

# Draft Genome Sequence of *Paecilomyces hepiali*, Isolated from *Cordyceps sinensis*

Yi Yu,<sup>a</sup> Wenting Wang,<sup>a</sup> Linping Wang,<sup>a</sup> Fang Pang,<sup>a</sup> Lanping Guo,<sup>c</sup> Lai Song,<sup>b</sup> Guiming Liu,<sup>b</sup> Chengqiang Feng<sup>a</sup>

Beijing Key Laboratory of Protection and Utilization of Chinese Medicine, Beijing Normal University, Beijing, China<sup>a</sup>; CAS Key Laboratory of Genome Sciences and Information, Beijing Institute of Genomics, Chinese Academy of Sciences, Beijing, China<sup>b</sup>; Resource Center of Chinese Materia Medica, China Academy of Chinese Medical Sciences, Beijing, China<sup>c</sup>

***Paecilomyces hepiali* is an endoparasitic fungus that commonly exists in the natural *Cordyceps sinensis*. Here, we report the draft genome sequence of *P. hepiali*, which will facilitate the exploitation of medicinal compounds produced by the fungus.**

Received 16 May 2016 Accepted 18 May 2016 Published 7 July 2016

Citation Yu Y, Wang W, Wang L, Pang F, Guo L, Song L, Liu G, Feng C. 2016. Draft genome sequence of *Paecilomyces hepiali*, isolated from *Cordyceps sinensis*. *Genome Announc* 4(4):e00606-16. doi:10.1128/genomeA.00606-16.

Copyright © 2016 Yu et al. This is an open-access article distributed under the terms of the [Creative Commons Attribution 4.0 International license](https://creativecommons.org/licenses/by/4.0/).

Address correspondence to Guiming Liu, liugm@big.ac.cn, or Chengqiang Feng, fengchengqiang@bnu.edu.cn.

*Paecilomyces hepiali* is an endoparasitic fungus that commonly exists in the natural *Cordyceps sinensis* anamorph stage. *C. sinensis* has been described as an exotic medicinal mushroom in traditional Chinese and Tibetan medicine (1). Data from the pure compounds are the most revealing in determining the effects of the fungus (2). The chemical constituents isolated from *C. sinensis* include cordycepin (3'-deoxyadenosine) and its derivatives (3). Many studies have demonstrated that cordycepin has significant pharmacological effects, including antiviral, insecticidal, antibacterial, immunostimulatory, and antitumor effects (2, 4, 5). The chemical profiles for *P. hepiali* mycelia are more similar to those for *C. sinensis* (3). *P. hepiali* contains pharmacologically active components similar to those of the natural *C. sinensis* (6).

*P. hepiali* was isolated from *Cordyceps sinensis* from Yushu, Qinghai Province, China. The genome of *P. hepiali* was sequenced using the Illumina HiSeq 2500 sequencing platform with 5 insert libraries (180 bp, 500 bp, 800 bp, 2 kbp, and 5 kbp), providing about 380-fold coverage of the genome. *De novo* assembly of the reads was performed using the SOAPdenovo2 (7). The gaps inside scaffolds were closed with GapFiller (8), resulting in 88 scaffolds with an  $N_{50}$  of 2,350 kbp and 660 contigs with an  $N_{50}$  of 178 kbp. Finally, a 34.6-Mb draft genome sequence with a 53.4% G+C content was obtained for *P. hepiali*.

Gene structure was predicted using an integrated approach. *De novo*, homologue-based, and RNA sequencing (RNA-seq)-based prediction models were conducted with Augustus, GeneWise, and TopHat, respectively. A total of 11,261 gene models were predicted, with an average length of 16,366,717 bp. tRNAs and rRNAs were predicted using tRNAscan-SE (9) and RNAmmer (10), respectively. The functions of genes were also annotated using COG, KEGG (11), and InterProScan (12). The genome contains 113 tRNAs and 23 rRNAs, representing 0.03% of the genome. A total of 6,083 genes were assigned to Clusters of Orthologous Groups (COGs), and 1,829 coding sequences (CDSs) were annotated into pathways using KAAS (13). The data offer a better understanding of *P. hepiali* biology and will facilitate the exploitation of medicinal compounds produced by the fungus.

