

Complete Genome Sequence of Lytic Bacteriophage LZ35 Infecting *Acinetobacter baumannii* Isolates

Zhonghe Guo, Honglan Huang, Xiaolin Wu, Yuchong Hao, Yanbo Sun

Department of Pathogen Biology, College of Basic Medical Sciences, Jilin University, Changchun, People's Republic of China

***Acinetobacter baumannii* is a Gram-negative opportunistic pathogen that is frequently associated with nosocomial infections. Bacteriophages infecting *A. baumannii* can be used as effective agents to control these infections. Here, we announce the complete genome sequence of the lytic bacteriophage LZ35 infecting *A. baumannii* isolates.**

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Address correspondence to Yanbo Sun, sunyb@jlu.edu.cn.

Acinetobacter baumannii, a nonfermentative, aerobic, Gram-negative bacillus, is a significant cause of opportunistic infections, such as ventilator-associated pneumonia, urinary tract infections, and bacteremia in intensive care units (1). Due to the increasing resistance to many classes of antibiotics among *A. baumannii* strains (2, 3), phage therapy (4, 5), an application of bacteriophages (phages) as an antibacterial agent, may be an alternative potential for controlling *A. baumannii* infection. Here, we present the complete genome sequence of lytic bacteriophage LZ35, which is infectious to *A. baumannii*.

Bacteriophage LZ35 was isolated from sewage effluents from the Jilin University First-Affiliated Hospital. The morphology of LZ35 was determined using transmission electron microscopy and demonstrated a 47-nm head diameter and a 56-nm-long contractile tail, which are characteristics of members of the family *Myoviridae*. Phage DNA was sequenced in an Illumina MiSeq 250-bp paired-end run with a 546-bp insert library at Suzhou Genewiz Biological Technology Co., Ltd. (Suzhou, People's Republic of China). After trimming with Trimmomatic (<http://usadellab.org/cms/?page=trimmomatic>) and removal of low-quality ends, 3,592,478 reads comprising a mean coverage of 17,647-fold were *de-novo* assembled using SSPACE (<http://baseclear.com/genomics/bioinformatics/basetools/SSPACE>) and GapFiller (<http://baseclear.com/genomics/bioinformatics/basetools/gapfiller>). Gene prediction and annotation of the phage LZ35 genome was performed using the RAST server (6). Phage LZ35 contained a 44,885-bp double-stranded DNA and a G+C content of 37.95%, with no tRNAs identified. A total of 83 coding sequences were identified. The analysis of the LZ35 genomic sequence reveals 78% and 77% matches with 97% and 99% identity to the *Acinetobacter* phage IME-AB2 (accession no. JX976549.1) (7) and *Acinetobacter* phage YMC-13-01-C62 (accession no. KJ817802.1) genomes, respectively.

The genomic data of lytic phage LZ35 will prove useful for comparative studies and for the future investigation of LZ35 acting as a potential alternative for controlling *A. baumannii* infection.

Accession number(s). The complete genome sequence of *A. baumannii* phage LZ35 has been submitted to the GenBank database under the accession number [KU510289](https://www.ncbi.nlm.nih.gov/nuclseq/KU510289). The version described in this paper is the first version, KU510289.1.

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