Draft Genome Resource for the Potato Powdery Scab Pathogen *Spongospora subterranea*

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Abstract

The Plasmodiophorida (Phytomyxea, Rhizaria) are a group of protists that infect plants. Of this group, *Spongospora subterranea* causes major problems for the potato industry by causing powdery scab and root galling of potatoes and as vector for the *Potato mop-top virus* (PMTV) (genus *Pomovirus*, family *Virgaviridae*). A single tuber isolate (SSUBK13) of this uncultivable protist was used to generate DNA for Illumina sequencing. The data were assembled to a draft genome of 28.08 Mb consisting of 2,340 contigs and an L50 of 280. A total of 10,778 genes were predicted and 93% of the BUSCO genes were detected. The presented genome assembly is only the second genome of a plasmodiophorid. The data will accelerate functional genomics to study poorly understood interaction of plasmodiophorids and their hosts.

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Genome Announcement

Spongospora subterranea belongs to the Plasmodiophorida (Phytomyxea, Rhizaria) (Neuhauser et al. 2014; Sierra et al. 2016). This soilborne, obligate biotrophic protist is present in most potato-growing regions worldwide (Gau et al. 2013). It causes powdery scab of potato tubers, root galling in its host, reduces tuber yield, and transmits the *Potato mop-top virus* (PMTV) (genus *Pomovirus*, family *Virgaviridae*) (Balendres et al. 2016; Falloon et al. 2016). It can also diminish the growth of other solanaceous crops, such as tomatoes (*Solanum lycopersicum*) (Balendres et al. 2018). Because of their intracellular growth and obligate host-dependent life style, genomic data for plasmodiophorids are rare, although those pathogens cause substantial crop damage, e.g., in brassicas, sugar beet, or gramineous crops (Schwelm et al. 2018). So far, the only publicly available whole-genome data of plasmodiophorids are from the clubroot pathogen *Plasmodiophora brassicae* (Bi et al. 2016; Schwelm et al. 2015). Therefore, the here-presented *S. subterranea* data now enables comparative analyses between plasmodiophorid plant pathogens.

The genome data described here were generated from DNA of purified sporosori (characteristic aggregates of haploid resting spores) from potatoes originating from Lower Saxony, Germany (IB2016400). Sporosori from an individual potato tuber were collected to generate a single tuber isolate (SSUBK13). To reduce the amount of bacterial contamination, sporosori were immersed in lysis solution (GeneJET plasmid miniprep kit; Thermo-Scientific) for 5 min before washing twice by centrifugation in sterile water. Sporosori were purified by sequential filtering using CellTrics strainers (Sysmex). Material between 20 and 50 μ m was collected and a subsample of the sporosori was visually checked for the presence of fungal and bacterial contamination prior DNA extraction. Sporosori were snap-frozen in

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liquid nitrogen and disrupted using a Fastprep 24 5G homogenizer (MP Biomedicals, LLC). For lysis, the material was incubated in cetyltrimethylammonium bromide buffer at 65°C for 10 to 12 h before iso-amylalcohol and chloroform extraction and isopropanol precipitation of the DNA. A yellow-brown pigment coprecipitated with the DNA was separated from the DNA via gel electrophoresis. The genomic DNA of a size larger than 10 kb was purified from the gel using the ZYMO large-fragment DNA gel purification kit (Zymo Research Corporation). A strand-specific paired-end library (2× 150 bp) was constructed and was sequenced using HiSeq 2500 technique (Illumina) at the Vienna Biocenter Core Facilities sequencing platform. Reads were quality checked using FastQC and sequencing adapters, and low-quality reads (sliding window 5 bp; average quality score <20) were removed using Trimmomatic v0.36 (Bolger et al. 2014). Only surviving read pairs in which each read was longer than 70 bp were processed further. To filter out reads from contaminating organisms (e.g., bacteria) and the host, reads that matched to the potato genome (Potato Genome Sequencing Consortium et al. 2011) or the National Center for Biotechnology Information (NCBI) bacterial reference database using Bowtie2 v2.2.6 (Langmead and Salzberg 2012) were removed. The so-filtered reads were assembled using Velvet v1.2.10 (Zerbino and Birney 2008) and SPAdes v3.11.1 (Bankevich et al. 2012). Velvet assembled (kmer set to 31) contigs (≥1,000 nt, coverage ≥13) were used in the SPAdes assembly (kmer settings 23, 33, 41, 55, 63) as untrusted contigs. From this assembly only contigs longer than 500 bp were used and were manually sorted based on coverage, blastN hit, and length. Contigs with coverage above 30 were selected when their BlastN hit did not exclusively match bacterial genomes or the potato genome. All other contigs were analyzed by BlastX, using the NCBI nr database and *S. subterranea* protein models (Schwelm et al. 2015) and contigs with their best hits matching only to *P. brassicae* or *S. subterranea* protein models were also kept. Gapfiller v1.10 (Boetzer and Pirovano 2012) and SSPACE-standard v3.0 (Boetzer et al. 2011) were used to improve scaffolding.

