

# Draft genome of *Paenibacillus ottowii* LIS04 and *Bacillus velezensis* IM14: dual biocontrol bacterial strains with broad-spectrum antifungal activity

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**ABSTRACT** *Paenibacillus* and *Bacillus* are gram-positive bacteria known for promoting plant growth and controlling agriculturally important phytopathogens. This study highlights the genomic traits of *Paenibacillus ottowii* LIS04 and *Bacillus velezensis* IM14, isolated from sorghum seeds and maize stigmas, respectively, with potential applications in developing innovative agricultural bioproducts.

**KEYWORDS** biological control, plant growth-promoting bacteria, bioproduct, genomics

*Paenibacillus ottowii* LIS04 and *Bacillus velezensis* IM14 are potent antagonists against several plant pathogens and exhibit plant growth-promoting properties (1–3). These strains produce antifungal secondary metabolites, including iturin, fengycin, and surfactin from *B. velezensis* IM14 and fusaricidin from *P. ottowii* LIS04 (4). Their compatibility highlights the potential for combined application in developing innovative bioproducts.

*P. ottowii* LIS04 was isolated from sorghum (*Sorghum bicolor* L.) seeds collected in the Cerrado biome (Brazilian savanna) of Sete Lagoas (19.28°S, 44.14°W), Minas Gerais, Brazil. The seeds were surface-disinfected and incubated in germination boxes at 22°C with a 12 hour photoperiod. The strain was obtained from seeds exhibiting bacterial colonies that visibly inhibited the mycelial growth of phytopathogenic fungi. Bacterial samples were collected using sterile platinum loops and streaked onto Tryptone Soy Agar (Kasvi, Brazil) plates. *B. velezensis* IM14 strain was isolated from maize (*Zea mays* L.) stigmas collected in the Cerrado region of Sete Lagoas. The stigmas were macerated and suspended in sterile saline solution (0.9%). Serial dilutions were plated on Potato Dextrose Agar and incubated at 28°C for 48 hours. Both strains were cryopreserved in a glycerol-based medium at –80°C prior to genome sequencing. *Paenibacillus ottowii* LIS04 and *B. velezensis* IM14 were deposited in the Embrapa Collection of Multifunctional and Phytopathogenic Microorganisms for Maize and Sorghum (CMMF) under accession BRM053425 (CMPC2456) and BRM046334 (CMPC2048), respectively.

The DNA extraction of the bacterial isolates was performed using pure cultures grown in Luria Bertani liquid medium at 28°C for 24 hours under agitation at 150 rpm. After this period, the culture was centrifuged at 16,000 rpm, and the supernatant was discarded. Genomic DNA from LIS04 and IM14 was extracted using the Wizard Genomic DNA Purification Kit (Promega, USA) and quantified with a Qubit 2.0 fluorometer (Life Technologies, USA). Whole-genome sequencing (WGS) libraries were prepared according to the BGISEQ-500 WGS protocol described by Huang et al. (5). Genome sequencing was performed on the Illumina HiSeq 4000 platform (Illumina, USA) at the Beijing Genomics

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The authors declare no conflict of interest.

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TABLE 1 Genomic features of *Paenibacillus ottowii* LIS04 and *Bacillus velezensis* IM14

Assembly	<i>P. ottowii</i>	<i>B. velezensis</i>
Strain ID	LIS04	IM14
Fast ANI placement (%)	96866	98296
Fast ANI reference (NCBI)	GCA_006874425.1	GCA_001461825.1
Library size	6,832,602	6,847,781
Number of contigs	164	55
Total sequence length (pb)	5,553,857	3,997,051
Total ungapped length (bp)	5,553,759	3,996,960
N50 (kb)	473,395	468,182
G+C content (%)	45.5	46.5
Genes	5,209	3,977
Protein-coding	4,932	3,794
Noncoding (RNA)	1	2
Genome completeness/contamination (%)	99.7/0.56	99.41/0
Coverage (×)	347	489
SRA identifiers	<a href="https://www.ncbi.nlm.nih.gov/sra/SRS23177414">SRS23177414</a>	<a href="https://www.ncbi.nlm.nih.gov/sra/SRS23197145">SRS23197145</a>
Genome assembly (NCBI)	<a href="https://www.ncbi.nlm.nih.gov/genome/assembly/GCF_030676815.1">GCF_030676815.1</a>	<a href="https://www.ncbi.nlm.nih.gov/genome/assembly/GCF_030676595.1">GCF_030676595.1</a>

