A draft genome of *Alliaria petiolata* (garlic mustard) as a model system for invasion genetics

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Abstract

The emerging field of invasion genetics examines the genetic causes and consequences of biological invasions, but few study systems are available that integrate deep ecological knowledge with genomic tools. Here we report on the *de novo* assembly and annotation of a genome for the biennial herb *Alliaria petiolata* (M. Bieb.) Cavara & Grande (Brassicaceae), which is widespread in Eurasia and invasive across much of temperate North America. Our goal was to sequence and annotate a genome to complement resources available from hundreds of published ecological studies, a global field survey, and hundreds of genetic lines maintained in Germany and Canada. We sequenced a genotype (EFCC-3-20) collected from the native range near Venice, Italy and sequenced paired-end and mate pair libraries at ~70× coverage. A *de novo* assembly resulted in a highly continuous draft genome ($N_{50} = 121\text{Mb}$; $L_{50} = 2$) with 99.7% of the 1.1Gb genome mapping to contigs of at least 50Kb in length. A total of 64,770 predicted genes in the annotated genome include 99% of plant BUSCO genes and 98% of transcriptome reads. Consistent with previous reports of (auto)hexaploidy in western Europe. Almost, we found that almost one third of BUSCO genes (390/1440) mapped to two or more scaffolds despite a genome-wide average of < 2% heterozygosity. The continuity and gene space quality of our draft genome assembly will enable genomic studies of *A. petiolata* to address questions relevant to invasion genetics and conservation efforts.

Introduction

Biological invasions are a threat to global biodiversity with significant impacts to human health and welfare (Mack et al., 2000). They also present opportunities for large-scale ‘natural’ experiments to study ecological and evolutionary processes in the wild (Mooney and Cleland, 2001; Sax et al., 2005). Despite a large body of ecological research and a growing number of evolutionary studies of invasive species, functional genetic and ‘omics approaches have been rare among studies of invasive species until recently (reviewed in (Barrett, 2015; Bock et al., 2015); but see e.g. (Barrett et al., 2016; Boheemen et al., 2017; Bourne et al., 2020)). Apart from studies using neutral markers to assess population structure (reviewed in (Dlugosch and Parker, 2008)), relatively little is known about the genetic causes and consequences of biotic invasions at the molecular level. A lack of genomic resources has hindered high-resolution genetic studies of most invasive species. Here, we report on a draft genome for the herbaceous biennial plant *Alliaria petiolata*, a plant invader in North America with potential to become a model system for the emerging field of invasion genetics.

Several factors favour *A. petiolata* as an emerging model system for invasion genetics (Colautti et al., 2014). First, it is widely distributed with variable ecological impacts throughout its range (Lankau et al., 2009; USDA, 2020). Second, it has a relatively simple two-year life cycle with well-defined life stages (seed, rosette, bolting, senescent). This simple life history facilitates measurements of lifetime fitness, natural selection, and their impacts on population dynamics in natural populations. Third, *A. petiolata* is a member of the Brassicaceae and therefore benefits from genetic resources available for well-studied species like *Brassica rapa* (canola) and the model plant *Arabidopsis thaliana*, providing opportunities for functional and comparative genomics. Fourth, high selfing rates produce naturally inbred seed families that can be maintained through single-seed descent. Fifth, *A. petiolata* has been the focal species in hundreds of field surveys and experimental studies, including influential studies testing the role of natural enemies (Lewis et al., 2006), competition (Prati and Bossdorf, 2004), the ‘novel weapons’ hypothesis (Callaway et al., 2008), competitive ability (Bossdorf et al., 2004), glucosinolate metabolism

(Haribal et al., 2001), and eco-evolutionary dynamics (Lankau et al., 2009). One mechanism that is especially well-studied is the production of secondary metabolites and their effects on soil microbiota, particularly the suppression of mycorrhizal fungi that form beneficial symbiotic networks among native plant roots (Anthony et al., 2017; Duchesneau et al., 2020). Understanding the genomic basis of such interactions with soil microbes will not only advance basic science but it also has potential applications in plant restoration and agriculture.

