Genome Sequencing of *Pleurozium schreberi*: The Assembled and Annotated Draft Genome of a Pleurocarpous Feather Moss

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**ABSTRACT** The pleurocarpous feather moss *Pleurozium schreberi* is a ubiquitous moss species which plays a fundamental role in many terrestrial ecosystems, for instance within the boreal forest, the Earth’s largest terrestrial biome, this species plays a significant role in driving ecosystem nitrogen and carbon inputs and fluxes. By hosting dinitrogen (N2)-fixing cyanobacteria, the moss-cyanobacteria symbiosis constitutes the main nitrogen input into the ecosystem and by the high productivity and the low decomposability of the moss litter, *P. schreberi* contributes significantly to build-up soil organic matter, and therefore long-term C sequestration. Knowledge on *P. schreberi* genome will facilitate the development of ‘omics’ and system’s biology approaches to gain a more complete understanding of the physiology and ecological adaptation of the moss and the mechanisms underpinning the establishment of the symbiosis. Here we present the de novo assembly and annotation of *P. schreberi* genome that will help investigating these questions. The sequencing was performed using the HiSeq X platform with Illumina paired-end and mate-pair libraries prepared with CTAB extracted DNA. In total, the assembled genome was approximately 318 Mb, while repetitive elements account for 28.42% of the genome and 15,992 protein-coding genes were predicted from the genome, of which 84.23% have been functionally annotated. We anticipate that the genomic data generated will constitute a significant resource to study ecological and evolutionary genomics of *P. schreberi*, and will be valuable for evo-devo investigations as well as our understanding of the evolution of land plants by providing the genome of a pleurocarpous moss.

**KEYWORDS** *Pleurozium schreberi* Genome sequencing Comparative genomic Annotation Genome assembly

*Pleurozium schreberi* (Brid.) Mitt. also known as the red-stemmed feather moss belongs to the order Hypnales, family Hylocomiaceae. *P. schreberi* has a wide global distribution and is one of the most common moss species dominating the ground layer in the boreal forest, sub-alpine and Arctic ecosystems (Nilsson and Wardle 2005; Lindo and Gonzalez 2010). *P. schreberi* serves a variety of key functions in its ecosystem; for instance, *P. schreberi* can host dinitrogen (N2)-fixing cyanobacteria, a symbiosis which serve as the major input of nitrogen (N) into boreal forests and thus play a vital role in primary productivity of this N limited ecosystem (Turetsky 2003; Lindo et al. 2013). High-resolution secondary ion mass spectrometry verified transfer of fixed N from the cyanobacteria to the moss host, further demonstrating that the symbiosis governs the main N entrance to boreal forests (Bay et al. 2013). While, the genetic and genomic of *P. schreberi* symbiotic cyanobacteria have been previously studied (Ininbergs et al. 2011; Warshan et al. 2016, 2017 and 2018), nothing is known about the moss genomic diversity and gene repertoire needed to form the symbiosis. In particular the genome will be a crucial tool for further studies on the symbiotic interaction with N2-fixing cyanobacteria where previous study, have shown the importance of moss secreted signaling molecules to attract the cyanobacteria (Bay et al. 2013). Further, the biomass of *P. schreberi* and *Hylocomium splendens* (another ubiquitous feather moss in the boreal forest) can account for up to a third of the total...
As boreal forests represent one of the largest terrestrial carbon (C) sinks on Earth, storing approximately 30% of total terrestrial C stocks (Goodale et al. 2002; Pan et al. 2013) the feather mosses play a significant role in the global C cycles. As such, the response of the feather moss cover to future climate change may influence whether and how the ecosystem will be regulated and the consequences for C sequestration by this biome (Lindo et al. 2013). The availability of P. schreberi genomic resource will facilitate population genetic study of P. schreberi at genomic level using genotyping-by-sequencing approaches. For instance, investigating the link between intraspecific genetic diversity and moss traits variations, such as its physiology, decomposition rate and C accumulation could reveal pattern of local adaptations that might have an impact on N and C cycles. Even though mosses are the second most diverse phylum of land plant with approximately 13,000 species (Goffinet et al. 2009), to date only two other moss species had their genome sequenced i.e., Sphagnum fallax (DOE-JGI, http://phytozome.jgi.doe.gov/; Shaw et al. 2016), and Physcomitrella patens (Rensing et al. 2008). The latter of which have become a very important model organism (Cove et al. 2009).

In this study, we present a draft genome of an axenic pleurocarpous feather moss P. schreberi. Indeed, the sequenced draft genome of the moss will be a fundamental resource for future research spanning from evolutionary to ecological aspects.

