysis was carried out at CJ Bioscience (Seoul, South Korea). PacBio sequencing data were assembled with SMRT Link v10.1.0.119588 according to the microbial assembly protocol (PacBio). All procedures were implemented according to the manufacturer's protocols. Default parameters were used for all software unless otherwise specified. The resulting draft genome (average coverage, 611.0×) contained three contigs of 2,790,095 bp (*N*<sub>50</sub>, 2,697,877 bp), with an overall G+C content of 38.7%. The genome was annotated by the NCBI Prokaryotic Genome Annotation Pipeline (PGAP) using the best-placed reference protein set method (GeneMarkS-21) (6). Genome annotation revealed 2,582 coding DNA sequences (CDSs) and 108 RNA sequences (22 rRNA genes and 86 tRNA genes). The genome of *Desemzia* sp. strain C1 contains putative genes encoding various oxidases involved in H<sub>2</sub>O<sub>2</sub> production, such as lactate oxidase (LOX), pyruvate oxidase, and three distinct NADH oxidases (7, 8).

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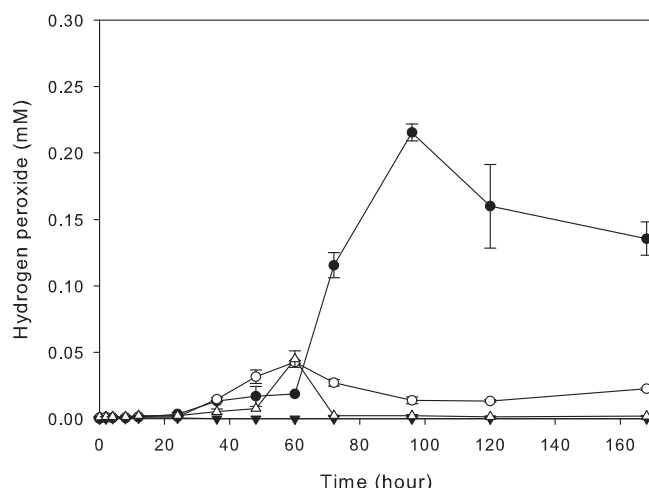
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**FIG 1** Production of hydrogen peroxide ( $\text{H}_2\text{O}_2$ ) by *Desemzia* sp. strain C1 and *Streptococcus oralis*. Data for *Desemzia* sp. strain C1 cultured in BHI broth (●) and TSBY (○) and *S. oralis* cultured in BHI broth (▼) and TSBY (△) are shown.

The current genome information can shed light on the understanding of  $\text{H}_2\text{O}_2$  production and resistance mechanisms in the bacterial system. In addition, the  $\text{H}_2\text{O}_2$ -producing putative gene encoding LOX could be a good candidate involved in the bacterial enzyme-mediated advanced oxidation processes to apply for degradation and detoxification of various organic pollutants.

**Data availability.** The draft genome of *Desemzia* sp. strain C1 has been deposited in GenBank under the BioProject accession number [PRJNA777376](https://doi.org/10.1099/00207713-49-1-185), the BioSample accession number [SAMN22852026](https://doi.org/10.1099/00207713-49-1-185), and the GenBank accession number [JAJIZP000000000](https://doi.org/10.1099/00207713-49-1-185). The raw reads can be accessed under the SRA accession number [SRR17868029](https://doi.org/10.1099/00207713-49-1-185). The version described in this paper is the first version, [JAJIZP010000000](https://doi.org/10.1099/00207713-49-1-185).

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