Explore the Genome Browsers created by G-OnRamp

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07/2017

# 1. Introduction

This exercise will explore the *Drosophila miranda* UCSC Assembly Hub and the JBrowse Hub created by G-OnRamp in order to illustrate how you can use the Genome Browsers to address interesting biological questions.

# 2. Use Galaxy to explore the genome assembly

Open a new web browser window and navigate to the G-OnRamp server at <http://cloud5.galaxyproject.org/>. (Note that this instance will only be available during the G-OnRamp workshop.) Log into your account and then import the “Drosophila miranda MSH22 UCSC Genome Browser” History from “Shared Data”.

**Q1. Which dataset in the History contains the sequences of the *D. miranda* whole genome assembly? How many scaffolds are in the *D.* *miranda* whole genome assembly?**

**Q2. Which scaffold is the largest in the *D. miranda* assembly? How long is the largest scaffold?** (Hint: run the **twoBitInfo** tool on the dataset “12: faToTwoBit on data 8”. Select the “Sequence lengths” option for the “Type of output file” field.)

**Q3. How many gaps are in the assembly? How many gaps are in each scaffold?**

[Hint: You can use the **Group** tool to group the gap items by column (*e.g.*, scaffold)]

# 3. Use the UCSC Assembly Hub to explore a genomic region

Open the *D. miranda* UCSC Genome Browser Assembly Hub. Navigate to position scaffold\_6:789,800-806,800. Scroll down to the track configuration section and hide the HISAT RNA-Seq alignment tracks (i.e., “SRR364798 HISAT s” and “SRR364800 HISAT s”) in order to simplify the display.

**Q4. How many Augustus gene predictions are in this region? How many of these Augustus gene predictions overlap with features in either the BLAST or BLAT alignment tracks?**

**Q5. For the Augustus gene predictions that do not overlap with features in the TBLASTN or BLAT alignment tracks, perform a NCBI BLASTP search of the translated protein sequence against the “Reference proteins (refseq\_proteins)” database. Based on the BLASTP search results, do these predictions correspond to protein-coding genes in *Drosophila*?**

(Hint: To obtain the protein sequence of a predicted gene, click on the feature in the Genome Browser image, and then click on the “Translated Protein” link. The NCBI BLAST web server is available at <https://blast.ncbi.nlm.nih.gov/Blast.cgi>.)

**Q6. Change the display mode of the BLAT alignment track to “full”. Click on the BLAT alignment to the transcript NM\_143661. What is the name of this transcript? How long is this transcript? Scroll down to the “FEATURES” section of the GenBank report for NM\_143661. Where is the coding region within this transcript?** (Hint: see the CDS section of the FEATURES table.)

**Q7. Scroll down to the “Genomic Alignments” section of the BLAT Alignment details page. What is the percent identity between the NM\_143661 and scaffold\_6? What is the orientation of the transcript alignment? Does the alignment include the entire length of the NM\_143661 transcript?**

**Q8. Click on the alignment statistics link (next to the “browser” link) under the “Genomic Alignments” section to view the transcript alignment. What do the capital blue and red letters within the cDNA and genomic sequences symbolize?**

**Q9. Go back to the Genome Browser view of the region at scaffold\_6:789,800-806,800. There is another set of BLAT and TBLASTN alignments at 804,000–805,500 that is also supported by the RNA-Seq data from the virgin males (SRR364798) and virgin females (SRR264800) samples. Based on the RNA-Seq read coverage tracks (“SRR364798 RNA-Seq” and “SRR364800 RNA-Seq”) and the regtools splice junction tracks (“SRR364798 Splice” and “SRR364800 Splice”), how many introns does this feature have?**

**Q10. Using the BLAT alignments and the procedure described above, characterize the feature located at 804,000-805,500 of scaffold\_6.**

**Q11. The RefSeq accession number for the A isoform of *CG32850* is NM\_166753, and the RefSeq accession number for the B isoform of *CG32850* is NM\_001272124. Can you find the BLAT alignments to these transcripts in the *D. miranda* assembly? Are the genes *Rad23*, *Zip102B*, and *CG32850* syntenic between *D. melanogaster* and *D. miranda*?** (Hint: refer to FlyBase for information of genes Rad23, Zip102B, and CG32850 in*D. melanogaster*)

# 4. Use the JBrowse to explore a genomic region

Go back to the web browser window with the G-OnRamp server. Import the “Drosophila miranda MSH22 JBrowse” History from “Shared Data”. Expand the “41: JBrowse Archive Creator” dataset and then click on the “View JBrowse” link to launch the JBrowse version of the *D. miranda* Genome Browser.

Enter the accession number for the B isoform of *CG32850* (NM\_001272124) into the search box of the JBrowse and then click on “Go” to navigate to *CG32850*.

**Q12. What is the orientation of the transcript relative to scaffold\_6 (1 for plus strand, -1 for minus strand)? How many alignment blocks are there? Where are these alignment blocks located on scaffold\_6?** (Hint: Right click on a feature in the Blat Alignment track and select the “View alignment” option to view the BLAT alignment.)

**Q13. Examine the “Augustus”, “GlimmerHMM”, and “SNAP” evidence tracks on JBrowse. Are there any genes predicted in this region? For the gene(s) predicted by Augustus, perform a NCBI BLASTP search of the translated protein sequence against the *D. melanogaster* protein sequences in the nr database. Based on the BLASTP search results, do the Augustus gene prediction(s) correspond to protein-coding genes in *Drosophila melanogaster*?**

[Hint: to obtain the protein sequence of a predicted gene, right clicking on the gene, and then select “View translated protein” option from the drop-down menu. You can use the “Organism” field (under the “Choose Search Set” section) of the NCBI BLASTP search interface to restrict the search to “*Drosophila melanogaster* (taxid:7227)”.]