



Draft Genome Sequence of a Highly Virulent Rabbit Staphylococcus aureus Strain

Zoltán Német,^a Ervin Albert,^a Tibor Nagy,^b Ferenc Olasz,^b Endre Barta,^b János Kiss,^b Ádám Dán,^c Krisztián Bányai,^d Katleen Hermans,^e Imre Biksi^a

Faculty of Veterinary Science, Szent István University, Department and Clinic for Production Animals, Üllő, Hungary^a; Innovation Centre and Agricultural Biotechnology Institute of the National Agricultural Research, Gödöllő, Hungary^b; Veterinary Diagnostic Directorate, National Food Chain Safety Office (NFCSO), Budapest, Hungary^c; Institute for Veterinary Medical Research, Hungarian Academy of Sciences, Budapest, Hungary^d; Department of Pathology, Bacteriology and Avian Diseases, Ghent University, Ghent, Belgium^e

We report the draft genome sequence of *Staphylococcus aureus* Sp17, a typical highly virulent (HV) rabbit strain. As current medicine apparently fails to effectively reduce disease and economical losses caused by this organism, it is essential to gain better insight on its genomic arrangement.

Received 4 June 2015 Accepted 5 June 2015 Published 9 July 2015

Citation Német Z, Albert E, Nagy T, Olasz F, Barta E, Kiss J, Dán Á, Bányai K, Hermans K, Biksi I. 2015. Draft genome sequence of a highly virulent rabbit *Staphylococcus aureus* strain. Genome Announc 3(4):e00461-15. doi:10.1128/genomeA.00461-15.

Copyright © 2015 Német et al. This is an open-access article distributed under the terms of the Creative Commons Attribution 3.0 Unported license.

Address correspondence to Zoltán Német, nemet.zoltan@aotk.szie.hu.

S taphylococcosis is a disease having major economic impact on industrial rabbit meat production. Infections caused by highly virulent (HV) *Staphylococcus aureus* strains result in severe clinical conditions; these strains are also frequently resistant to antimicrobials. An outbreak of an HV *S. aureus* strain hinders profitable production and frequently necessitates culling the entire flock (1).

A typical HV strain identified as Sp17, originating from a Spanish rabbit farm with severe staphylococcal mastitis problems, was used for whole-genome sequencing (WGS). This strain belongs to the typical highly virulent rabbit *S. aureus* clone, as it shows the mixed biotype CV-C (2), is sensitive to phages of phage group II (3A, 3C, and 71), shows the multiplex PCR pattern specific for highly virulent *S. aureus* strains (3), and has pulsed-field gel electrophoresis type N2 and *spa* type t645 (4).

Total DNA of the strain was subjected to 2×300 -bp pairedend Illumina MiSeq sequencing at the Department of Biochemistry, Faculty of Medicine, University of Szeged, Hungary. A total of 3.96 million read pairs were recorded, and the estimated coverage of the whole genome is $700 \times$.

The estimated coverage of the subsets of reads was adjusted to $30 \times$, and they were assembled *de novo* using MIRA version 4.0.2 (5), A5 pipeline version 20130326 (6), and SeqMan NGen version 4.1.2 build 25 (DNAStar version 10). Scaffolds were built from different assemblies using Mauve version 2.3.1 (7) as a Geneious version 8.1.2 (8) plugin. This resulted in a total of 10 scaffolds containing 2,684,832 nucleotides. The average G+C content is 32.7%.

Scaffolds were submitted to the RAST annotation server (9). The taxon was set to "*Staphylococcus aureus*" (1280.2034), the genetic code to "11 (*Archaea, Bacteria*)," the annotation scheme to "ClassicRAST," and "preserve gene calls," "automatically fix errors," "fix frameshifts," and "backfill gaps" to "no." We obtained 2,395 annotated genes, 50 tRNAs, and 5 rRNAs.