**Nucleotide sequence accession numbers.** This whole-genome shotgun project has been deposited at DDBJ/EMBL/GenBank under the accession no. [LNDK00000000](https://www.ncbi.nlm.nih.gov/nuclink/LNDK00000000). The version described in this paper is version LNDK02000000.

## ACKNOWLEDGMENTS

This work was supported by the Co-Building Special Project of the Beijing Municipal Education Commission (grant no. SYS100270430), the National Natural Science Foundation of China (grants 41476166 and 81130070), and the Rare traditional Chinese medicine resources sustainable utilization of capacity building (grant 2060302).

## FUNDING INFORMATION

This work, including the efforts of Chengqiang Feng, was funded by Co-building Special Project of Beijing Municipal Education Commission (SYS100270430) and the Rare Traditional Chinese medicine resources sustainable utilization of capacity building (2060302). This work, including the efforts of Guiming Liu, was funded by the National Natural Science Foundation of China (41476166 and 81130070).

## REFERENCES

- Panda AK, Swain KC. 2011. Traditional uses and medicinal potential of *Cordyceps sinensis* of Sikkim. *J Ayurveda Integr Med* 2:9–13. doi:10.4103/0975-9476.78183.
- Yoshikawa N, Kunitomo M, Kagota S, Shinozuka K, Nakamura K. 2009. Inhibitory effect of cordycepin on hematogenic metastasis of B16-F1 mouse melanoma cells accelerated by adenosine-5'-diphosphate. *Anticancer Res* 29:3857–3860.
- Ng TB, Wang HX. 2005. Pharmacological actions of cordyceps, a prized folk medicine. *J Pharm Pharmacol* 57:1509–1519. <http://dx.doi.org/10.1211/jpp.57.12.0001>.
- Yoshikawa N, Nakamura K, Yamaguchi Y, Kagota S, Shinozuka K, Kunitomo M. 2004. Antitumor activity of cordycepin in mice. *Clin Exp Pharmacol Physiol* 31(Suppl 2):S51–S53. <http://dx.doi.org/10.1111/j.1440-1681.2004.04108.x>.
- Choi S, Lim MH, Kim KM, Jeon BH, Song WO, Kim TW. 2011. Cordycepin-induced apoptosis and autophagy in breast cancer cells are independent of the estrogen receptor. *Toxicol Appl Pharmacol* 257:165–173. <http://dx.doi.org/10.1016/j.taap.2011.08.030>.
- Zhu JS, Halpern GM, Jones K. 1998. The scientific rediscovery of an ancient Chinese herbal medicine: *Cordyceps sinensis*: part I. *J Altern Complement Med* 4:289–303. <http://dx.doi.org/10.1089/acm.1998.4.3.289>.

7. 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, Yiu SM, Peng S, Xiaoqian Z, Liu G, Liao X, Li Y, Yang H, Wang J, Lam TW, Wang J. 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>.
8. Boetzer M, Pirovano W. 2012. Toward almost closed genomes with Gap-Filler. *Genome Biol* 13:R56. <http://dx.doi.org/10.1186/gb-2012-13-6-r56>.
9. 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>.
10. 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>.
11. Kanehisa M, Goto S. 2000. KEGG: Kyoto Encyclopedia of Genes and Genomes. *Nucleic Acids Res* 28:27–30. <http://dx.doi.org/10.1093/nar/28.1.27>.
12. Jones P, Binns D, Chang HY, Fraser M, Li W, McAnulla C, McWilliam H, Maslen J, Mitchell A, Nuka G, Pesseat S, Quinn AF, Sangrador-Vegas A, Scheremetjew M, Yong SY, Lopez R, Hunter S. 2014. InterProScan 5: genome-scale protein function classification. *Bioinformatics* 30:1236–1240. <http://dx.doi.org/10.1093/bioinformatics/btu031>.
13. Moriya Y, Itoh M, Okuda S, Yoshizawa AC, Kanehisa M. 2007. KAAS: an automatic genome annotation and pathway reconstruction server. *Nucleic Acids Res* 35:W182–W185. <http://dx.doi.org/10.1093/nar/gkm321>.