Recent efforts to develop *A. petiolata* as a model system include the Global Garlic Mustard Field Survey (GGMFS), which mobilized 164 participants from 16 countries across North America and Europe to collect field data and seed samples across Europe and North America, resulting in thousands of seed families from 383 distinct populations (Colautti et al., 2014). A subset of inbred lines collected across North America have been maintained through single-line descent in labs in Germany (Bossdorf Lab, University of Tübingen) and Canada (Colautti Lab, Queen's University). Adding to these resources, we here report on a draft genome of a single *A. petiolata* genotype from Europe, annotated with RNA sequencing of leaf and root tissue.

Methods & Materials

Study Organism and Line Derivation

Alliaria petiolata (M. Bieb.) Cavara & Grande is a biennial herbaceous plant in the Thlaspidaceae tribe of the Brassicaceae family. It was introduced to North America prior to 1868, when it was discovered in Long Island, New York (Nuzzo, 1993). By 1948 it was reported on the West Coast of North America and has established in at least 37 U.S. States and five Canadian provinces from the Atlantic coast to the Pacific (Cavers et al., 1979; USDA, 2020). As the only species of *Alliaria* with a broad distribution, *A. petiolata* is relatively easy to identify in natural habitats owing to its white flowers and dentated peltate leaves with long petioles. It is considered a noxious weed across most of its introduced range, due in part to impacts on native plants and tree regeneration in deciduous forest ecosystems (Cipollini and Cipollini, 2016; Stinson et al., 2007). Two genome types have been identified, including diploids ($2n = 14$) in Eastern Europe and Western Asia and hexaploids ($2n = 42$ and $2n = 36$) in Central Europe and North America (Esmailbegi et al., 2018; Weiss-Schneeweiss and Schneeweiss, 2003).

S_0 generation

All source material for genome and transcriptome sequencing originated from a single individual grown from seed collected as part of the GGMFS (Colautti et al., 2014). The inbred line used in this study is from population EFCC3, collected in 2011 from a small forest fragment (~2,000m²) surrounded by agricultural land, about 75km northwest of Venice, Italy (UTM 45.71°N × 11.72°E). The specific seed line was sampled at 3.2m along a 10m sampling transect originating at the edge of population EFCC3. This inbred line (code EFCC3-3-20) is currently maintained along with other GGMFS seed collections by two of the coauthors in replicate collections in Tübingen, Germany (Bossdorf) and Kingston, Ontario, Canada (Colautti).

S_1 generation

In July 2012, ten seeds of the EFCC3-3-20 genotype from the original S_0 field collection were removed from cold storage (4 °C), surface washed with a mild detergent and rinsed with distilled H₂O before surface sterilizing in 10% bleach for 10 minutes. Sterilized seeds were again rinsed with distilled H₂O before placing on filter paper saturated with distilled H₂O and sealed in a petri dish with paraffin wax.

We stratified seeds in the dark at 10 °C for ~90 days and thereafter inspected weekly until emerging radicles were observed. Germinating seeds were transplanted into 4" plastic pots containing a peat soil mixed with vermiculite that was watered to saturation and placed under shade cloth in the Horticulture Greenhouses at the University of British Columbia. We let seedlings establish in soil, watered as needed, for four weeks before a small amount (~5mm x 2mm) of young leaf meristem tissue was harvested from a single individual and immediately preserved in liquid nitrogen. Roughly 25 to 50mm³ of this tissue was divided into two separate 2mL screw-cap tubes, each containing two stainless steel ball bearings of 2mm diameter. Tubes were flash-frozen in liquid nitrogen and used for genomic DNA purification and genome sequencing.

A second individual from the same inbred family was transplanted to an 8" plastic pot and fertilized with 20/20/20 N/P/K fertilizer to encourage rosette growth before being moved outside from October 2012 to April 2013 for cold vernalization at the University of British Columbia Horticulture Greenhouses. In April 2013, we moved the plant back into the greenhouse and sprayed with 2% insecticidal soap to remove pests. Once inside the greenhouse, the plant was left to mature and set seed autonomously via self-pollination. Mature siliques were harvested in July 2013 and seeds were stored in paper envelopes at 4 °C.