A search for similar sequences deposited in GenBank was conducted using BLAST. The best match was *S. aureus* subsp. *aureus* 21337 (accession no. NZ_JHPZ0000000.1). Pairwise alignment showed 98.56% similarity of these sequences, which indicates a close relationship between these strains.

A detailed analysis of the virulence genes in these sequences will be conducted to identify key elements responsible for the remarkable pathogenicity of this strain, in order to facilitate the development of effective treatment methods and/or preventive measures against HV *S. aureus* infections in rabbits.

Nucleotide sequence accession numbers. This whole-genome shotgun project has been deposited at DDBJ/EMBL/GenBank under the accession no. LBCS00000000. The version described in this paper is version LBCS01000000.

ACKNOWLEDGMENTS

This research was financed by a research fund granted by Tetrabbit Ltd., Baja, Hungary.

We acknowledge NIIF for awarding us access to resources based in Hungary at Szeged.

Publication of the results of this study was supported by the 9877-3/ 2015/FEKUT grant of the Hungarian Ministry of Human Resources.

REFERENCES

- Hermans K, Devriese LA, Haesebrouck F. 2003. Rabbit staphylococcosis: difficult solutions for serious problems. Vet Microbiol 91:57–64. http:// dx.doi.org/10.1016/S0378-1135(02)00260-2.
- Devriese LA. 1984. A simplified system for biotyping *Staphylococcus aureus* strains isolated from animal species. J Appl Bacteriol 56:215–220. http:// dx.doi.org/10.1111/j.1365-2672.1984.tb01341.x.
- Vancraeynest D, Haesebrouck F, Hermans K. 2007. Multiplex PCR assay for the detection of high virulence rabbit *Staphylococcus aureus* strains. Vet Microbiol 121:368–372. http://dx.doi.org/10.1016/j.vetmic.2006.12.011.
- Vancraeynest D, Haesebrouck F, Deplano A, Denis O, Godard C, Wildemauwe C, Hermans K. 2006. International dissemination of a high virulence rabbit *Staphylococcus aureus* clone. J Vet Med B Infect Dis Vet Public Health 53:418–422. http://dx.doi.org/10.1111/j.1439-0450.2006.00977.x.
- Chevreux B, Wetter T, Suhai S. 1999. Genome sequence assembly using trace signals and additional sequence information, p 45–56. *In* Computer science and biology. Proceedings of the German Conference on Bioinformatics, GCB '99. GCB, Hannover, Germany.

- Tritt A, Eisen JA, Facciotti MT, Darling AE. 2012. An integrated pipeline for *de novo* assembly of microbial genomes. PLoS One 7:e42304. http:// dx.doi.org/10.1371/journal.pone.0042304.
- 7. Darling AE, Mau B, Perna NT. 2010. progressiveMauve: multiple genome alignment with gene gain, loss and rearrangement. PLoS One 5:e11147. http://dx.doi.org/10.1371/journal.pone.0011147.
- 8. Kearse M, Moir R, Wilson A, Stones-Havas S, Cheung M, Sturrock S, Buxton S, Cooper A, Markowitz S, Duran C, Thierer T, Ashton B, Meintjes P, Drummond A. 2012. Geneious basic: an integrated and ex-

tendable desktop software platform for the organization and analysis of sequence data. Bioinformatics 28:1647–1649. http://dx.doi.org/10.1093/bioinformatics/bts199.

 Aziz RK, Bartels D, Best AA, DeJongh M, Disz T, Edwards RA, Formsma K, Gerdes S, Glass EM, Kubal M, Meyer F, Olsen GJ, Olson R, Osterman AL, Overbeek RA, McNeil LK, Paarmann D, Paczian T, Parrello B, Pusch GD, Reich C, Stevens R, Vassieva O, Vonstein V, Wilke A, Zagnitko O. 2008. The RAST server: Rapid Annotations using Subsystems Technology. BMC Genomics 9:75. http://dx.doi.org/10.1186/1471-2164-9-75.