Institute (BGI), Shenzhen, China. Sequencing was performed using a 150 bp paired-end strategy. Raw reads were processed with Trimmomatic v0.38 (6) to remove low-quality and adapter sequences (Phred score <20). The quality-filtered reads were then assembled *de novo* using SPAdes v3.12.0 (7). Assembly quality was assessed with QUAST v5.0.2 (8), and genome completeness was evaluated using BUSCO v5.3.1 (Benchmarking Universal Single-Copy Orthologs) (9).

The key genomic features of LIS04 and IM14 are summarized in Table 1. The genomic sequences were submitted to NCBI and annotated using the Prokaryotic Genome Annotation Pipeline v6.6.1 (10). The annotation process identified 4,928 coding sequences, 92 rRNAs, and 109 tRNAs in LIS04, while IM14 contained 3,786 coding sequences, 22 rRNAs, and 84 tRNAs.

For taxonomic inference based on genomic data, the Type (Strain) Genome Server (<https://tygs.dsmz.de>) was utilized to identify the *Paenibacillus* and *Bacillus* species most closely related to LIS04 and IM14. The results confirmed that LIS04 belongs to *P. ottowii*, while IM14 corresponds to *B. velezensis*.

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Gisele de Fátima Dias Diniz, Conceptualization, Data curation, Formal analysis, Investigation, Methodology, Software, Validation, Visualization, Writing – original draft, Writing – review and editing | Luciano Viana Cota, Conceptualization, Formal analysis, Funding acquisition, Project administration, Resources, Supervision, Validation, Visualization | Ubiraci Gomes de Paula Lana, Data curation, Formal analysis, Methodology, Resources, Software, Validation, Visualization | José Edson Fontes Figueiredo, Conceptualization, Data curation, Formal analysis, Methodology, Writing – original draft, Writing – review and editing | Lourenço Vitor Silva Ferreira, Data curation, Methodology, Software, Visualization | Vera Lúcia dos Santos, Conceptualization, Formal analysis, Funding acquisition, Supervision, Validation, Visualization | Christiane Abreu de Oliveira-Paiva, Conceptualization, Formal analysis, Funding acquisition, Investigation, Project administration, Resources, Supervision, Validation, Visualization, Writing – original draft, Writing – review and editing.

## DATA AVAILABILITY

The draft genome sequences of *Paenibacillus ottowii* LIS04 and *Bacillus velezensis* IM14 strains were deposited in GenBank BioProject [PRJNA1001033](https://www.ncbi.nlm.nih.gov/bioproject/PRJNA1001033). The Whole Genome Shotgun project has been deposited at DDBJ/ENA/GenBank under the accession nos. [NZ\\_JAUUUV000000000.1](https://www.ncbi.nlm.nih.gov/nuccore/NZ_JAUUUV000000000.1) and [NZ\\_JAUUUY000000000.1](https://www.ncbi.nlm.nih.gov/nuccore/NZ_JAUUUY000000000.1), with BioSample IDs [SAMN36791032](https://www.ncbi.nlm.nih.gov/biosample/SAMN36791032) and [SAMN36791029](https://www.ncbi.nlm.nih.gov/biosample/SAMN36791029), for LIS04 and IM14, respectively. The raw sequencing reads for LIS04 and IM14 have been submitted to the SRA database and are available under accession numbers [SRS23177414](https://www.ncbi.nlm.nih.gov/sra/SRS23177414) and [SRS23197145](https://www.ncbi.nlm.nih.gov/sra/SRS23197145). The versions presented here represent the first releases of the LIS04 and IM14 genomes.

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