S₂ generation

In May 2016, we removed a subset of 10 seeds of the S₁ generation from cold storage and germinated in a 60mm × 15mm petri dish containing filter paper covered with a mixture of autoclaved ProMix soil and silica sand (1:9 ratio). We added distilled water until saturation and thereafter petri dishes were sealed with paraffin wax before storing in the dark at 4° C. Of these, six seedlings germinated and were retained for transcriptome sequencing, divided into one of two treatments. The first true leaf from each of the three plants in the experimental treatment were cut with scissors. We used a Kimwipe tissue saturated with either 0.4mM jasmonic acid (JA) dissolved in 10% ethanol (treatment) or 10% ethanol alone (control), adhered directly to maintain contact the cut site (treatment) or uncut leaf (control). We replaced the saturated Kimwipe every 8h to maintain the signal. After 48h of treatment, we harvested seedlings and preserved treated leaves and root tissue in liquid nitrogen, to be used for RNA purification and transcriptome sequencing.

DNA Isolation & Library Construction

In September 2013, frozen tissue from the S₁ genotype was pulverized and extracted using a Cetyltrimethyl Ammonium Bromide (CTAB) protocol (Clarke, 2009) with the following modifications. After pulverising tissue in a bead mill homogenizer at 60 Hz for 60s, we added 1mL chilled wash buffer (200mM Tris-HCl pH 8.0, 50mM EDTA, 250mM NaCl) and incubated on ice for 10min. The purpose of this wash step is to remove secondary metabolites after disruption of cell walls but prior to cell lysis with CTAB. Following this initial wash step, we spun tubes in a microcentrifuge at 4000 g and 4° C for 10 minutes, then discarded the supernatant and added another 1mL of wash buffer. This was repeated once more for a total of three wash cycles until no coloration was visible in the supernatant. After final discard of the supernatant and addition of warm lysis buffer as per the CTAB protocol, we vortexed tubes briefly to resuspend plant cells. After completion of the CTAB protocol, pellets were dissolved in 50 µL of reverse osmosis (RO) H₂O and sent to Centre d'expertise et de services Génome Québec (Génome Québec) for library preparation and sequencing.

analyze_blastPlus_topHit_coverage.pl, *TrinityStats.pl*). Additionally, we mapped read pairs to the transcriptome assembly to assess read content using **bowtie2** (version 2.3.3.1) with default parameters (Langmead and Salzberg, 2012). We used **Transdecoder** to predict open reading frames in transcripts before using **Trinotate** to annotate and analyze assembled transcripts (Haas et al., 2013).

As a first step in annotation, we established a detailed repeat library. Miniature Inverted Transposable Elements (MITES) represent the most abundant transposable elements in plants and were identified using **MITE Tracker** (Crescente et al., 2018). Long Terminal Repeat (LTR) elements are less common but occupy a larger proportion of the genome, and were identified using the **GenomeTools** package (Gremme et al., 2013). To reduce the number of false positive LTR transposons, only those that contained PPT (poly purine extract) or PBS (primer binding sites) were kept and the rest filtered. We further filtered the LTR candidates to eliminate three main sources of false positives: tandem local repeats such as centromeric repeats, local gene clusters derived from recent gene duplications, and two other transposable elements located in adjacent regions. We also identified elements with nested insertions. After processing known MITES and LTR elements, we identified additional repetitive sequences using **RepeatModeler** against a transposase database and excluded gene fragments using **ProtExcluder**.

