

# ERGA Assembly Report

v24.10.15

Tags: ERGA-BGE

TxID	446453
ToLID	<b>inOsmFulv5</b>
Species	<i>Osmylus fulvicephalus</i>
Class	Insecta
Order	Neuroptera

Genome Traits	Expected	Observed
Haploid size (bp)	625,527,197	674,675,484
Haploid Number	8 (source: ancestor)	8
Ploidy	2 (source: ancestor)	2
Sample Sex	XY	XY

## EBP metrics summary and curation notes

Obtained EBP quality metric for pri: 7.8.Q68

The following metrics were automatically flagged as below EBP recommended standards or different from expected:

### Curator notes

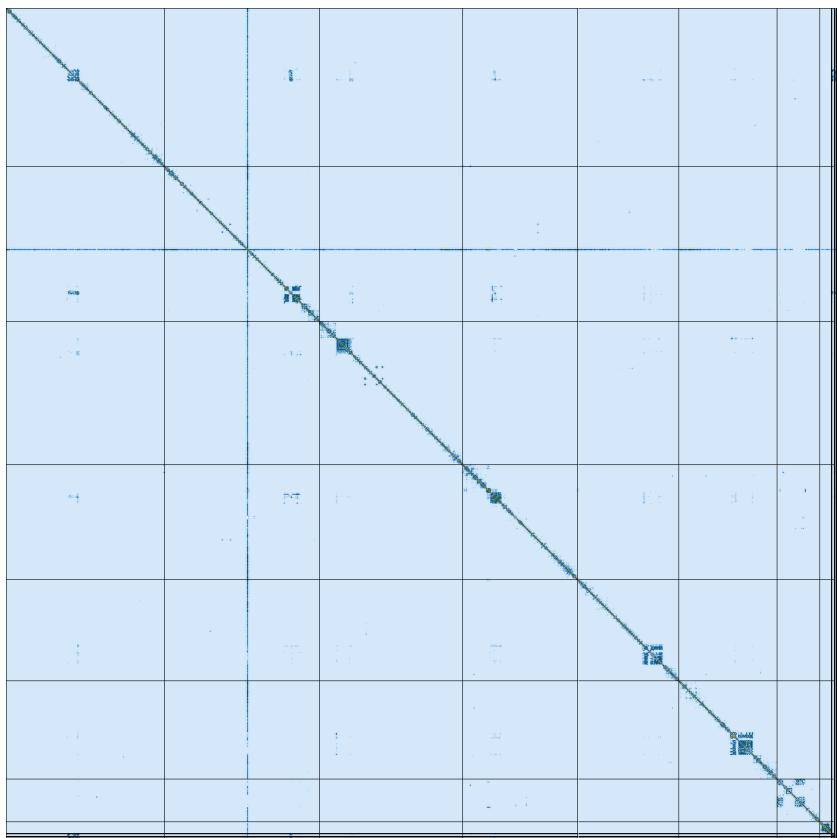
. Interventions/Gb: 9  
. Contamination notes: "No contaminations were detected with FCS-GX."  
. Other observations: "PacBio reads were subsampled to 60X. Reads were assembled with hifiasm (yield: 488M, N50: 49.1Mb, 176 contigs). FCS-GX detected no contaminations. Purge\_dups was run on the contigs but only the the dups.bed file was used in the manual curation step, as it removed too much sequence and telomere motifs. Due to the low HiC quality (2.8M cis-read pairs >40Kb) the scaffolding was manually done in the first manual curation round. In this step the information from tiara, mitohifi, blobtools and purge\_dups was used and 110 contigs (15M) were removed. SUPER\_8 and SUPER\_11 had a very low coverage profile - roughly half coverage. A synteny analysis with GCA\_958496175, GCA\_958496155, GCF\_905475395.1 and GCA\_020423425.1 suggests that SUPER\_8 is the X chromosome. Super\_11 is the Y chromosomes. Furthermore, the chromosome structure seems to be very stable within these insects (6 "closely" related species). All of them suggest the following joins: SUPER\_5 AND SUPER\_7 (both telomere motifs available), SUPER\_2 AND SUPER\_10, SUPER\_6 and SUPER\_9 (one telomere motif). Then these weird repeats (centromers?) are always within the chromosomes. "

