



HAL
open science

New WGS data and annotation of the heterosomal vs. autosomal localization of *Ostrinia scapularis* (Lepidoptera, Crambidae) nuclear genomic scaffolds

Louise Brousseau, Sabine Nidelet, Rejane Streiff

► To cite this version:

Louise Brousseau, Sabine Nidelet, Rejane Streiff. New WGS data and annotation of the heterosomal vs. autosomal localization of *Ostrinia scapularis* (Lepidoptera, Crambidae) nuclear genomic scaffolds. *Data in Brief*, 2018, 20, pp.644-648. 10.1016/j.dib.2018.08.011 . hal-02618976

HAL Id: hal-02618976

<https://hal.inrae.fr/hal-02618976>

Submitted on 25 May 2020

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Distributed under a Creative Commons Attribution 4.0 International License



ELSEVIER

Contents lists available at ScienceDirect

Data in Brief

journal homepage: www.elsevier.com/locate/dib

Data Article

New WGS data and annotation of the heterosomal vs. autosomal localization of *Ostrinia scapularis* (Lepidoptera, Crambidae) nuclear genomic scaffoldsLouise Brousseau^{a,b,*}, Sabine Nidelet^a, Réjane Streiff^a^a CBGP, INRA, CIRAD, IRD, Montpellier SupAgro, Univ Montpellier, Montpellier, France^b IRD, UMR DIADE Diversité - Adaptation - Développement, 911 Avenue Agropolis, BP64501, 34394 Montpellier, France

ARTICLE INFO

Article history:

Received 7 May 2018

Received in revised form

9 July 2018

Accepted 3 August 2018

Available online 9 August 2018

Keywords:

Ostrinia scapularis

Genome

NGS

HiSeq2500

Depth analysis

AD-ratio

Structural annotation

Sex-chromosome

Z-heterosome

Autosomes

ABSTRACT

Here, we introduce new whole-genome shotgun sequencing and annotation data describing the autosomal vs. Z-heterosomal localization of nuclear genomic scaffolds of the moth species *Ostrinia scapularis*. Four WGS libraries (corresponding to 2 males and 2 females) were sequenced with an Illumina HiSeq2500 sequencing technology, and the so-called 'AD-ratio' method was applied to distinguish between autosomal and Z-heterosomal scaffolds based on sequencing depth comparisons between homogametic (male) and heterogametic (female) libraries. A total of 25,760 scaffolds (corresponding to 341.69 Mb) were labelled as autosomal and 1273 scaffolds (15.29 Mb) were labelled as Z-heterosomal, totaling about 357 Mb. Besides, 4874 scaffolds (29.07 Mb) remain ambiguous because of a lack of AD-ratio reproducibility between the two replicates. The annotation method was evaluated *a posteriori*, by comparing depth-based annotation with the exact localization of known genes. Raw genomic data have been deposited and made accessible via the EMBL ENA BioProject id PRJEB26557. Comprehensive annotation is made accessible via the LepidoDB database (http://bipaa.genouest.org/sp/ostrinia_scapularis/download/genome/v1.2/).

© 2018 The Authors. Published by Elsevier Inc. This is an open access article under the CC BY license (<http://creativecommons.org/licenses/by/4.0/>).

* Correspondence to: French Institute for Sustainable Development (IRD), UMR DIADE, 911 Av Agropolis BP64501, 34394 Montpellier, France.

E-mail address: louise.brousseau@ird.fr (L. Brousseau).

<http://dx.doi.org/10.1016/j.dib.2018.08.011>

2352-3409/© 2018 The Authors. Published by Elsevier Inc. This is an open access article under the CC BY license (<http://creativecommons.org/licenses/by/4.0/>).

Specifications Table

Subject area	Biology
More specific subject area	Genomics, Bioinformatics
Type of data	DNA sequence reads and annotation table
How data was acquired	Shotgun whole genome sequencing (Illumina HiSeq2500)
Data format	Raw (2 × 125 raw reads) and Analyzed (Scaffolds annotation)
Experimental factors	Wild-type specimens collected in the field
Experimental features	Genome: DNA sequencing
Data source location	Abbeville, France (50° 8'11.03"N; 1°49'49.22"E)
Data accessibility	Raw reads are accessible through the EMBL ENA BioProject id PRJEB26557 (https://www.ebi.ac.uk/ena/data/view/PRJEB26557) Annotation data are made accessible via the LepidoDB database (http://bipaa.genouest.org/sp/ostrinia_scapularis/download/genome/v1.2/)
Related article	B. Gschloessl, F. Dorkeld, P. Audiot, A. Bretaudeau, C. Kerdelhué, R. Streiff (2018) <i>De novo</i> genome and transcriptome resources of the Adzuki bean borer <i>Ostrinia scapularis</i> (Lepidoptera: Crambidae). <i>Data in Brief</i> . doi 10.1016/j.dib.2018.01.073 [1]

