Draft Genome Sequence of Dematiaceous Coelomycete Pyrenochaeta sp. Strain UM 256, Isolated from Skin Scraping

Su Mei Yew,^a Chai Ling Chan,^a Tuck Soon Soo-Hoo,^a Shiang Ling Na,^a Seong Siang Ong,^a Hamimah Hassan,^a Yun Fong Ngeow,^a Chee Choong Hoh,^b Kok Wei Lee,^b Wai Yan Yee,^b Kee Peng Ng^a

Tropical Infectious Diseases Research and Education Centre (TIDREC), Department of Medical Microbiology, University Malaya Medical Centre, Faculty of Medicine, University of Malaya, Kuala Lumpur, Malaysia^a; Codon Genomics SB, Jalan Bandar Lapan Belas, Pusat Bandar Puchong, Selangor Darul Ehsan, Malaysia^b

Pyrenochaeta, classified under the order *Pleosporales*, is known to cause diseases in plants and humans. Here, we report a draft genome sequence of a *Pyrenochaeta* sp. isolated from a skin scraping, with an estimated genome size of 39.4 Mb. Genes associated with the synthesis of proteases, toxins, plant cell wall degradation, and multidrug resistance were found.

Received 3 March 2013 Accepted 19 April 2013 Published 30 May 2013

Citation Yew SM, Chan CL, Soo-Hoo TS, Na SL, Ong SS, Hassan H, Ngeow YF, Hoh CC, Lee KW, Yee WY, Ng KP. 2013. Draft genome sequence of dematiaceous coelomycete *Pyrenochaeta* sp. strain UM 256, isolated from skin scraping. Genome Announc. 1(3):e00158-13. doi:10.1128/genomeA.00158-13.

Copyright © 2013 Yew et al. This is an open-access article distributed under the terms of the Creative Commons Attribution 3.0 Unported license.

Address correspondence to Kee Peng Ng, kpng@ummc.edu.my.

F ungi belonging to the *Pyrenochaeta* species are dematiaceous coelomycetes that have morphological characteristics similar to those in the *Phoma* species. These fungi have been reported to be important plant pathogens causing root disease in numerous plants, leading to significant plant loss (1, 2). In humans, the diseases caused by *Pyrenochaeta* are mostly onychomycosis and keratitis (3–5). A draft genome sequence of *Pyrenochaeta* sp. strain UM 256 was generated and is reported here to reveal the genes present in *Pyrenochaeta* sp., which are of interest to researchers from different fields. This fungus was isolated from a skin scraping and was identified molecularly using the universal primers ITS1 and ITS4.

Genomic DNA from *Pyrenochaeta* sp. UM 256 was extracted for the generation of a single-end library using the Roche 454 GS FLX+ system and a 3-kb insert size paired-end library using Roche 454 GS Junior. A total of 1,225,229 single-end reads and 199,554 paired-end reads were generated with 27× sequencing depth. The sequence reads were assembled using the GS *de novo* Assembler version 2.70 (Newbler, Roche), and 286 contigs (\geq 500 bp) were generated. The contigs were then scaffolded into 254 scaffolds (\geq 1,000 bp) with an N₅₀ size of 482 kb. The genome size was estimated to be 39.4 Mb with 50.35% G+C content. GeneMark-ES v2.3e was used to predict 12,545 protein-coding genes with an exon frequency of 2.75 exons per gene from repeatmasked scaffolds (6). The scaffolds were searched against the NCBI Swiss-Prot database, resulting in the annotation of 7,754 (61.81%) genes.

Putative genes of interest found in the UM 256 genome include those involved in the production of proteases, such as aspergillopeptidases and metalloproteinases, which are known to be virulence factors associated with tissue destruction (7); mycotoxins, such as the secondary metabolites gliotoxin, aflatoxin, and sterigamatocystin (8, 9); and enzymes involved in plant cuticle and cell wall degradation, such as cutinase, feruloyl esterase, and glucanase (10, 11). These putative proteins suggest that UM 256 might be a phytopathogenic fungus that is able to invade host tissues and cause immunosuppression. Also found were genes associated with multidrug resistance (*cdr1*, *cdr2*) conferring resistance to azole antifungal agents, and genes encoding proteins resistant to quinidine, flurocytosine, and benomyl/methotrexate, suggesting multidrug resistance (12–17); this is corroborated by the results of *in vitro* antifungal susceptibility tests performed in our laboratory.

Nucleotide sequence accession numbers. The nucleotide sequence of the *Pyrenochaeta* sp. UM 256 genome has been deposited in DDBJ/EMBL/GenBank under the accession no. AOUM00000000. The version described in this paper is the first version, accession no. AOUM01000000.

ACKNOWLEDGMENTS

This study was supported by research grant UM.C/625/1/HIR/MOHE/ MED/31 and Postgraduate Research Grants (PPP) PV051.2012 from the University of Malaya.

