Draft Genome Sequence of *Cercospora arachidicola*, Causal Agent of Early Leaf Spot in Peanuts

Valerie A. Orner  
*National Peanut Research Laboratory, valerie.orner@ars.usda.gov*

Emily G. Cantonwine  
*Valdosta State University*

Xinye Monica Wang  
*National Peanut Research Laboratory*

Amr Abouelleil  
*Broad Institute of MIT and Harvard*

James Bochicchio  
*Broad Institute of MIT and Harvard*

See next page for additional authors

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Early leaf spot caused by *Cercospora arachidicola* S. Hori (teleomorph *Mycosphaerella arachidis* Deighton) is one of two important leaf spot diseases in peanut (*Arachis hypogaea* L.) responsible for significant economic loss to the industry (1, 2). Infections by *C. arachidicola* appear as small necrotic lesions on the leaves, petioles, or stems, which may be followed by premature defoliation, and, if left unmanaged on susceptible cultivars, can severely decrease yield (1). An effective, yet expensive, disease management strategy consists of multiple fungicide applications throughout the growing season (3). Other strategies such as strip-tillage instead of conventional tillage (4) or weather forecast models that predict disease outbreaks (5) can help minimize the number of fungicide treatments. However, the development of leaf-spot-resistant cultivars that require no fungicide application would be the most desirable means of control (6). The recent completion of the peanut genome ([http://www.peanutbase.org](http://www.peanutbase.org)) will aid breeding programs, but the negligible amount of *C. arachidicola* genetic information hinders progress. Currently, *C. arachidicola* entries in the NCBI-GenBank database total 8,077 bp in 21 sequences, with half of these entries corresponding to rRNA and the rest only 61 bp each. The genome sequence of *C. arachidicola* will provide relevant information for the advancement of leaf-spot resistant cultivars, be a useful resource to aid in the selection of target genes for disease control, and contribute to the study of genetic diversity of *C. arachidicola*.

A single-spore isolate of *C. arachidicola* from an infected peanut plant near Tifton, Georgia, USA, was grown on potato dextrose agar (Difco, Franklin Lakes, NJ, USA) for 6 months. The fungus was removed from the agar and ground using a Kleco tissue pulvzerizer (Garcia Machine, Visalia, CA, USA). Genomic DNA was extracted using phenol/chloroform/isoamyl alcohol followed by isopropanol precipitation (7) and cleaned using the GeneJET gel Extraction kit (Thermo, Fisher Scientific, Waltham, MA, USA). Sequencing and assembly were performed by the Broad Institute of MIT and Harvard using an Illumina HiSeq 2500 whole-genome shotgun approach. A total of 465,511,514 reads with an estimated genome coverage of $>100\times$ were assembled de novo using ALLPATHS (8) and generated 796 contigs ($>400$ bp each) with an average size of 40,930 bp that assembled into 491 scaffolds ($>1,000$ bp each). The average size of the scaffolds was 67,710 bp with a total of 33,245,410 bp and a maximum length of 1,387,526 bp. Most hits from BLAST analysis corresponded to the genera *Pseudocercospora* and *Passalora*.

**Nucleotide sequence accession numbers.** This whole-genome shotgun project has been deposited in DDBJ/ENA/GenBank under the accession number LIHB00000000. The version described in this paper is the first version, LIHB01000000.

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