





## Genome Sequence of *Enterococcus pernyi*, a Pathogenic Bacterium for the Chinese Oak Silkworm, *Antheraea pernyi*

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We report the draft genome assembly of *Enterococcus pernyi*. The genome sequence is 3.09 Mb in length with a G+C content of 38.35%. It covers 3,153 genes with an average length of 854 bp, and contains 65 tRNAs, 13 small RNAs, and 18 rRNAs. Moreover, it contains 9 genomic islands with an average length of 14,058 bp and 3 prophages with an average length of 37,430 bp.

Received 30 December 2015 Accepted 4 April 2016 Published 19 May 2016

Citation Sun Y, Li X, Wang G, Wang Y, Jiang Y, Liu Y, Yu Z, Qin L. 2016. Genome sequence of *Enterococcus pernyi*, a pathogenic bacterium for the Chinese oak silkworm, *Antheraea pernyi*. Genome Announc 4(3):01764-15. doi:10.1128/genomeA.01764-15.

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he Chinese oak silkworm (Antheraea pernyi [Guérin-Méneville, 1855]), which belongs to Lepidoptera: Saturniidae, is the most well-known wild silkworm for insect food and silk production. Rearing of the Chinese oak silkworm has a history of about 400 years in China (1). Empty-gut disease is one of the most important diseases in A. pernyi, and this disease seriously affects the yield of the tussah cocoon and causes great economic losses. The pathogen of the disease was established as *Streptococcus pernyi* sp. nov., based on morphological, physiological, biochemical, and serological characteristics (2). However, S. pernyi sp. nov. has been reclassified and renamed as Enterococcus pernyi, based on phylogenic analysis of 16S rRNA and tufgene sequences (3–5). Now, the pathogen of empty-gut disease in A. pernyi has been defined as E. pernyi in the taxonomy of the National Center for Biotechnology Information. The genome of E. pernyi was sequenced to gain a better understanding of the taxonomic status of this bacterium and provide more genomic information for further studies to prevent and cure the disease.

The complete genome sequence was determined by Illumina Solexa technology at Novogene Bioinformatics Technology Co., Ltd. (Beijing, China). Sequence assembly was performed using SOAPdenovo version 2.04 (6). Coding sequences (CDSs) were predicted using GeneMarkS software (7) and further annotated into databases through BLASTp, including NCBInr, COG, GO, KEGG, Swiss-Prot, and TrEMBL. tRNAscan (8), RNAmmer (9), and Rfam (10) were used to predict tRNAs, rRNAs, and small RNAs, respectively. Gene islands and prophages were predicted using IslandPath-DIOMB (11) and PHAST software (12).

The genome size of *E. pernyi* is 3.09 Mb with a G+C content of 38.35%. A total of 626 Mb of clean data were generated, reaching a genome coverage depth of over 200-fold. Sequences were assembled into 23 contigs with a total length of 3,181,210 bp (largest, 603,828 bp, and smallest, 654 bp) and with an  $N_{50}$  contig size of 370,188 bp. Finally, there were a total of 9 scaffolds with a total length of 3,188,572 bp (largest, 3,086,269 bp, and smallest, 654 bp) and with an  $N_{50}$  scaffold size of 3,086,269 bp. The genome contains 3,153 CDSs with an average length of 854 bp, which repre-

sent 84.48% of the whole genome. The annotation results showed that only 224 CDSs (7.1%) were not annotated into any databases; there were 2,916, 1,537, 1,577, 1,487, 1,242, and 2,812 CDSs annotated into NCBInr, COG, GO, KEGG, Swiss-Prot, and TrEMBL, respectively. Meanwhile, 65 tRNAs, 18 rRNAs, and 13 small RNAs were identified. Furthermore, the genome contains 9 genomic islands with an average length of 14,058 bp, and contains 3 prophages with an average length of 37,430 bp. There was no clustered regularly interspaced short palindromic repeat identified in the genome.

**Nucleotide sequence accession number.** The whole-genome sequences of *E. pernyi* have been deposited at DDBJ/EMBL/GenBank under the accession number LPVT00000000.

## **ACKNOWLEDGMENTS**

This work was supported by the National Modern Agriculture Industry Technology System Construction Project (silkworm and mulberry) (CARS-22), the Cultivation Plan for Youth Agricultural Science and Technology Innovative Talents of Liaoning Province (2014040), the Scientific Research Project for the Education Department of Liaoning Province (2014476), the Magnitude Science and Technology Projects of Liaoning Province, and the Liaoning Province Climbing Scholars Support Plan (1102).

