

MITOGENOME ANNOUNCEMENT

OPEN ACCESS  Check for updates

## The complete mitogenome of *Fusarium equiseti*

Xiaotang Sun, Mengshuang Shu, Pengmei Shang and Ruqiang Cui

College of Agronomy/Key Laboratory of Crop Physiology, Ecology and Genetic Breeding, Ministry of Education, Jiangxi Agricultural University, Nanchang, Jiangxi, China

### ABSTRACT

The complete mitochondrial genome of plant pathogenic fungus, *Fusarium equiseti*, was sequenced. The circular molecule is 53,411 bp long with a GC content of 32.81%. It contains 22 protein-coding genes, 4 ribosomal RNA (rRNA), and 24 transfer RNA (tRNA) genes. Phylogenetic reconstructions confirmed that it has the closest relationship with *Fusarium equiseti*. The mitogenome analysis of *Fusarium equiseti* provides a molecular basis for further studies on molecular systematics and evolutionary dynamics.

### ARTICLE HISTORY

Received 26 July 2019  
Accepted 3 August 2019

### KEYWORDS

*Fusarium equiseti*; mitogenome; phylogenetic analysis

The significantly important plant pathogens of species belonging to the genus *Fusarium* cause diseases in many crops and wild plants. The isolate of *Fusarium equiseti* is a strongly virulent strain, which can cause leaves wilting. We sequenced the complete mitogenome (mitochondrial genome) of the *Fusarium equiseti* strain 2018BL08 (GenBank accession number MN199625) isolated from *Nelumbo nucifera* in Guangchang City of Jiangxi Province (26°47'49" N, 116°18'11" E) and the specimens were stored at the Plant Pathology Lab in Jiangxi Agricultural University. DNA isolation using an improved extraction method (Chen et al. 2011) and libraries were sequenced on the Illumina Hiseq 4000 (Borgström et al. 2011) (Shanghai BIOZERON Co., Ltd, Shanghai, China) with a 150 bp paired-end read.

The filtered reads were assembled using ABySS (Simpson et al. 2009) and verifying the assembly and completing the circle. ORF Finder (<http://www.ncbi.nlm.nih.gov/gorf/gorf.html>) was used to identify any potential genes in the uncoding regions. The mitochondrial genes were annotated using homology alignments and de novo prediction. Transfer RNA (tRNA) genes and ribosome RNA (rRNA) genes were predicted by tRNAscan-SE (Lowe and Eddy 1997) and rRNAmer 1.2 (Lagesen et al. 2007).

The genome has a total length of 53,411 bp and the nucleotide composition of the mitogenome is: 34.05% of A, 33.13% of T, 17.81% of G, and 15.01% of C. It contains 22 protein-coding genes, 4 ribosomal RNA (rRNA), and 24 transfer RNA (tRNA) genes. The tRNA genes contain codons for all 20 standard amino acids. Most amino acids are represented by only one tRNA gene, however, two trnL (trnL-UAA and trnL-UAG), two trnG (trnG-ACC and trnG-UCC), two trnR

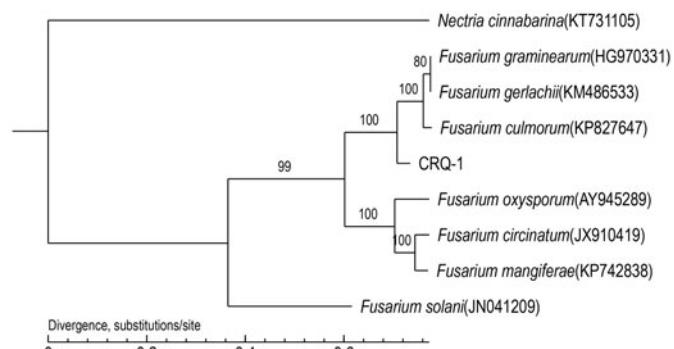


Figure 1. Phylogenetic analysis of 9 members of Nectriaceae based on core protein-coding genes.

(trnR-ACG and trnR-UCU), and two trnS (trnS-GCU and trnS-UGA) genes are found in this mitochondrial genome.

Nine members of Nectriaceae are included in the phylogenetic analysis, including eight taxa of *Fusarium*. Maximum likelihood (ML) were used to construct the phylogenetic trees with all protein-coding genes and rRNA by PhyML v3.0 (<http://www.atgc-montpellier.fr/phym/>) (Liu et al. 2019).

As shown in Figure 1, *Fusarium graminearum* (HG970331), *Fusarium gerlachii* (KM486533), and *Fusarium culmorum* (KP827647) are determined as sisters of *Fusarium equiseti* with strong support. High bootstrap and posterior probability values show that presented relations are stable. The mitochondrial genome of *Fusarium equiseti* will contribute to the understanding of phylogeny.

**CONTACT** Ruqiang Cui  [cuiruqiang@jxau.edu.cn](mailto:cuiruqiang@jxau.edu.cn) College of Agronomy/Key Laboratory of Crop Physiology, Ecology and Genetic Breeding, Ministry of Education, Jiangxi Agricultural University, No. 1101 Zhimin road, Nanchang City, Jiangxi Province 330045, China

© 2019 The Author(s). Published by Informa UK Limited, trading as Taylor & Francis Group.  
This is an Open Access article distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/4.0/>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

## Disclosure statement

No potential conflict of interest was reported by the authors.

## Funding

This work was supported by the Jiangxi Key Research and Development Program [No. 2016ACF60014].

## References

- Borgström E, Lundin S, Lundeberg J. 2011. Large scale library generation for high throughput sequencing. PLoS One. 6:e19119.
- Chen J, Guan R, Chang S, Du T, Zhang H, Xing H. 2011. Substoichiometrically different mitotypes coexist in mitochondrial genomes of *Brassica napus* L. PLoS One. 6:e17662.
- Lagesen K, Hallin P, Rodland EA, Staerfeldt HH, Rognes T, Ussery DW. 2007. RNAmmer: consistent and rapid annotation of ribosomal RNA genes. Nucleic Acids Res. 35:3100–3108.
- Liu J, Champer J, Langmüller AM, Liu C, Chung J, Reeves R, Luthra A, Lee YL, Vaughn AH, Clark AG, et al. 2019. Maximum likelihood estimation of fitness components in experimental evolution. Genetics. 211: 1005–1017.
- 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.
- Simpson JT, Wong K, Jackman SD, Schein JE, Jones SJ, Birol I. 2009. ABySS: a parallel assembler for short read sequence data. Genome Res. 19:1117–1123.