The annotation pipeline **maker** (Campbell et al. 2014) was used to identify gene models and predict functional annotations in the draft genome. Both *est2genome* and *protein2genome* modes were used initially to make *ab initio* gene predictions from EST and protein evidence, respectively. The EST evidence was based on our own transcriptome data whereas the protein evidence was gathered from the reference proteomes of six closely related plants available from the SwissProt database: *Arabidopsis thaliana*, *Glycine max*, *Brassica oleracea*, *Medicago truncatula*, *Brassica napus*, and *Brassica rapa*. From the first round of annotation, high confidence models were predicted by the *maker2zff* command with default minimums (50% of splice sites confirmed by EST alignment, 50% of exons match an EST alignment, 50% of exons overlap any evidence, and maximum AED of 0.5). These predictions were used to train the *ab initio* gene predictor **snapp** (Korf, 2004). A second round of **maker** was run using the hidden Markov model (HMM) from **snapp** rather than the *est2genome* mode. All other settings were the same as for the first run, with the transcripts now being used only as evidence to support *ab initio* gene predictions. Two more rounds of annotation and gene prediction improvement followed. An **Augustus** gene prediction file was also generated for use as a second *ab initio* prediction (Stanke and Morgenstern, 2005). For the final round of annotation in **maker**, the final HMM file from **snapp**, the *A. petiolata* species **Augustus** library and gene prediction file were all used in addition to the following settings: always complete, single exons, and using correct EST fusion.

We used Benchmarking Universal Single-Copy Orthologs (**BUSCO**) v4.1 (Simão et al., 2015) to assess the assembly quality of the final draft genome. We used plant lineage delineation from the EmbryophytaDB V10 database, focusing on universal orthologs present in >90% of lineages, resulting in a total of 1,440 BUSCO orthologs. We used **minimap2** check for assembly contiguity and synteny with the model plant *Arabidopsis thaliana* using the TAIR 10 assembly with up to 20% sequence divergence. Plots generated with the R package **pafr**.

The genome annotation files were then curated through **deFusion** to resolve fused gene annotation problems (Wang, 2020), as well as **EvidenceModeler** (Haas et al., 2008) to combine *ab initio* gene predictions and protein and transcript alignments into weighted consensus gene structures. The

functional annotations were then created using NCBI BLAST+ and InterProScan (Jones et al., 2014) by adding new names, domains, and putative functions to improve the utility of the genome database.

Data Availability

Raw data used for genome assembly, transcriptome assembly, and the final draft genome can be found in the NCBI SRA database (accession numbers TBD) under BioProject **SUB9096116** (BioSample Accession: **SAMN17958863**). Analysis scripts are available on GitHub (https://github.com/turkrhen/snapping_turtle_genome_scripts).

Results and Discussion

Whole Genome Shotgun (WGS) sequencing of genomic DNA produced 45.8Gb from 229 million HiSeq reads and an additional 7.9Gb from 15.8 million MiSeq reads. This represents an estimated 68× average coverage of the genome with an estimated genome size of 1.07Gb. This is similar to published size of 1.35Gb estimated by flow cytometry (Barow and Meister, 2002). Initial assembly with **ALLPATHS-LG** resulted in 16,743 contigs longer than 1Kb. Technically, these are scaffolded contigs linked using mate pairs, however we refer to these as contigs to avoid confusion with the scaffolds created by merging heterozygous loci in the **redundans** program. The final assembled genome was 1.08 Gb long across 694 scaffolds larger than 1Kb, and more than 50% of the genome is contained in the ten longest scaffolds (Table 1).

Cytological studies of *A. petiolata* suggest variation in chromosome number and ploidy (Cavers et al., 1979; Weiss-Schneeweiss and Schneeweiss, 2003), and whether the hexaploids are allo- or auto-polyploids. Approximately 36% of contigs (6,045 of 16,743) remain heterozygous after **redundans** assembly; however, average identity within these contigs was 94.8%, meaning that an average of just 2% of sites in the genome assembly are heterozygous. Similarly, just 3.2% of **BUSCO** genes mapped to more than two scaffolds (46 of 1440), consistent with a low level of heterozygosity and minimal duplication of housekeeping genes.

This relatively low level of heterozygosity is consistent with a diploid or highly-inbred autopolyploid. A relatively simple genome combined with geographic variation in ploidy make *A. petiolata* an appealing species to study the role of polyploidy in local adaptation and range expansion, which is an active area of research (e.g. (Pandit et al., 2011; Payseur and Rieseberg, 2016; te Beest et al., 2012)).