## **FUNDING INFORMATION**

This work, including the efforts of Li Qin, was funded by National Modern Agriculture Industry Technology System Construction Project (Silkworm and Mulberry) (CARS-22). This work, including the efforts of Zhiguo Yu, was funded by Liaoning Province Climbing Scholars Support Plan (1102). This work, including the efforts of Yiren Jiang, was funded by Cultivation Plan for Youth Agricultural Science and Technology Innovative Talents of Liaoning Province (2014040). This work, including the efforts of Yiren Jiang, was funded by Scientific Research Project for Education Department of Liaoning Province (2014476). This work, including the efforts of Li Qin, was funded by Magnitude Science and Technology Projects of Liaoning Province.

The funders had no role in study design, data collection and interpretation, or the decision to submit the work for publication.

## **REFERENCES**

- 1. Liu Y, Li Y, Li X, Qin L. 2010. The origin and dispersal of the domesticated Chinese oak silkworm, *Antheraea pernyi*, in China: a reconstruction based on ancient texts. J Insect Sci 10:180. http://dx.doi.org/10.1673/031.010.14140.
- Wang DS, Zhan LK, Yu XB, Zhang CF. 1980. Studies on empty-gut disease of tussah II. Identification of the causative agent of empty-gut disease of tussah, *Streptococcus pernyi* sp. nov. Acta Microbiol Sin 20: 225–229. http://dx.doi.org/10.13343/j.cnki.wsxb.1980.03.001.
- Wang LL, Qin F, Song C, Zhou ZY. 2010. Reclassification of the pathogen for empty-gut disease of Chinese oak silkworm, *Antheraea pernyi*. J Food Agric Environ 8:156–158.
- 4. Shang CF, Qin L, Zhao ZJ, Song C, Li SY, Jiang DF, Fan Q. 2011. Identification of the pathogen for *Antheraea pernyi* empty-gut disease by using 16S rRNA gene sequence. Sci Sericulture 37:931–936. http://dx.doi.org/10.13441/j.cnki.cykx.2011.05.026.
- Li XR, Xing J, Li BY, Wang P, Liu JX. 2012. Use of tuf as a target for sequence-based identification of Gram-positive cocci of the genus *Entero*coccus, Streptococcus, coagulase-negative Staphylococcus, and Lactococcus. Ann Clin Microbiol Antimicrob 11:31. http://dx.doi.org/10.1186/1476 -0711-11-31.
- 6. Li RQ, Zhu HM, Ruan J, Qian WB, Fang XD, Shi ZB, Li YR, Li ST, Shan G, Kristiansen K, Li SG, Yang HM, Wang J, Wang J. 2010. De novo

- assembly of human genomes with massively parallel short read sequencing. Genome Res 20:265–272. http://dx.doi.org/10.1101/gr.097261.109.
- Besemer J, Lomsadze A, Borodovsky M. 2001. GeneMarkS: a self-training method for prediction of gene starts in microbial genomes. Implications for finding sequence motifs in regulatory regions. Nucleic Acids Res 29:2607–2618. http://dx.doi.org/10.1093/nar/29.12.2607.
- Lowe TM, Eddy SR. 1997. tRNAscan-SE: a program for improved detection of transfer RNA genes in genomic sequence. Nucleic Acids Res 25: 955–964. http://dx.doi.org/10.1093/nar/25.5.0955.
- Lagesen K, Hallin P, Rødland EA, Staerfeldt H-H, Rognes T, Ussery DW. 2007. RNAmmer: consistent and rapid annotation of ribosomal RNA genes. Nucleic Acids Res 35:3100–3108. http://dx.doi.org/10.1093/ nar/gkm160.
- Gardner PP, Daub J, Tate JG, Nawrocki EP, Kolbe DL, Lindgreen S, Wilkinson AC, Finn RD, Griffiths-Jones S, Eddy SR, Bateman A. 2009. Rfam: updates to the RNA families database. Nucleic Acids Res 37: D136–D140. http://dx.doi.org/10.1093/nar/gkn766.
- 11. Hsiao W, Wan I, Jones SJ, Brinkman FS. 2003. IslandPath: aiding detection of genomic islands in prokaryotes. Bioinformatics 19:418–420. http://dx.doi.org/10.1093/bioinformatics/btg004.
- 12. Zhou Y, Liang Y, Lynch KH, Dennis JJ, Wishart DS. 2011. PHAST: a fast phage search tool. Nucleic Acids Res 39:W347–W352. http://dx.doi.org/10.1093/nar/gkr485.