Value of the data

- This article enriches and updates the annotation of *Ostrinia scapularis* (Lepidoptera) nuclear genome recently published by Gschloessl et al. [1] with an accurate annotation of the chromosomal localization of the scaffolds constituting the nuclear genome assembly.
- The new genomic data acquired here (whole-genome shotgun sequencing of four libraries, two males and two females) will enrich the public sequence database for this species.
- WGS sequencing depth analysis is a promising method to retrieve the autosomal or heterosomal localization of assembly fragments (scaffolds or contigs) obtained through *de novo* assembly.
- The annotation data released here provide key information about the autosomal vs. Z-heterosomal localization of scaffolds described in Gschloessl et al. [1].
- Such annotation is of great value for future evolutionary studies, as genome-wide population genomics analyses (e.g. continent-scale phylogeography, host-plant adaptation studies etc.) may be dramatically sensitive to the confounding influence of autosomal and heterosomal evolutionary histories (because of different inheritance, ploidy levels, recombination rates, effective population size and genetic drift).

1. Data

The dataset described here is composed of new whole-genome sequencing (WGS) data (paired-end sequencing of four libraries with an Illumina HiSeq2500 sequencing technology) and a new annotation of autosomal and heterosomal scaffolds of the nuclear genome of the moth species *Ostrinia scapularis*. These new data are complementary to the scaffold-level nuclear genome (hereafter *OSCA*) recently assembled for this species [1]. Specifically, we applied the *AD-ratio* method originally developed by Bidon et al. [2] to compare sequencing depth between male and female libraries, and we introduce an accurate labelling (autosomal vs. Z-heterosomal) of the scaffolds of *OSCA* genomic reference which enriches preliminary structural and functional annotations described in Gschloessl et al. [1].

2. Experimental design, materials, and methods

2.1. Species model

O. scapularis (i.e. the Adzuki bean borer) is a phytophagous moth species living on a variety of dicotyledon plants (e.g. *Humulus lupulus*, *Artemisia vulgaris*, *Cannabis sativa*) across Europe, and phylogenetically close to the European corn borer (*O. nubilalis*), a major maize pest worldwide. In this species, 31 pairs of chromosomes are expected (30 autosomal pairs and one heterosomal pair) with a ZZ/ZW sex determination: males are homogametic (ZZ) and females are heterogametic (ZW).

2.2. Sampling and DNA extraction

O. scapularis diapausing larvae were collected in stems of wild mugwort in northern France (Abbeville, Picardie) and stored in 95% ethanol at -20°C . Genomic DNA (gDNA) was extracted using BioBasic '96-well plate animal genomic DNA mini-preps' extraction kits (Euromedex) according to manufacturer's instructions. gDNAs were quantified using a NanoDrop 8000 Spectrophotometer (Thermo Scientific). The sex of each sample was characterized according to the molecular method described in Orsucci et al. [3]: sex-linked microsatellite markers, ONW1 and ONZ1 (specific of W and Z heterochromosomes respectively), were amplified simultaneously using the Multiplex PCR Master Mix (Qiagen). Four specimens were finally retained among the best quality DNAs: two males (IDs 12098 and 12114) and two females (IDs 12099 and 12111).

2.3. Library preparation and sequencing

Four libraries (one per sample) were prepared according to the 'TruSeq Nano DNA Library Preparation Guide' (Illumina, <https://support.illumina.com/downloads/truseq-nano-dna-library-prep-guide-15041110.html>), starting with 100 ng of gDNA per library. According to manufacturer's instruction, library preparation workflow included: (1) gDNA fragmentation with a Covaris S220, (2) libraries end-repair and sizing, (3) 3' ends adenylation, (4) adapters ligation, (4) DNA fragments enrichment, and (5) libraries normalization and pooling. At the end of the enrichment step (4), libraries quality and quantity were evaluated using both a 'DNA 1000 ship' ('Agilent Technologies 2100 Bioanalyzer') and a 'KAPA Library Quantification Kit' (KAPA Biosystems). Libraries varied between 515 and 533 bp in size, and were pooled 2 by 2 (one male and one female library per pool) in 20 nM equimolar mixture. Shotgun libraries were sequenced with an Illumina HiSeq2500 paired-end (2×125 bp) sequencing technology by the ISO9001:2008 Montpellier Genomix facility (MGX, France, <http://www.mgx.cnrs.fr>).