S.M.Y., K.P.N., Y.F.N., and H.H. conceived the project and contributed to the writing and editing of the manuscript. K.P.N., C.L.C., S.L.N., and S.S.O. were responsible for the isolation, identification, and DNA extraction. S.M.Y., C.C.H., K.W.L., and W.Y.Y. performed the genome sequencing and bioinformatics analysis.

REFERENCES

- Campbell RN, Schweers VH, Hall DH. 1982. Corky root of tomato in California caused by *Pyrenochaeta lycopersici* and control by soil fumigation. Plant Dis. 66:657–661.
- Lević J, Petrović T, Stanković S, Krnjaja V, Stanković G. 2012. Different symptoms in maize root caused by *Pyrenochaeta terrestris* and the fungal colony properties. Rom. Agric. Res. 29:339–347.
- English MP. 1980. Infection of the finger-nail by *Pyrenochaeta unguis-hominis*. Br. J. Dermatol. 103:91–94.
- Ferrer C, Pérez-Santonja JJ, Rodríguez AE, Colom MF, Gené J, Alio JL, Verkley GJ, Guarro J. 2009. New *Pyrenochaeta* species causing keratitis. J. Clin. Microbiol. 47:1596–1598.
- 5. Verkley GJ, Gené J, Guarro J, Pérez-Santonja JJ, Rodríguez AE, Colom MF, Alio JL, Ferrer C. 2010. Pyrenochaeta keratinophila sp. nov., isolated from an ocular infection in Spain. Rev. Iberoam. Micol. 27:22–24.
- Ter-Hovhannisyan V, Lomsadze A, Chernoff YO, Borodovsky M. 2008. Gene prediction in novel fungal genomes using an *ab initio* algorithm with unsupervised training. Genome Res. 18:1979–1990.

- Abad A, Fernández-Molina JV, Bikandi J, Ramírez A, Margareto J, Sendino J, Hernando FL, Pontón J, Garaizar J, Rementeria A. 2010. What makes *Aspergillus fumigatus* a successful pathogen? Genes and molecules involved in invasive aspergillosis. Rev. Iberoam. Micol. 27:155–182.
- Cramer RA, Gamcsik MP, Brooking RM, Najvar LK, Kirkpatrick WR, Patterson TF, Balibar CJ, Graybill JR, Perfect JR, Abraham SN, Steinbach WJ. 2006. Disruption of a nonribosomal peptide synthetase in *Aspergillus fumigatus* eliminates gliotoxin production. Eukaryot. Cell 5:972–980.
- 9. Keller NP, Kantz NJ, Adams TH. 1994. Aspergillus nidulans verA is required for production of the mycotoxin sterigmatocystin. Appl. Environ. Microbiol. 60:1444–1450.
- 10. De Vries RP, Vankuyk PA, Kester HC, Visser J. 2002. The *Aspergillus niger faeB* gene encodes a second feruloyl esterase involved in pectin and xylan degradation and is specifically induced in the presence of aromatic compounds. Biochem. J. **363**:377–386.
- Pauly M, Andersen LN, Kauppinen S, Kofod LV, York WS, Albersheim P, Darvill A. 1999. A xyloglucan-specific *endo-β-*1,4-glucanase from *Aspergillus aculeatus*: expression cloning in yeast, purification and characterization of the recombinant enzyme. Glycobiology 9:93–100.
- 12. Ben-yaacov R, Knoller S, Caldwell GA, Becker JM, Koltin Y. 1994.

Candida albicans gene encoding resistance to benomyl and methotrexate is a multidrug resistance gene. Antimicrob. Agents Chemother. **38**: 648–652.

- 13. Ghannoum MA, Rice LB. 1999. Antifungal agents: mode of action, mechanisms of resistance, and correlation of these mechanisms with bacterial resistance. Clin. Microbiol. Rev. 12:501–517.
- Krishnamurthy S, Gupta V, Prasad R, Panwar SL, Prasad R. 1998. Expression of *CDR1*, a multidrug resistance gene of *Candida albicans*: transcriptional activation by heat shock, drugs and human steroid hormones. FEMS Microbiol. Lett. 160:191–197.
- 15. Sá-Correia I, Tenreiro S. 2002. The multidrug resistance transporters of the major facilitator superfamily, 6 years after disclosure of *Saccharomyces cerevisiae* genome sequence. J. Biotechnol. **98**:215–226.
- Sanglard D, Ischer F, Monod M, Bille J. 1997. Cloning of *Candida albicans* genes conferring resistance to azole antifungal agents: characterization of *CDR2*, a new multidrug ABC transporter gene. Microbiology 143(Pt 2):405–416.
- Vlanti A, Diallinas G. 2008. The *Aspergillus nidulans* FcyB cytosinepurine scavenger is highly expressed during germination and in reproductive compartments and is downregulated by endocytosis. Mol. Microbiol. 68:959–977.