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

Nikolay Alabi, 1 Yihan Wu, 1 Oliver Bossdorf, 2 Loren H. Rieseberg, 3 and Robert I. Colautti 1, 4

1Biology Department, Queen’s University, Kingston, ON K7L 3N6, Canada
2Institute of Evolution and Ecology, University of Tübingen, Tübingen 72076, Germany
3Department of Botany and Biodiversity Research Centre, University of British Columbia, Vancouver, BC V6T 1Z4, Canada

*Corresponding author: Email: robert.colautti@queensu.ca

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 and 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 (EFCC3-3-20) collected from the native range near Venice, Italy, and sequenced paired-end and mate pair libraries at ~70 x coverage. A de novo assembly resulted in a highly continuous draft genome (N50 = 121 Mb; L50 = 2) with 99.7% of the 1.1 Gb genome mapping to scaffolds of at least 50 Kb 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, we found that almost one-third of BUSCO genes (390/1440) mapped to two or more scaffolds despite <2% genome-wide average heterozygosity. The continuity and gene space quality of our draft assembly will enable molecular and functional genomic studies of *A. petiolata* to address questions relevant to invasion genetics and conservation strategies.

Keywords: *Alliaria petiolata*; EFCC3; invasion genetics; garlic mustard; Illumina; mate pairs

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, van 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. Perhaps this is partly due to a lack of genomic resources, which 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 favor *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 2-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 combined influence 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 laboratory experiments, 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. 2021). Understanding the genomic basis of such interactions with soil microbes 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 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 and materials

Study organism and line derivation

*Alliaria petiolata* (M. Bieb.) Cavara and Grande is a biennial herbaceous plant in the Thlaspiideae 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 plant growth and tree regeneration in deciduous forest ecosystems (Stinson et al. 2007; Cipollini and Cipollini 2016). Two ploidy numbers 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 (Weiss-Schneeweiss and Schneeweiss 2003; Esmailbegi et al. 2018). The haploid genome size was previously estimated to be 1.35 Gb based on flow cytometry (Barow and Meister 2002).

**S₀ 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 (~2000 m²) surrounded by agricultural land, about 75 km northwest of Venice, Italy (UTM 45.71°N x 11.72°E). The specific seed line was sampled at 3.2 m along a 10 m sampling transect originating at the edge of population EFCC3. This inbred line (code EFCC3-3-20) is currently available along with other genetic lines as part of the GGMFS seed collections maintained by two of the coauthors in replicate collections in Tübingen, Germany (Bossdorf) and Kingston, Ontario, Canada (Colautti).

**S₁ generation**

In July 2012, 10 seeds of the EFCC3-3-20 genotype from the original S₀ 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 min. 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 (~ 5 mm x 2 mm) of young leaf meristem tissue was harvested from a single individual and immediately preserved in liquid nitrogen. Roughly 25–50 mm³ of this tissue was divided into two separate 2 mL screw-cap tubes, each containing two stainless steel ball bearings of 2 mm diameter. These samples were 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 for cold vernalization from October 2012 to April 2013 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 60 mm x 15 mm 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 0.4 mM 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 8 h to maintain the signal. After 48 h 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 and 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 pulverizing tissue in a bead mill homogenizer at 60 Hz for 60 s, we added 1 mL chilled wash buffer (200 mM Tris-HCl pH 8.0, 50 mM EDTA, 250 mM NaCl) and incubated on ice for 10 min. 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 cold vernalization from October 2012 to April 2013 at the University of British Columbia Horticulture Greenhouses. 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.
We used four separate sequencing libraries for genome assembly: (i) One whole-genome shotgun sequencing library using the Illumina TruSeq DNA v1 preparation kit with a target fragment length of 150 bp. (ii and iii) Two Illumina Nextera MatePair libraries with target lengths of 5 and 10 Kb. These three libraries were multiplexed and sequenced on a single flowcell of Illumina HiSeq 2000 using 2 x 100 bp paired-end (PE) sequencing chemistry. (iv) Target fragment lengths of 450 bp using the Illumina TruSeq DNA v1 and sequenced on Illumina MiSeq with 2 x 250 bp paired-end reads.