## Quality metrics table

Metrics	Pre-curation pri	Curated pri
Total bp	666,151,682	674,675,484
GC %	36.31	36.29
Gaps/Gbp	90.07	53.36
Total gap bp	12,000	7,200
Scaffolds	123	24
Scaffold N50	104,575,600	116,168,846
Scaffold L50	3	3
Scaffold L90	16	6
Contigs	183	60
Contig N50	22,977,000	51,542,635
Contig L50	8	5
Contig L90	43	16
QV	70.6914	68.5927
Kmer compl.	90.8782	90.8268
BUSCO sing.	93.3%	94.4%
BUSCO dupl.	1.6%	0.4%
BUSCO frag.	1.4%	1.4%
BUSCO miss.	3.7%	3.8%

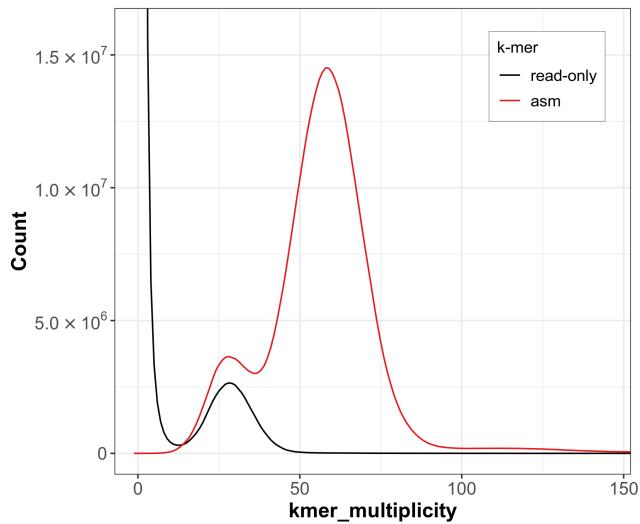
BUSCO: 5.8.3 (euk\_genome\_min, miniprot) / Lineage: endopterygota\_odb12 (genomes:76, BUSCOs:3754)

# HiC contact map of curated assembly

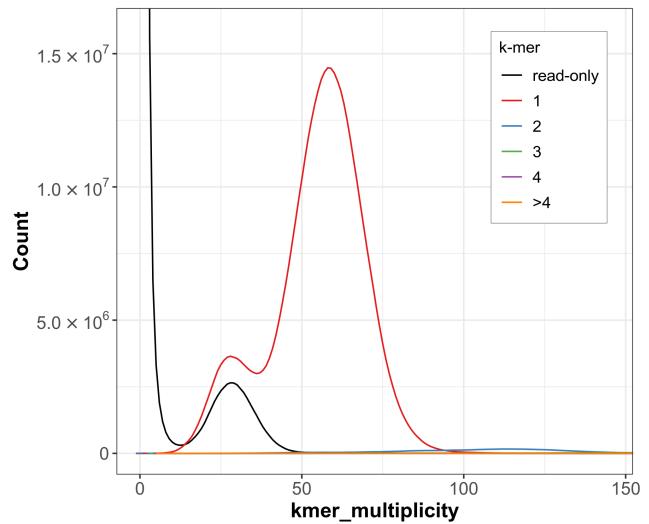


pri [LINK]