**MATERIALS AND METHODS**

**Sample collection**

Pleuroziunm schreberi is a pleurocarpous moss with typical tissue growth such as leafy gametophores with phyllids, rhizoids and reproductive organs (Figure 1A, B), protonemal tissue (Figure 1C) as well as gametangia (Figure 1D). Intact P. schreberi sporophytes were randomly collected June 26, 2016 on Blidö (59°52.20.06’N, 18°90.26.41’E), an island located in the Stockholm archipelago, Sweden. An axenic line was generated by first separating the septa from the sporophyte with forceps and then placing in 99.9% ethanol for 1 min, subsequently moved to 1 min in 5% sodium hypochlorite followed by 4 rounds of rinsing in sterile water. The sterilized spores where placed on BCD media (Cove et al. 2009) + 5% sucrose for approximately 2 weeks. Axenic germinated spores were transferred to BCD and placed in an incubator at 24°C with constant white light at 30 W/m² to allow the gametophore to growth. Pictures of the P. schreberi tissues were taken using the Zeiss Stemi 2000-C Stereo microscope equipped with an AxioCam HR microscope camera (Zeiss, Oberkochen, Germany).

**Genome sequencing and assembly**

DNA extraction was performed from gametophores originating from a single spore isolate according to Schlink and Reski (2002) with the following modifications; 10% CTAB was used during the extraction and DNA was re-dissolved in water. After the DNA quality check using the Bioanalyzer 2100 system (Agilent Technologies, Santa Clara, USA), 180bp average fragment size Illumina TruSeq PCR-free library (Illumina, Cambridge, UK), 3kb and 5-8kb insert size Nextera mate-pair libraries (Illumina) were constructed at SciLifeLab (Stockholm, Sweden) following manufacturer’s recommendations. Sequencing proceeded on the Illumina HiSeq X platform, where the Illumina TruSeq PCR-free library and the Nextera mate-pair libraries were sequenced on separate lanes at 2x150bp. Preliminaries de novo assemblies were conducted with NouGAT, NGI open universal Genome Assembly Toolbox pipeline (https://github.com/vezzi/NouGAT) using Abyss (Simpson et al. 2009), Allpaths (Butler et al. 2008), and SOAPdenovo (Luo et al. 2012) assemblers with default settings. The quality of the selected assembly was assessed using BUSCO version 3 (Simão et al. 2015) with the eukaryota database (release 9) in protein mode. We estimated genome size using
The gene models were carried out using the MAKER pipeline, version 3.0.1.1 (Cantarel et al. 2008; Holt and Yandell 2011). The gene models incorporate ab initio gene prediction, homology-based prediction and RNA-seq assisted prediction. Prior to ab initio gene prediction, repeat regions of the moss were masked based on repeat annotation results. Homology-based prediction and RNA-seq assisted prediction used protein sequences collected on the Uniprot database (Magrane and Consortium 2011) selecting those belonging to the Swiss-prot section and nine P. schreberi protein sequences. The input data used included protein sequences and reference proteins. Finally, a second round of annotation using MAKER with ab initio tools Augustus, SNAP, GeneMark and EvidenceModeler (Haas et al. 2008) and evidence build (proteins and transcripts) allowed creating an ab initio evidence-driven gene build. After a visual analysis of the ab initio evidence-driven gene build, spurious ORFs among the gene structure were discovered in some loci, and fixed using in-house scripts (https://github.com/NBISweden/GAAS). k-mer counting of quality and barcode trimmed reads using Jellyfish v2.2.10 (Marçais and Kingsford 2011). K-mer frequency distributions of 21-, 25-, 31-, 41-, 65-mers were generated, then we used findGSE (Sun et al. 2017) to estimate genome size and repeat content.

Repeat content analysis

To identify repeats in the genome assembly, a custom species-specific repeat library was created, using the RepeatModeler package 1.0.8 (Smit 2010) with default parameters. As repeats can be part of actual protein-coding genes, the candidate repeats were vetted against the Uniprot/Swiss-prot protein set (minus transposons) to exclude any nucleotide motif stemming from low-complexity coding sequences. From the repeat library, identification of repeat sequences present among the genome was performed using RepeatMasker 4.0.3 (Smit et al. 2010) and RepeatRunner (Magrane and Consortium 2011).