Table 1. Assembly statistics for the *Alliaria petiolata* genome

Statistic	Value
# contigs (>= 1000b)	694
# contigs (>= 50,000b)	227
Total length	1,075,010,735
Total length (>= 50,000b)	1,071,536,925
Largest contig	485,611,451
GC (%)	37.2
N ₅₀	121,941,980
N ₇₅	40,840,077
L ₅₀	2

L₇₅ 5
Mean Sequence Length 1,549,006.82

Most (98.7%) of the essential single-copy genes from **BUSCO** mapped to our assembly (Fig. 1), with 71.5% occurring only once in the assembled genome. Similarly, 98% of sequence reads from the transcriptome experiment mapped successfully to the assembled genome. Sequencing of the transcriptome libraries yielded a total of 68.1 Gb from 272.5 million paired reads. Trimming sequence reads for quality reduced usable data by less than 2%.

Figure 1. Percentage of predicted single-copy plant genes from **BUSCO** that are found one (light blue) or more times (dark blue), or are missing (red) or fragmented (yellow) in the annotated genome assembly of *Alliaria petiolata*.

Our *de novo* transcriptome assembly included 699,048 putative isoform “transcripts” representing 535 Mb with N50 of 1,233 base pairs. The minimum transcript length was 201 as set by the Trinity default parameter while 1382 (~1.98%) of transcripts were longer than 5Kb. These transcripts clustered into 350,672 hypothetical genes with an average of 2.18 isoforms and 26,910 (~7.67%) of hypothetical genes having more than five isoforms. A BLAST search of hypothetical genes to the SwissProt protein database matched 10,352 proteins with at least 90% coverage of the query sequence, including 7,930 proteins with 100% coverage (Table 2).

Our TE annotation analysis identified 8,220 unique sequences across the *A. petiolata* genome. Of these, 112 were classified as LTRs with relatively recent origin (99% similarity), 240 as relatively old (85%), and 7,137 were classified as miniature inverted transposable elements (MITE). An additional 731 sequences were found to match the DNA transposase family. After masking TE sequences the final gene set from **maker** included 64,770 gene predictions with an average of 6 exons and an average exon length of 251b (Table 2).

A dot-plot comparison of gene synteny with the model plant *Arabidopsis thaliana* (TAIR 10) revealed large blocks of orthologous sequence (Figure 2). However, the arrangement of synteny blocks shows a complete re-arrangement of *A. thaliana* chromosomes when mapped to the *A. petiolata* contigs. Despite a high level of asynteny that is characteristic of the Brassicaceae family, the conservation of large synteny blocks can help to identify candidate genes and genetic loci of interest for understanding plant invasions. Future research could also investigate whether gene rearrangements occur among geographically and historically isolated populations of *A. petiolata*, and whether this genomic architecture has played an important role in the spread of the species.

Table 2. Summary statistics of genes annotated for the *Alliaria petiolata* genome assembly.

Statistic	Value
Number of genes	64,770
Number of exons	408,155
Number of introns	343,385
Overlapping genes	9,669
Contained genes	1,464
Total gene length	210,804,785
Total exon length	102,316,121
Total intron length	109,175,434
Total CDS length	842,788
% of genome covered by genes	19.6
% of genome covered by CDS	7.8
Mean mRNAs per gene	1
Mean exons per mRNA	6
Mean introns per mRNA	5
Mean gene length	3,255
Mean intron length	318
Mean exon length	251
Mean CDS length	1301

Figure 2. Dot-plot showing blocks of synteny between scaffolds of the *Alliaria petiolata* assembly (x-axis), and five chromosomes of the model plant *Arabidopsis thaliana*. Blue lines show aligned sequences with up to 20% divergence. Vertical dotted lines denote separation of the major scaffolds of the *A. petiolata* assembly.

Conclusions

There is a growing interest in the genetic causes and consequences of range expansion and biological invasion. The field of invasion genetics has emerged from ecological and evolutionary studies of invasive species but lacks well-developed model systems. The draft genome and gene annotation reported here represents an important link from the many field and experimental studies of *A. petiolata* to the genetic architecture of adaptation and invasion. High levels of self-fertility and the resultant low levels heterozygosity observed in the genome will be beneficial for future projects linking ecologically important phenotypes to specific genes. The genomic resources reported here complement available seed resources, experimental findings, and field data to accelerate genomic and molecular studies of *A. petiolata* as a candidate for a model system in invasion genetics.

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