The *S. subterranea* SSUBK13 genome draft presented here consists of 2,340 scaffolds with a total length of 28.08 Mb (229 N/100 kbp) and 1,150 scaffolds above 5 kb. The largest scaffold was 262,481 nt long (N50 value of 28,677; L50 value of 280). The GC content of 45.7% is lower than the 58% for *P. brassicae* (Schwelm et al. 2015) but comparable to the 46% GC content of the genome of the rhizarian *Bigelowiella natans* (Curtis et al. 2012). After repeat-masking (RepeatMasker program), a total of 10,778 genes were predicted using the BRAKER1 v2.0 pipeline (Hoff et al. 2016), a TopHat v2.1.0 (Kim et al. 2013)-created BAM-file, and available RNAseq reads (Schwelm et al. 2015; European Nucleotide Archive [ENA] BioProject PRJEB9159). The completeness of the genome assembly was estimated using BUSCO v2.0.1 (Simão et al. 2015), identifying 93% (88% complete, 5% partial) of the Eukaryota set. For *P. brassicae*, the only other sequenced Phytomyxea to date, 97% (94% complete, 3% partial) of BUSCO proteins are present in the isolate e3 (Schwelm et al. 2015). The presented genome draft contains the complete ribosomal DNA sequence (scaffold 678). The mitochondrial sequence was partially assembled from the unfiltered reads using MITObim (Hahn et al. 2013). Of the 10,780 predicted genes 1,991 could be categorized by KAAS (Moriya et al. 2007) to a KEGG category and 4,402 by eggNOG (Huerta-Cepas et al. 2016) to a COG (clusters of orthologous groups) category other than S, which is unknown function. SignalP v4.1 (Petersen et al. 2011) predicted 697 secreted proteins, of which 102 were classified as potential effectors by the fungal effector predictor EffectorP 2.0 (Sperschneider et al. 2018). Using dbCAN (Yin et al. 2012), 300 carbohydrate active enzymes (CAZymes) were predicted (e value < e⁻⁰⁵).

Table 1. Metrics of the *Spongospora subterranea* SSUBK13 genome draft

Feature	Numbers
Assembled genome size	28,084,886 nt
Number of scaffolds (>1 kb)	2,340 (2,255)
N50	28,677
L50	280
Largest scaffold	262,481 nt
Scaffolds >25 kb	339
BUSCOs	93% (271 complete, 15 partial)
Number of predicted genes	10,780
Number of predicted CAZymes	300
Number of predicted secreted proteins	697

The whole-genome sequence assembly (accessions OUQQ01000001 to OUQQ01002340) and the complete data set used in this report, including the Illumina raw reads have been deposited at the European Nucleotide Archive (ENA) under BioProject PRJEB26377 (Table 1).

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Author-Recommended Internet Resource

RepeatMasker program: <http://www.repeatmasker.org>