Gene prediction

The gene models were carried out using the MAKER pipeline, version 3.0.1.1 (Cantarel et al. 2008; Holt and Yandell 2011). The gene models incorporate ab initio gene prediction, homology-based prediction and RNA-seq assisted prediction. Prior to ab initio gene prediction, repeat regions of the moss were masked based on repeat annotation results. Homology-based prediction and RNA-seq assisted prediction used protein sequences collected on the Uniprot database (Magrane and Consortium 2011) selecting those belonging to the Swiss-prot section and nine P. schreberi transcriptomes coming from the JGI study ID number Gs0110198. The transcriptome of P. schreberi (biosample ID Gb0144502- Gb0144504) had already been assembled by JGI using Rnnotator 3.4.0 (Martin et al. 2010) and completed with Velvet 1.2.07 (Zerbino and Birney 2008). The six other transcriptomes (ID Gb0144505- Gb0144510) have been assembled using HISAT2 (version 2.1.0) (Kim et al. 2015) and StringTie 1.2.2.2 (Pertea et al. 2015) with default parameters. MAKER ab initio training was performed using Augustus (Stanke et al. 2008), GeneMark (Besemer and Borodovsky 2003) and SNAP (Bromberg and Rost 2007) with transcripts and high-confidence protein sequences. For ab initio training purpose, a gene set was created by selecting the best gene models based on i) genes have to be complete (i.e. start/stop codons mandatory); ii) no similarity over 85 is allowed among genes of the set; iii) Annotation Edit Distance (AED) scores have to be inferior to 0.3; iv) genes have to be at a distance of 1,000 bp from each other. In total, 4,264 genes were selected and used for Augustus and SNAP training process. GeneMark has also been trained using splicing sites information of nine transcriptome assemblies. Then, an evidence-guided build was computed by allowing the MAKER software to construct gene models directly from both aligned transcript sequences and reference proteins. Finally, a second round of annotation using MAKER with ab initio tools Augustus, SNAP, GeneMark and EvidenceModeler (Haas et al. 2008) and evidence build (proteins and transcripts) allowed creating an ab initio evidence-driven gene build. After a visual analysis of the ab initio evidence-driven gene build, spurious ORFs among the gene structure were discovered in some loci, and fixed using in-house scripts (https://github.com/NBISweden/GAAS).

Table 1: Statistics of Pleurozium schreberi genome assembly

| Total length (bp) | 318,338,550 |
|-------------------|------------|
| Number of scaffolds | 2,695 |
| Number of uncalled bases (N's) | 98,291,915 |
| Number of N-regions | 35,733 |
| NG50 (bp) | 204,181 |
| GC-content (%) | 26.4 |

Table 2: Repeat statistics by RepeatMasker and RepeatRunner

| Number of repeats | RepeatMasker | RepeatRunner |
|-------------------|--------------|--------------|
| Total size (kb)   | 255,686      | 484          |
| Mean size (bp)    | 90,223       | 271.39       |
| Percentage of the genome (%) | 35.26       | 560.73       |

In the case of loci containing proteins and/or transcripts without ab initio prediction, the gene models were taken from the evidence build.

Gene annotation

The functional inference for genes and transcripts were performed using the translated CDS features of each coding transcript. Each predicted protein sequences were blasted against the Uniprot/Swissprot database to retrieve the gene name and the protein function as well as run against InterProscan 5.21-60 (Jones et al. 2014) to retrieve functional information from 21 different sources (Table S1).

Gene name inference was performed with the best blast hit approach using the Uniprot reference data set (Kim et al. 2015). Only the hits with an e-value inferior to 10E-7 were taken into account. TRNAs have been predicted by using tRNAscan v1.3.1 (Lowe and Eddy 1997) with default settings and other non-coding RNAs by using the RNA family database Rfam (Nawrocki et al. 2015) with conserved eukaryotic models only.

Analyses of potential whole genome duplication in P. schreberi

Whole genome duplication (WGD) events can be detected using synonymous substitution rates (Ks) among pairs of paralogous genes. WGD event(s) within P. schreberi were estimated using the FASTKs pipeline (https://github.com/mrmckain/FASTKs). Protein-coding genes were blasted against themselves using an e-value cutoff of 10E-5 and then putative pairs were filtered using FASTKs default parameters. Amino acid sequences for putative paralog pairs were then aligned using MUSCLE v3.8.31 (Edgar 2004), and back translated to CDS using PAL2NAL v14 (Suyama et al. 2006). Ks were estimated for the aligned pairs using codeml in PAML v4.8 (Yang 2007) using the same parameters used by McKain et al. (2012). All Ks values >0.1 were excluded for analysis to avoid the incorporation of allelic variants and to prevent the fitting of a component to infinity, while Ks values >5.0 were removed because of Ks saturation (Vanneste et al. 2015). Normal mixture models were estimated for Ks values using the mclust v 5.4.3 (Scruca et al. 2016) in R (v3.5.1). We evaluated mixture models with between one and nine components, and the best fit model was chosen using the Bayesian Information Criterion (BIC).

