

# Draft Genome Sequence of *Psychrobacter piscatorii* Strain LQ58, a Psychrotolerant Bacterium Isolated from a Deep-Sea Hydrothermal Vent

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**Here, we report the 3.1-Mb draft genome sequence of *Psychrobacter piscatorii* strain LQ58, isolated from a deep-sea hydrothermal vent on the East Pacific Rise. The sequence will provide further insight into the environmental adaptation of psychrotolerant bacteria and the development of novel cold-active enzymes for industrial application.**

Received 11 January 2016 Accepted 14 January 2016 Published 3 March 2016

**Citation** Zhou M, Dong B, Liu Q. 2016. Draft genome sequence of *Psychrobacter piscatorii* strain LQ58, a psychrotolerant bacterium isolated from a deep-sea hydrothermal vent. *Genome Announc* 4(2):e00044-16. doi:10.1128/genomeA.00044-16.

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Members of the genus *Psychrobacter* are psychrotolerant or psychrophilic bacteria, which are widely distributed in marine and terrestrial environments (1–3). They have been considered good sources of cold-active enzymes for industrial and environmental uses (4–6). *Psychrobacter piscatorii* strain LQ58 (=MCCC 1A10701) is an aerobic, psychrotolerant, chemo-organotrophic bacterium, isolated from a deep-sea hydrothermal vent sediment sample collected on the East Pacific Rise at a depth of 2,906 m (3.10 S, 102.55 W). Growth was observed at temperatures between 0°C and 35°C with an optimum at 22°C in the presence of 2% NaCl. Strain LQ58 showed highest 16S rDNA sequence similarity with *P. piscatorii* T-3-2 (99.3%) (7), followed by *Psychrobacter aquaticus* CMS56 (98.6%) (8). *P. piscatorii* was reported to exhibit an extraordinarily high catalase activity (9). To date, there is no reference *P. piscatorii* genome sequence available publicly. Here, we report the draft genome sequence of strain LQ58, the first released *P. piscatorii* genome sequence, which will provide further insight into the environmental adaptation of psychrotolerant bacteria and the development of novel cold-active enzymes for industrial application.

Extraction of genomic DNA from strain LQ58 was carried out using a bacterial DNA kit (Omega) according to the manufacturer's instructions. The genome of strain LQ58 was sequenced by whole-genome shotgun sequencing using the Illumina MiSeq system at Shanghai Majorbio Bio-pharm Technology Co., Ltd. (Shanghai, China). A total of 804,651,728 reads were generated, representing an approximately 260-fold coverage of the genome. These reads were assembled using SOAPdenovo version 2.0 (10) and resulted in 139 contigs with an  $N_{50}$  of 47,940 bp. The size of the longest contig was 149,861 bp. The draft genome of strain LQ58 was analyzed and annotated using the NCBI Prokaryotic Genome Annotation Pipeline (PGAP) ([http://www.ncbi.nlm.nih.gov/genome/annotation\\_prok](http://www.ncbi.nlm.nih.gov/genome/annotation_prok)). Genes of interest were manually evaluated.

The draft genome sequence of strain LQ58 was 3,089,314 bp in length, with a G+C content of 44.1%. The genome contains 2,676 predicted open reading frames and 2,562 predicted protein-

coding sequences). There were 33 tRNA genes, 3 rRNA genes, and 1 noncoding RNA (ncRNA) gene predicted from this assembly.

*In silico* DNA-DNA hybridization (DDH) and average nucleotide identity values of LQ58 against *P. aquaticus* CMS56 were  $22.6 \pm 2.36\%$  and 79.3% (11), respectively, which was significantly lower than the recommended cutoff value for species delineation (12, 13).

Sequence analysis revealed that the LQ58 genome encodes 37 lipolytic enzymes, 58 peptidases, 2 formate dehydrogenases, 12 alcohol dehydrogenases, and 7 glycosyltransferases for potential biotechnology applications. LQ58 genome also encodes 3 cold-shock proteins (CspA, CspC and CspD), 1 capsular polysaccharide synthesis gene cluster, and 1 GroES-GroEL operon associated with cold stress (14). In addition, compared with other sequenced *Psychrobacter* genomes, LQ58 contains more catalase genes, including 1 HP1I-like monofunctional catalase gene and 1 HP1-like bifunctional catalase gene (15), which may be favorable for its adaptation to an oxidative environment.

**Nucleotide sequence accession numbers.** This whole-genome shotgun project has been deposited at DDBJ/EMBL/GenBank under the accession number **LNDJ00000000**. The version described in this paper is the first version, LNDJ01000000.

## ACKNOWLEDGMENTS

This work was supported by grants from the National Basic Research Program of China (973 Program, no. 2012CB722209), the Natural Science Foundation of Fujian Province (no. 2015J01136). The funders had no role in study design, data collection and interpretation, or the decision to submit the work for publication.

## FUNDING INFORMATION

This work, including the efforts of Meixian Zhou, was funded by National Basic Research Program of China (2012CB722209). This work, including the efforts of Meixian Zhou, was funded by Natural Science Foundation of Fujian Province (Fujian Provincial Natural Science Foundation) (2015J01136).