2.4. Bioinformatics pipeline

The bioinformatics pipeline is detailed in [Supplementary file 1](#). In brief, raw reads were cleaned and mapped against the reference nuclear genome *OSCA* as follow:

- (1) Reads that did not pass *Illumina* chastity filter (i.e. purity filter PF) were discarded with *zcat* and *grep*.
- (2) phiX control reads were removed by mapping raw reads against phiX genome with *bowtie2* [4]: only unmapped reads were used in the following.
- (3) Individual bases of low quality (*phred-score* < 25) were masked using *fastq_masker* (http://hanonlab.cshl.edu/fastx_toolkit/) with parameters $-q\ 25 -Q\ 33$.
- (4) Reads of low quality and orphan reads were discarded using *sickle* [5], with parameter $-q\ 25$.
- (5) Cleaned reads were mapped against the reference nuclear genome *OSCA* [1], separately for each library, using *bwa aln* and *bwa sampe* [6].
- (6) SAM files (.sam) were converted into sorted BAM with *samtools view* (with parameter $-q\ 20$ to discard multiple mapped reads) and *samtools sort* [7].

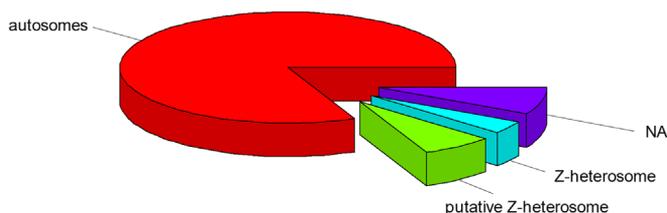


Fig. 1. Total assembly length (~ 420 Mb) partitioning into autosomal (~ 342 Mb), Z-heterosomal (~ 15 Mb), putatively Z-heterosomal (~ 29 Mb) and un-annotated (~ 33 Mb) genomic regions.

(7) Multiple BAM (.mpileup) were generated with samtools mpileup and converted into tabular 'synchronized' files with 'mpileup2sync' perl script originally implemented in Popoolation2 [8].

Synchronized files were further handled with perl and R to estimate base depth, per-scaffold mean depth, and to compute *AD-ratios* between homogametic and heterogametic libraries, see [Supplementary file 2](#). The *AD-ratio* method [2] is conceptually based on the simple assumption that the ratio of sequencing depth between homogametic (here, male ZZ) and heterogametic (here, female ZW) libraries - standardized by the number of mapped reads for each library - would be 1 for autosomal scaffolds, 2 for Z-heterosomal scaffolds and 0 for W-heterosomal scaffolds, (see [Supplementary file 3](#) for additional information about scaffold-specific *AD-ratio* estimation). Note here that the W-chromosome is absent from the *OSCA* nuclear reference which was drawn from a single male (ZZ) [1].

A total of 25,760 scaffolds (corresponding to 341.69 Mb) were identified as autosomal and 1273 scaffolds (15.29 Mb) were identified as Z-heterosomal, totaling about 357 Mb, [Fig. 1](#). Besides, 4874 scaffolds (29.07 Mb) were ambiguously annotated because of a lack of *AD-ratio* reproducibility between the two replicates, putting thus the emphasis on the necessity to use two independent biological replicates. Last, 18,831 scaffolds remained un-annotated, because of insufficient mapping depth (< 4X in average) in one or several libraries. They represent ~ 33 Mb, which corresponds to ~ 8% of total assembly length, indicating that the subset of un-annotated scaffold is largely enriched in short scaffolds. Comprehensive annotation is provided in [Supplementary file 4](#) and is made publically-available via the LepidoDB database (http://bipaa.genouest.org/sp/ostrinia_scapularis/download/genome/v1.2/).

2.5. A posteriori validation

We compared the 'blinded' annotation based on *AD-ratios* for scaffolds holding either autosomal or Z-heterosomal known candidate genes, including six olfactory receptors, *OR1* to *OR6* [9,10], and two Z-linked genes, *Tpi* and *Kettin* respectively [11]. To do that, candidate genes were localized by blastn against the nuclear reference *OSCA* using the program blastall (*e-value* < 10⁻²⁰). The Z vs. autosomal annotation based on *AD-ratios* was totally consistent with the actual Z or autosomal localization of the candidate genes, even in cases of a lack of reproducibility between replicates ([Supplementary file 5](#)).