Comparative genomic

A comparative genomic analysis was performed on three different moss genomes (Physcomitrella patens, Sphagnum fallax and P. schreberi) using OrthoMCL 2.0.9 (Li et al. 2003) with mcl-14-137 (Enright et al. 2002) and BLAST+ 2.2.28 (Camacho et al. 2009) to infer homologous (both orthologous and paralogous) relationships among a set of protein sequences. The input data used included protein sequences from P. schreberi, P. patens (v1.39) and S. fallax (v0.5). For each moss genome, the CDS were filtered to keep only the longest coding sequence for each gene, and the corresponding sequences were extracted, translated and filtered to retain those larger than 50 amino acids (Table S2). An all-against-all BLASTP analysis of the sequences of these moss species led to the identification of candidate homologs. BLAST hits with an e-value >10E-5 and for which the query and the hit sequence had <50% overlap of their gene length, were excluded. Gene families...
characterized by the OrthoMCL analysis were processed to infer the presence/absence of the genes along a species tree to provide an evolutionary view of the gene flux in those genomes.

**Data availability**

The raw data are deposited in NCBI with SRA accessions numbers: SRR8297981 and SRR8297982 (BioProject accession number PRJNA509035). The BioSample is available with accession number SAMN10578977 at NCBI. The assembled genome is available with the accession number YACF00000000 at NCBI. All of the annotation files are available at github: https://github.com/Pycnophila/Pleurozium_schreberi_annotated_genome_files. Supplemental material available at Figshare: https://doi.org/10.6084/m9.figshare.7773951.

**RESULTS AND DISCUSSION**

**Genome assembly**

Concerning the initial genome assemblies of *Pleurozium schreberi*, the Allpaths assembler showed the best assembly statistics (i.e., highest NG30 value and lowest number of scaffolds) compared to assemblies done by Abyss and SOAPdenovo (Table S3). Quality control of the Allpaths assembly including coverage distribution, GC content, contig length and median coverage are shown in Figure S1. Low coverage contigs showed deviation in GC content, which might be a sign of contamination, and therefore they were removed from the assembly (Figure S1C). The total length of the assembly was 318.34 Mb in 2,695 scaffolds with an NG30 score of 204 Kb, and a genomic GC content of 26.4% (Table 1). The repeat regions were estimated to account for 28.42% of the genome (90 Mb; Table 2), and BUSCO analysis captured 90.1% of genes in the eukaryota database with 75.6% complete, 14.5% fragmented and 9.9% missing genes (Table 3). Even if the genome size and number of repetitive elements in *P. schreberi* appears smaller compared to *P. patens*, they are similar to the estimates of *S. fallax* assembly (genome size of 475.8 and 396 Mb, and percentage of repetitive elements of 57% and 32% for *P. patens* and *S. fallax*, respectively; Shaw et al. 2016; Lang et al. 2018). When comparing to the k-mer analysis, the predicted genome size and percentage of repetitive elements are reasonably close to our assembly (Table S4). Genome size estimates ranged from 296.9 to 319.4 Mb, and repetitive elements ranged from 12 to 26% of the genome (Table S4). Nevertheless, the length of our assembly is small compared to genome size predicted by flow cytometry that indicated a genome 1.3x – 2.5x larger (ranging from 418 – 809 Mb; Voglmayr 2000; Bainard 2011). A high proportion of repeat sequences in *P. schreberi* genome could result in an underestimation of genome size, but k-mer analysis provides an indication that this was not the case, since the percentage of predicted repeated elements was very similar to the one estimated by our repeat analysis (Table 2 and Table S4). Alternatively, since the BUSCO analysis indicates that our genome assembly was incomplete, the discrepancy observed with the flow cytometry estimates could be partially explained by the fraction of coding sequence missing from our data. Besides, considering that the genome size estimations by flow cytometry originate from gametophyte cells (Voglmayr 2000; Bainard 2011), a most likely explanation to the extreme variation observed in genome sizes seems to be endopolyploidy in gametophyte tissues of *P. schreberi*. In addition, different isolates could have different ploidy levels, particularly in widespread species like *P. schreberi*. This appears to be a common phenomenon in this species and in mosses in general (Bainard and Newman 2010). Throughout the assembly of a plant genome many difficulties are incurred (Griesmann et al. 2018) and it is therefore not surprising that this assembly is less complete at the first draft stage, though future work should be conducted to improve these assemblies.