RNA isolation and library construction

In March 2017, we pulverized frozen leaf and root tissue for RNA purification in the same manner as the DNA extraction protocol outlined in the previous section. After pulverizing the tissue, we extracted whole RNA from each plant separately using Invitrogen’s TRIzol reagent, following the manufacturer's protocol (Pub No. MAN001271 Rev. A).0). We sequenced four of the six extractions with the highest RNA yields at Génome Québec using the KAPA rRNA-depletion Kit (plant) followed by the Illumina TruSeq LT multiplex kit for sequencing on a single lane of Illumina HiSeq 2500 with 2 x 125 bp paired-end reads. Eight separate libraries were sequenced based on tissue and treatment as described above (see S2 Generation): Control Leaf (CL), Control Root (CR), Treated Leaf (TL), and Treated Root (TR), with two replicates each.

Data processing methods

Raw sequencing data were processed and demultiplexed by Génome Québec, and copied to the Frontenac cluster hosted by the Centre for Advanced Computing (CAC) at Queen’s University and the Cedar cluster maintained by Simon Fraser University on behalf of Compute Canada. The CAC maintains the Rosalind Franklin Cluster for Analysis of Complex Genomes, which is a 256-core computing cluster with 2 TB of RAM. We used this hardware for the memory-intensive steps of de novo genome assembly, with the remaining analyses completed using shared Frontenac and Cedar clusters, and on personal computers. All FASTQ files from both experiments passed quality controls using fastqc (version 0.11.5) (Andrews 2010). We used the raw, demultiplexed FASTQ files for de novo assembly, but the transcriptome data were pre-processing using cutadapt (Martin 2011) to trim adapters and remove reads shorter than 25 b prior to assembly.

Our genome assembly pipeline involved two main steps. First, we used ALLPATHS-LG (Gnerre et al. 2011) version R52488 to assemble contigs from both the HiSeq and MiSeq paired-end libraries and then to link contigs into scaffolds using the 5 and 10 Kb MatePair libraries. We filtered reads that aligned to a draft cpDNA genome but otherwise did not pre-process reads as the ALLPATHS-LG pipeline includes filtering steps. Mitochondrial DNA sequences were manually removed from the final assembly. The analysis parameters included PLOIDY = 2 and HAPLOIDIFY = TRUE to perform a diploid genome assembly. Although our genome is likely hexaploid, polyploid models are not supported by ALLPATHS-LG and low heterozygosity is expected given the high selfing rates in natural populations. Second, we joined scaffolds from ALLPATHS-LG into larger mega-scaffolds using redundans (Przyshcz and Gabaldón 2016) version 0.13c, with the following parameters: identity = 0.9, qers = 5, joins = 5, limit = 1, linkratio = 0.7, mapq = 10, minlength = 1000, and overlap = 0.75. Thus, overlapping scaffolds were collapsed only if they overlapped by at least 75% with an average identity of at least 90%. We repeated this script four times with output scaffolds of the prior run acting as input scaffolds given the long run-time required (~28 days). This second round of assembly combines scaffolds with overlapping similarity, resulting in mega-scaffolds that can span across multiple chromosomes, and other regions of high recombination.

