Complete Genome Sequences of 10 Lactococcal Skunavirus Phages Isolated from Cheddar Cheese Whey Samples in Canada

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ABSTRACT We report the complete genome sequences of 10 virulent phages of the Skunavirus genus (Siphoviridae) that infect Lactococcus lactis strains used for cheddar cheese production in Canada. Their linear genomes range from 28,969 bp to 31,042 bp with GC contents of 34.1 to 35.1% and 55 to 60 predicted open reading frames (ORFs).

Lactococcus lactis strains are added to milk to manufacture a wide variety of cheeses worldwide. The most common cause of slow milk fermentation, which leads to low-quality fermented products, is virulent phages infecting these strains (1). Lactococcal phages are classified into several groups (2), with phages belonging to the Skunavirus genus (formerly 936) being the most common (3). Constant phage monitoring in dairy factories is essential for adapting antiphage measures and preventing fermentation failure (4). Here, we report the genomic characterization of 10 new virulent phages (FB3, FB6, FB10, FB14, GL7, GP13, GP14, GP15, RH6, and RH10) of the Skunavirus genus. Phages were isolated from 2007 to 2019 from whey samples obtained from a Canadian cheddar cheese factory.

L. lactis strains were grown at 30°C in M17 medium with 0.5% (wt/vol) lactose (LM17). Phages were isolated using the double-layer plaque assay (5) on LM17 medium supplemented with 10 mM CaCl2. Phage genomic DNA was extracted using phenol-chloroform (6) from high-titer (>10⁹ PFU/ml) filtered (0.45-μm filter) lysates. Sequencing libraries were prepared using a Nextera XT DNA library preparation kit and sequenced with Illumina MiSeq (250-nucleotide paired-end reads). Reads were cleaned using Trimomatic v0.36 (7) and assembled to obtain circular complete sequences using Ray v3.0.1 (8) with k-mer sizes of 21, 31, 41, 51, 71, and 91 and SPAdes v3.13 (9). Open reading frames were predicted using GeneMark (prokaryotic) v3.25 (10), the PECAAN annotation tool (https://discover.kbrinsgd.org/autoannotate/), and Geneious v11.1.5 (11), with the following principles: genes started with ATG, GTG, or TTG codons and were preceded by a Shine-Dalgarno sequence similar to 5’-AGAAAGGAGGT-3’ (12). Coding sequences of 30 or more amino acids were annotated, and deduced proteins were searched for function using BLAST v2.10.0 and a cutoff E value of 0.001. We searched tRNAs with ARAGORN v1.2.38 (13) and tRNAscan-SE 2.0 (14). Annotations were also manually curated by comparing them with other Skunavirus genomes. Unless defined, default parameters were used for all software.

The genome size (from 28,969 to 31,042 bp), number of predicted open reading frames (ORFs) (55 to 60), and GC content (34.1 to 35.1%) for each phage are reported in Table 1. The percentage of deduced proteins with an assigned function ranged from


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<table>
<thead>
<tr>
<th>Phage name</th>
<th>Isolation mo-yr</th>
<th>L. lactis host strain</th>
<th>Genome length (bp)</th>
<th>No. of ORFs</th>
<th>GC content (%)</th>
<th>GenBank/SRA accession no.</th>
<th>No. of reads</th>
<th>Most closely related phage(s)</th>
<th>ANI (%)</th>
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<td>34.8</td>
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</table>

*a* This study.

*b* NA, no country available.
32.1 to 39.0%. Cos sites were found in the 10 phages by sequence homology and were identical (5’-CACAAGGACT-3’) to other Skunavirus phages (15). The average nucleotide identities (ANI) were calculated with a BLAST+ analysis in JSpeciesWS v3.7.3 (16) and are listed in Table 1.

Phages FB10, GL7, GP13, GP14, GP15, RH6, and RH10 possess an early-expressed gene that codes for a methyltransferase. These methylases likely protect the viral genome against a specific host endonuclease during intracellular replication. Phage FB14 also carries a methyltransferase-coding gene, but it is located in the late-expressed region. This methylase may perform regulatory functions (17, 18). Six genomes contained tRNA-Pro and tRNA-Trp. Phages GP13 and RH10 had only tRNA-Pro, and phages GP14 and GP15 did not carry any tRNA.

**Data availability.** The phages are available at www.phage.ulaval.ca. The genome sequences and raw data are available under the GenBank and SRA accession numbers reported in Table 1.

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**REFERENCES**