## REFERENCES

- Matsuyama H, Minami H, Sakaki T, Kasahara H, Watanabe A, Onoda T, Hirota K, Yumoto I. 2015. *Psychrobacter oceani* sp. nov., isolated from

- marine sediment. *Int J Syst Evol Microbiol* 65:1450–1455. <http://dx.doi.org/10.1099/ijs.0.000118>.
2. Romanenko LA, Tanaka N, Frolova GM, Mikhailov VV. 2009. *Psychrobacter fulvigenes* sp. nov., isolated from a marine crustacean from the sea of Japan. *Int J Syst Evol Microbiol* 59:1480–1486. <http://dx.doi.org/10.1099/ijs.0.007195-0>.
  3. Bakermans C, Ayala-del-Río HL, Ponder MA, Vishnivetskaya T, Gilichinsky D, Thomashow MF, Tiedje JM. 2006. *Psychrobacter cryohalolentis* sp. nov. and *Psychrobacter arcticus* sp. nov., isolated from Siberian permafrost. *Int J Syst Evol Microbiol* 56:1285–1291. <http://dx.doi.org/10.1099/ijs.0.64043-0>.
  4. Kim HW, Wi AR, Jeon BW, Lee JH, Shin SC, Park H, Jeon SJ. 2015. Cold adaptation of a psychrophilic chaperonin from *Psychrobacter* sp. and its application for heterologous protein expression. *Biotechnol Lett* 37:1887–1893. <http://dx.doi.org/10.1007/s10529-015-1860-y>.
  5. Petrovskaya LE, Novototskaya-Vlasova KA, Kryukova EA, Rivkina EM, Dolgikh DA, Kirpichnikov MP. 2015. Cell surface display of cold-active esterase EstPc with the use of a new autotransporter from *Psychrobacter cryohalolentis* K5(T). *Extremophiles* 19:161–170. <http://dx.doi.org/10.1007/s00792-014-0695-0>.
  6. Bujacz A, Rutkiewicz-Krotewicz M, Nowakowska-Sapota K, Turkiewicz M. 2015. Crystal structure and enzymatic properties of a broad substrate-specificity psychrophilic aminotransferase from the Antarctic soil bacterium *Psychrobacter* sp. B6. *Acta Crystallogr D Biol Crystallogr* 71:632–645. <http://dx.doi.org/10.1107/S1399004714028016>.
  7. Yumoto I, Hirota K, Kimoto H, Nodasaka Y, Matsuyama H, Yoshimune K. 2010. *Psychrobacter piscatorii* sp. nov., a psychrotolerant bacterium exhibiting high catalase activity isolated from an oxidative environment. *Int J Syst Evol Microbiol* 60:205–208. <http://dx.doi.org/10.1099/ijs.0.010959-0>.
  8. Shivaji S, Reddy GS, Suresh K, Gupta P, Chintalapati S, Schumann P, Stackebrandt E, Matsumoto GI. 2005. *Psychrobacter vallis* sp. nov. and *Psychrobacter aquaticus* sp. nov., from Antarctica. *Int J Syst Evol Microbiol* 55:757–762. <http://dx.doi.org/10.1099/ijs.0.03030-0>.
  9. Kimoto H, Yoshimune K, Matsuyma H, Yumoto I. 2012. Characterization of catalase from psychrotolerant *Psychrobacter piscatorii* T-3 exhibiting high catalase activity. *Int J Mol Sci* 13:1733–1746. <http://dx.doi.org/10.3390/ijms13021733>.
  10. Luo R, Liu B, Xie Y, Li Z, Huang W, Yuan J, He G, Chen Y, Pan Q, Liu Y, Tang J, Wu G, Zhang H, Shi Y, Liu Y, Yu C, Wang B, Lu Y, Han C, Cheung DW. 2012. SOAPdenovo2: an empirically improved memory-efficient short-read *de novo* assembler. *GigaScience* 1:18. <http://dx.doi.org/10.1186/2047-217X-1-18>.
  11. Reddy GS, Ara S, Singh A, Kumar Pinnaka A, Shivaji S. 2013. Draft genome sequence of *Psychrobacter aquaticus* strain CMS 56<sup>T</sup>, isolated from a cyanobacterial mat sample collected from water bodies in the McMurdo dry valley region of Antarctica. *Genome Announc* 1(6):e00918-13. <http://dx.doi.org/10.1128/genomeA.00918-13>.
  12. Meier-Kolthoff JP, Auch AF, Klenk HP, Göker M. 2013. Genome sequence-based species delimitation with confidence intervals and improved distance functions. *BMC Bioinformatics* 14:60. <http://dx.doi.org/10.1186/1471-2105-14-60>.
  13. Richter M, Rosselló-Móra R. 2009. Shifting the genomic gold standard for the prokaryotic species definition. *Proc Natl Acad Sci USA* 106:19126–19131. <http://dx.doi.org/10.1073/pnas.0906412106>.
  14. Mykytczuk NC, Trevors JT, Foote SJ, Leduc LG, Ferroni GD, Twine SM. 2011. Proteomic insights into cold adaptation of psychrotrophic and mesophilic *Acidithiobacillus ferrooxidans* strains. *Antonie Van Leeuwenhoek* 100:259–277. <http://dx.doi.org/10.1007/s10482-011-9584-z>.
  15. Chelikani P, Fita I, Loewen PC. 2004. Diversity of structures and properties among catalases. *Cell Mol Life Sci* 61:192–208. <http://dx.doi.org/10.1007/s00018-003-3206-5>.