Sequences from the transcriptome experiment were cleaned and assembled with trinity (Grabherr et al. 2011) following protocols outlined in the software documentation and in Haas et al. (2013). We used default parameters and the quality of the assembly was analyzed using the custom perl scripts included in the trinity package to examine full-length transcripts and scaffolds (i.e., Contig N length in analyze_blastPlus_topHit_coverage.pl, TrinityStats.pl). In addition, 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) were identified using MITT Tracker (Crescente et al. 2018). Long Terminal Repeat (LTR) elements 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 tract) or PBS (primer binding sites) within a 300 bp radius 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: A. thaliana, Glycine max, Brassica oleracea, Medicago truncatula, Brassica napus, and B. rapa. From the first round of annotation, high confidence models were predicted by the maker2jff 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 SNAP (Korf 2004). A second round of MAKER was run using the hidden Markov model (HMM) from SNAP 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 SNAP, 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 1440 BUSCO orthologs. We used minimap2 to check for assembly contiguity and synteny with the model plant A. thaliana using the TAIR 10 assembly and up to 20% sequence divergence. We used the R package pafR to generate dot-plots for the four largest scaffolds of A. petiolata and all chromosomes of A. thaliana.

The genome annotation files were curated through deFusion to resolve fused gene annotation problems (Wang et al. 2021). We also used 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.

**Results and discussion**

Most (98.7%) of the essential single-copy genes from BUSCO mapped to our assembly (Figure 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%.

Whole Genome Shotgun (WGS) sequencing produced 45.8 Gb from 229 million HiSeq reads and an additional 7.9 Gb from 15.8 million MiSeq reads. This represents an estimated 68× average coverage of the genome with a predicted genome size of 1.07 Gb. This is 80% of the published size of 1.35 Gb, which was a rough estimate based on flow cytometry (Barow and Meister 2002). Initial assembly with ALLPATHS-LG resulted in 16,743 scaffolds longer than 1 Kb. These scaffolds were linked using mate pairs. We further scaffolded by merging heterozygous loci using the redundans program. The final assembled genome was 1.08 Gb long across 694 scaffolds larger than 1 Kb; more than 75% of the genome is contained in the five 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), but are ambiguous to whether the hexaploids are allo- or auto-polyploids. Approximately 11% of scaffolds (143 of 1,332) remained heterozygous after redundans assembly, compared to 36% (6045 of 16,743) after ALLPATHS-LG but prior to redundancy reduction. The average identity within these scaffolds was 99.61%, meaning that an average of just 0.4% of sites on the heterozygous scaffolds were still heterozygous after redundans assembly. Heterozygosity was similarly low (98.0%) following ALLPATHS-LG assembly but prior to the redundans step, indicating that homozygosity is a feature of the genome rather than an artifact of redundancy reduction. Finally, 27% of BUSCO genes (390 of 1440) were duplicated across the assembly (390 of 1440) but just 3.2% of genes mapped to more than two scaffolds (46 of 1440), indicating that the redundans step was successful in collapsing homologous chromosomes.

This relatively low level of heterozygosity is consistent with a high-inbred autopolyploid or diploid. Geographical variation in ploidy (Weiss-Schneeweiss and Schneeweiss 2003; Esmailbegi et al. 2018) with relatively simple genome 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; te Beest et al. 2012; Payseur and Rieseberg 2016).

![Figure 1](image-url) 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 1233 base pairs. The minimum transcript length was 201 as set by the Trinity default parameter while 1382 (~1.98%) of transcripts were longer than 5 Kb. 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 7930 proteins with 100% coverage (Table 2). The large difference in the number of protein and gene transcript is likely a function of the Trinity algorithm and the ribosome-depletion library preparation method, which includes all RNA transcripts, not just mRNA.

Our TE annotation analysis identified 8220 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 7137 were classified as 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 251 bp (Table 2).

A dot-plot comparison of gene synteny with the model plant A. thaliana (TAIR 10) (Berardini et al. 2015) revealed large blocks of orthologous sequence (Figure 2). However, the arrangement of synteny blocks along chromosomes shows a complete rearrangement of A. thaliana chromosomes when mapped to the A. petiolata scaffolds. Despite a high level of synteny that is characteristic of the Brassicaceae family, the conservation of large synteny blocks holds promise for future research to identify...
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 the four largest scaffolds of the Alliaria petiolata assembly (x-axis) and five chromosomes of the model plant A. 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.

candidate genes and genetic loci of interest for understanding plant invasions. Future research could also investigate whether ploidy variation and 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.