**Genome annotation**

The final annotation of the *P. schreberi* genome from the MAKER annotation pipeline included 15,992 protein-coding genes (Table 4). The functional annotation using InterProScan to detect motifs, domains, signatures, and BLASTP on the Uniprot/Swissprot database resulted in putative function annotation of 13,470 proteins (84.23% of the CDS; Table S5). Further, the best blast hit approach using the Uniprot/Swissprot database assigned a name to 12,295 genes (76.88%; Table S5).

**Whole genome duplication in *P. schreberi***

Our Ks-based analysis of paralogs was able to detect a peak of Ks values between 0.5 and 2.0, which support one WGD in *P. schreberi* (Figure 2A). Ks analyses conducted on *P. schreberi* and 24 other Hypnales transcriptomes, were unable to detect a clear evidence of WGD event (Johnson et al. 2016), but our analysis showed a stronger signal of WGD. When Gaussian distributions were fitted to the Ks values the best fit was of 4 components, but no obvious second large peak of Ks values was observed (Figure 2A). This would have been an indication of a second WGD event in *P. schreberi*, as observed in *P. patens* and *S. fallax* where several rounds of WGD and/or multiple large-scale duplication events have occurred (Devos et al. 2016; Lang et al. 2018). When considering that the number of protein-coding genes in the genome of *P. schreberi* is 2x smaller than in *P. patens* and 1.7x smaller than in *S. fallax* (32,926 and 26,939 CDS for *P. patens* and *S. fallax*, respectively), the fact that only one WGD event occurred in *P. schreberi* could explain the difference observed.

**Comparative genome analysis**

The comparison of gene content of the *P. schreberi* genome with the two other moss genomes available (*P. patens* and *S. fallax*) was conducted to illustrate potential gene flux between the moss species. We found 6,753 gene families shared between the moss species (Figure 2B, Table S2), and 5,109 gene families unique to *P. schreberi* (361 in-paralogs and
The comparative analysis suggests a large evolutionary difference between the three species, while Figure 2B shows that *P. schreberi* is sharing nearly 60% of gene families with *P. patens* and/or *S. fallax*, and that the remaining 40% of gene families identified in *P. schreberi* are unique for the species. Besides, it appears that more gene families are shared between *S. fallax* and *P. patens* (1,736 shared gene families) than with *P. schreberi* (Figure 2B). The gene flux analysis over the evolutionary time line for *P. schreberi*, *S. fallax* and *P. patens* provides some indications to this observation by predicting that most of the gene gain occurred in the most recent common ancestor between the three moss species (8,652 gene families; Figure 2C), and that after the split with *P. patens*, *P. schreberi* gain 361 gene families and lose the 1,736 gene families that are in common between *P. patens* and *S. fallax* (Figure 2C). Nevertheless, this large putative gene loss in the genome might be due to the incompleteness of our draft assembly (Table 3). Moreover, due to the limited number of moss genomes available, this result remains highly speculative especially considering that in the moss phylogeny, *P. schreberi*, *P. patens* and *S. fallax* are evolutionarily quite distant from each other (Liu et al. 2019).

CONCLUSIONS

Illumina HiSeq X combining mate-pair and low error rate short-read sequencing enabled *de novo* assembly of the genome of a pleurocarpous moss species, *Pleurozium schreberi*.

Here, we provide a draft assembly that will be useful for long-read sequencing to improve genome scaffolding and completeness of *P. schreberi* genome. Further, a large part of the genes was functionally annotated (84.23%) and preliminaries gene ontology and comparative analyses were undertaken, which will ease future functional genomic studies. Considering that *P. schreberi* occupies a key role in the ecosystem functioning of cold biomes, the data generated provide the genomic resources to deepen our understanding of the physiology of this ecologically important moss species.

ACKNOWLEDGMENTS

We would like to thank colleagues at the Oak Ridge National Laboratory, Duke University, and the DOE Joint Genome Institute for access to prepublication genome and annotation of *Sphagnum fallax* to use in comparisons against gene families. The Nilsson-Ehle Endowments from The Royal Physiographic Society of Lund for financial support. The authors would also like to acknowledge support from Science for Life Laboratory (SciLifeLab, Sweden), the National Genomics Infrastructure (NGI, Stockholm, Sweden), Nation Bioinformatics Infrastructure (NBIS; Stockholm, Sweden) and Uppmax (Uppsala, Sweden), for providing assistance in massive parallel sequencing as well as the computational infrastructure.

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Communicating editor: S. Mathews