Acknowledgments

This work was supported by a grant from the ANR Adapt-Ome (ANR-13-BSV7-0012) funded by the French National Research Agency and coordinated by Réjane Streiff. We thank Bernhard Gschoessl and Anthony Bretaudeau for access and data entry in the LepidoDB database. We are grateful to Montpellier GenomiX (MGX, France) for providing sequencing facility, and to the Genotoul bioinformatics platform Toulouse Midi-Pyrenees (Bioinfo Genotoul, Toulouse, France) for providing computing and storage resources.

Transparency document. Supplementary material

Transparency document associated with this article can be found in the online version at <https://doi.org/10.1016/j.dib.2018.08.011>.

Appendix A. Supplementary material

Supplementary files associated with this article can be found in the online version at <https://doi.org/10.1016/j.dib.2018.08.011>.

References

- [1] B. Gschloessl, F. Dorkeld, P. Audiot, A. Bretaudeau, C. Kerdelhué, R. Streiff, De novo genome and transcriptome resources of the Adzuki bean borer *Ostrinia scapularis* (Lepidoptera: crambidae), Data Brief 17 (2018) 781–787. <http://dx.doi.org/10.1016/j.DIB.2018.01.073>.
- [2] T. Bidon, N. Schreck, F. Hailer, M.A. Nilsson, A. Janke, Genome-wide search identifies 1.9 Mb from the polar bear Y chromosome for evolutionary analyses, Genome Biol. Evol. 7 (2015) 2010–2022. <http://dx.doi.org/10.1093/gbe/evv103>.
- [3] M. Orsucci, P. Audiot, A. Pommier, C. Raynaud, B. Ramora, A. Zanetto, D. Bourguet, R.P. insects, P. Streiff, Host specialization involving attraction, avoidance and performance, in two phytophagous moth species, J. Evol. Biol. 29 (2016) 114–125. <http://dx.doi.org/10.1111/jeb.12766>.
- [4] B. Langmead, S.L. Salzberg, Fast gapped-read alignment with Bowtie 2, Nat. Methods. 9 (2012) 357–359. <http://dx.doi.org/10.1038/nmeth.1923>.
- [5] N.A. Joshi, J.N. Fas, Sickle: A Sliding-window, Adaptive, Quality-based Trimming tool for FastQ Files (Version 1.33). Available at (<https://github.com/najoshi/sickle>).
- [6] H. Li, R. Durbin, Fast and accurate short read alignment with Burrows–Wheeler transform, Bioinformatics 25 (2009) 1754–1760. <http://dx.doi.org/10.1093/bioinformatics/btp324>.
- [7] H. Li, B. Handsaker, A. Wysoker, T. Fennell, J. Ruan, N. Homer, G. Marth, G. Abecasis, R. Durbin, G.P.D.P. Subgroup, The sequence alignment/map format and SAMtools, Bioinformatics 25 (2009) 2078–2079. <http://dx.doi.org/10.1093/bioinformatics/btp352>.
- [8] R. Kofler, R.V. Pandey, C. Schlötterer, PoPoolation2: identifying differentiation between populations using sequencing of pooled DNA samples (Pool-Seq), Bioinformatics 27 (2011) 3435–3436. <http://dx.doi.org/10.1093/bioinformatics/btr589>.
- [9] J.-M. Lassance, S.M. Bogdanowicz, K.W. Wanner, C. Löfstedt, R.G. Harrison, Gene genealogies reveal differentiation at sex pheromone olfactory receptor loci in pheromone strains of the European corn borer *Ostrinia nubilalis*, Evolution 65 (2011) 1583–1593. <http://dx.doi.org/10.1111/j.1558-5646.2011.01239.x>.
- [10] Y. Yasukochi, N. Miura, R. Nakano, K. Sahara, Y.P. insects, P. Ishikawa, Sex-linked pheromone receptor genes of the European corn borer *Ostrinia nubilalis* are in tandem arrays, PLoS One 6 (2011) e18843. <http://dx.doi.org/10.1371/journal.pone.0018843>.
- [11] T. Malausa, L. Leniaud, J.-F. Martin, P. Audiot, D. Bourguet, S. Ponsard, S.-F. Lee, R.G. Harrison, E. Dopman, Molecular differentiation at nuclear loci in French host races of the European corn borer *Ostrinia nubilalis*, Genetics 176 (2007) 2343–2355. <http://dx.doi.org/10.1534/genetics.107.072108>.