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 represent 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 of 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.

Data availability

Raw data used for genome assembly, transcriptome assembly, and the final draft genome is available from DDBJ/ENA/GenBank under BioProject PRJNA702530, including the WGS project AHZTW000000000 from BioSample SAMN17958863, with sequence reads deposited in SRR13765007 to SRR13765010. The transcriptome analysis BioSamples SAMN17984654—SAMN17984657 and raw sequence reads are also available (SRR13765003 to SRR13765006). The latest genome assembly scripts are available on GitHub (https://github.com/ColauttiLab/Ap_Genome_Assembly).

Acknowledgments

The authors are grateful for bioinformatics support from C Grassa, J Stafford, and H Schmider, hardware support from C MacPhee, and wetlab assistance from M. Todesco, W. Chen, and A. Siew. They also thank M. Todesco and D. Galanti for comments that improved the manuscript.

Funding

This work was supported by the Natural Sciences and Engineering Council of Canada (NSERC) Discovery grants to R.I.C. and L.H.R.

Conflicts of interest

The authors declare that there is no conflict of interest in this work.

Literature cited

Andrews S. 2010. FastQC A Quality Control tool for High Throughput Sequence Data [Online] [WWW Document]. (Accessed: 2020 November 19). http://www.bioinformatics.babraham.ac.uk/projects/fastqc/

Anthony MA, Frey SD, Stinson KA. 2017. Fungal community homogenization, shift in dominant trophic guild, and appearance of novel taxa with biotic invasion. Ecosphere. 8:e01951.doi:10.1002/ecs2.1951.

Barow M, Meister A. 2002. Lack of correlation between AT frequency and genome size in higher plants and the effect of nonrandomness of base sequences on dye binding. Cytometry. 47:1–7. doi:10.1002/cyto.10030.

Barrett SCH. 2015. Foundations of invasion genetics: the Baker and Stebbins legacy. Mol Ecol. 24:1927–1941. doi:10.1111/mec13014.

Barrett SCH, Colautti RI, Dlugosch KM, Rieseberg LH. 2016. Invasion Genetics: The Baker and Stebbins Legacy. Hoboken, NJ: Wiley.

Berardini TZ, Reiser L, Li D, Mezheritsky Y, Muller R, et al. 2015. The Arabidopsis information resource: making and mining the “gold standard” annotated reference plant genome. Genesis. 53: 474–485. doi:10.1002/dvg.22877.
Northeastern Natural. 14:73–88. doi:10.1656/1092-6194(2007)14[73:IOGMI]2.0.CO;2.
Te Beest M, Le Roux JJ, Richardson DM, Brysting AK, Suda J, et al. 2012. The more the better? The role of polyploidy in facilitating plant invasions. Ann Bot. 109:19–45. doi:10.1093/aob/mcr277.
USDA, N. 2020. The PLANTS Database. Greensboro, NC: National Plant Data Team. (Accessed: 2020 April 10). http://plants.usda.gov.
van Boheemen LA, Lombaert E, Nurkowski KA, Gauffre B, Rieseberg LH, et al. 2017. Multiple introductions, admixture and bridgehead invasion characterize the introduction history of Ambrosia artemisiifolia in Europe and Australia. Mol Ecol. 26:5421–5434. doi:10.1111/mec.14293.
Wang W, Zhang X, Dai D-Q. 2021. DeFusion: a denoised network regularization framework for multi-omics integration. Brief Bioinform. 22:bbab057. doi:10.1093/bib/bbab057.
Weiss-Schneeweiss H, Schneeweiss GM. 2003. Karyological investigations of selected angiosperms from Georgia and Azerbaijan. Acta Biologica Cracoviensia Series Botanica. 45:49–56.

Communicating editor: T. Pyhajarvi