# K-mer spectra of curated assembly

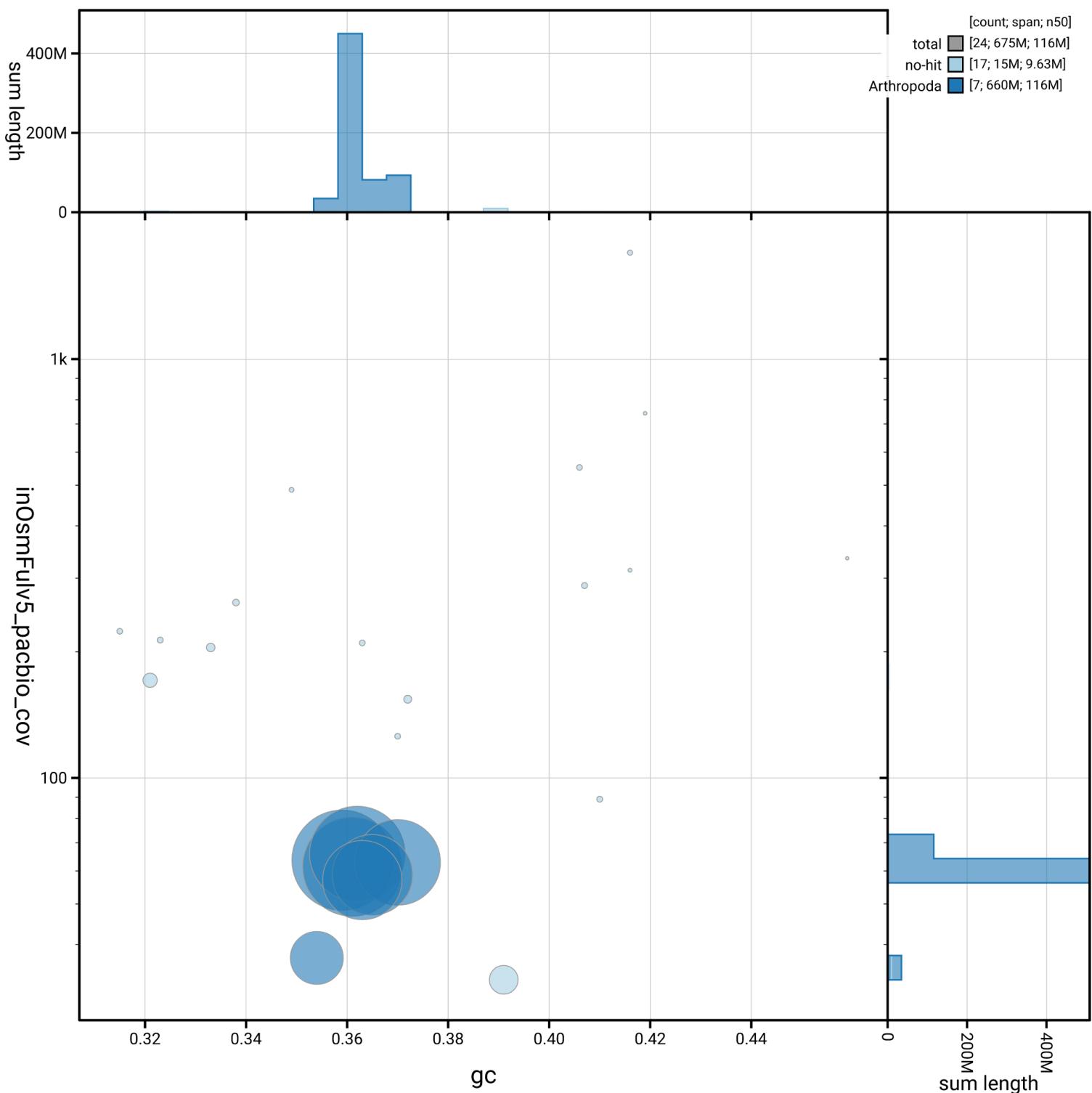


Distribution of k-mer counts coloured by their presence in reads/assemblies



Distribution of k-mer counts per copy numbers found in asm

# Post-curation contamination screening



**pri.** Bubble plot circles are scaled by sequence length, positioned by coverage and GC proportion, and coloured by taxonomy. Histograms show total assembly length distribution on each axis.

## Data profile

	Data	HiFi	HiC
Coverage	164x	49x	

## Assembly pipeline

- **Hifiasm**
  - |\_ ver: 0.25.0-r726
  - |\_ key param: 13
- **purge\_dups**
  - |\_ ver: 1.2.6
  - |\_ key param: NA

## Curation pipeline

- **GRIT\_Rapid**
  - |\_ ver: 1a3d79a8
  - |\_ key param: NA
- **HiGlass**
  - |\_ ver: 0.10.4
  - |\_ key param: NA

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