

### Part 3 -

Load the attached mouse files to Galaxy. They are ungroomed single-end FASTQ files with Illumina 1.5 phred encoding from a ChIP-seq experiment and downsampled to a part of chromosome 19. In Galaxy, run the **FASTQ Groomer** tool to convert the reads to fastqsanger format. Then, use **Trimmomatic** to require a phred score greater than or equal to 20. Align the trimmed reads to the mm9 reference with **Map with BWA**. Finally, run **MACS2 callpeak** on the experimental ChIP-seq with the control output as the control.

**\*On Trimmomatic, the sliding window trimming operation was used with 4 bases average across, and 20 average quality minimum.**

- Retrieve the peaks in tabular format. Find the interval chr19:37,340,169-37,340,716. List the value in the fold\_enrichment column.

**27.13470**

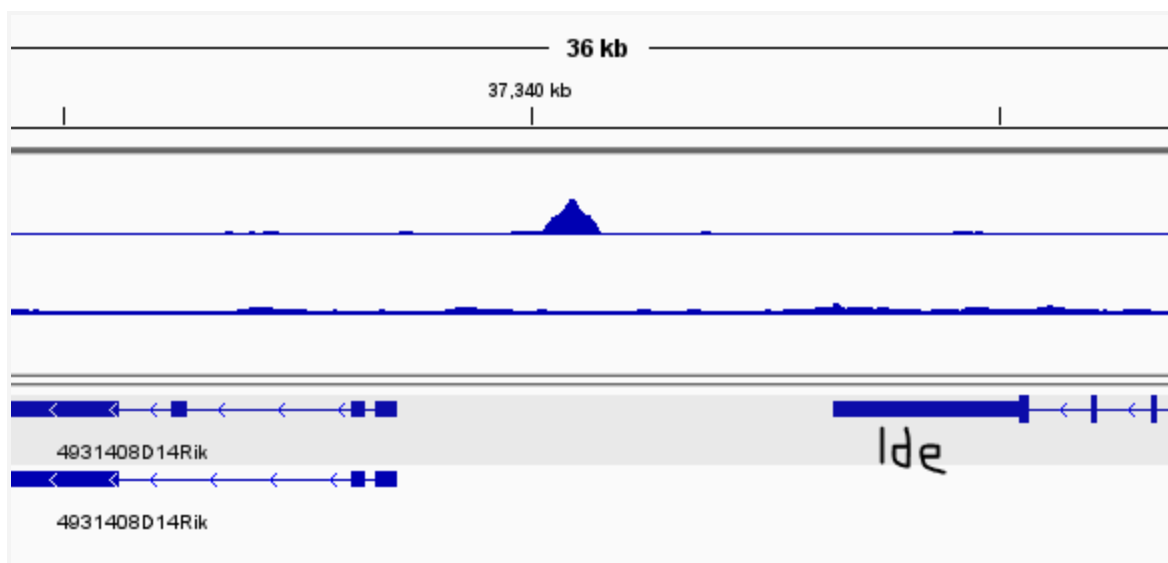
- Load both bedgraph files into IGV, mm9. Go to the interval from Part 3a. What is the nearest transcript?

**The nearest transcript from Refseq genes is the gene 4931408D14Rik.**

- Relative to the nearest two genes, where (upstream, exon, intron, downstream) is the MACS peak?

**The MACS peak is between the two nearest genes of 4931408D14Rik and Ide. It is downstream from 4931408D14Rik and upstream from Ide.**

- Submit a screenshot from IGV showing both the MACS peak and a small portion of the nearest two genes.



\*Fuller view also attached