Present for Sally

MBL, from Ian, May 2014

• Go to Gert's website:

Genome assembly Xentr v7.1

The "tracks" file below is a bigWig file that can be used to visualize *H3K4me3, TBP, H3K4me1, RNAPII, H3K27me3, RNAseq, MethylCap and Input tracks* of multiple stages (stages 9, 12, 16 and 30) at the genome browser at NIMR. Alternatively, browse genes for H3K4me3 and H3K27me3 profiles at our gene profile page.

File Name:	Date Created:	File Size:
mostconserved_gmap_k12.bed.gz	7/17/2013	5,814 KB
Veenstra_tracks_JGI-7-13.wig.gz	7/15/2013	1 KB
ktev_v3-4_JGI7-1_all.bed.gz	12/4/2013	1,460 KB
ktev_v3-4_JGI7-1_genename.bed.gz	6/14/2013	999 KB

File Name:	Date Created:	File Size:
CpGI_Gardiner-Garden_xenTro2.bed.gz	3/23/2011	1,488 KB
CpGI_Takai_Jones_xenTro2.bed.gz	3/23/2011	212 KB
DNAme_St12_500mM_xenTro2.wg.gz	3/23/2011	5,473 KB
DNAme_St12_700mM_xenTro2.wg.gz	3/23/2011	5,561 KB

Newer time-series data

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- All his data are aligned already wig files are a lot like bam files or bedgraph files
- The newer stuff is aligned to Trop genome v7.1, which is not available on the UCSC browser
- So instead you have to go to some other browser to see the reads – he links to it on his lab resources page. It's the NIMR link

• NIMR link in the box:

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- Unfortunately the NIMR browser sucks, and there is something wrong with Gert's site such that you can't upload those newer tracks there.
- I just wrote him about it.

Older data

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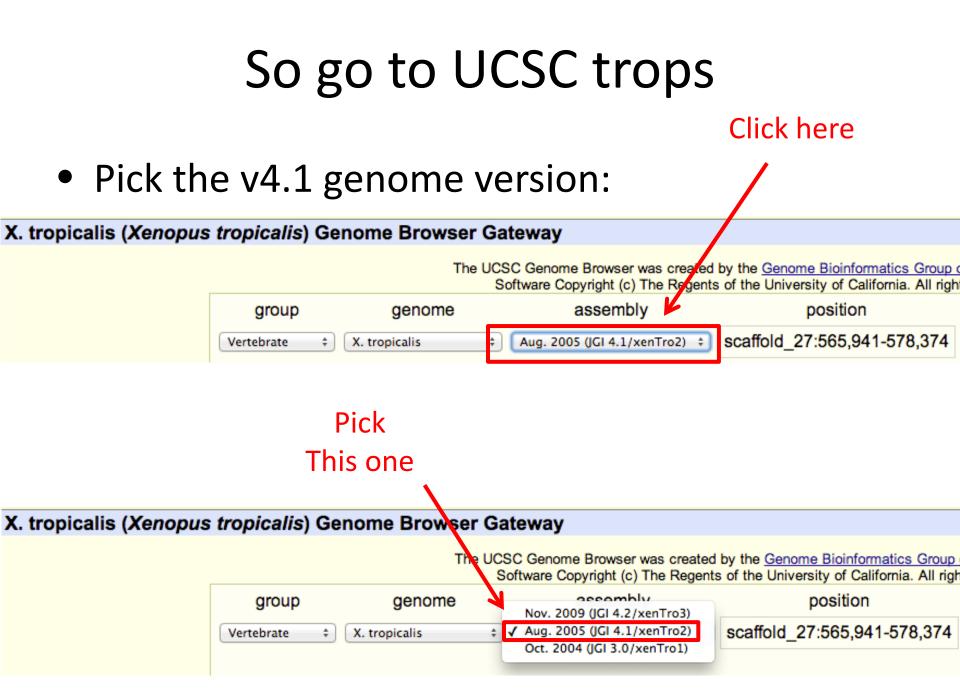
• Older data are aligned to trops v4.1

Genome assembly Xentr v7.1

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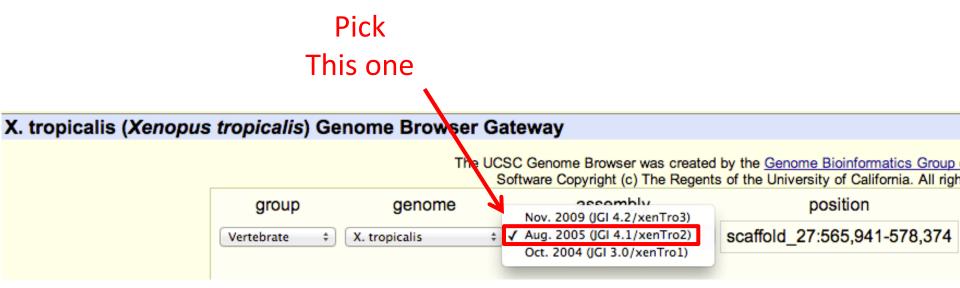
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So go to UCSC trops

- Pick the v4.1 genome version:
- Because Gert already aligned these, if you load them to the wrong genome build the numbers won't agree and it won't work
- For bedgraph and wig files, this mistake happens a lot. Pay attention to genome version!



Get the track links from Gert's site

Genome assembly Xentr v7.1

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Right-click, "copy link address"

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Go back to the UCSC browser for trops v4.1

X. tropicalis + Aug. 2005 (JGI 4.1/xenTro2) + scaffold_27:565,941-578,374 enter position, Click here to reset the browser user interface settings to their defaults. track search add custom tracks track hubs configure tracks and display	ene symbol or search terms submit

Add track to browser

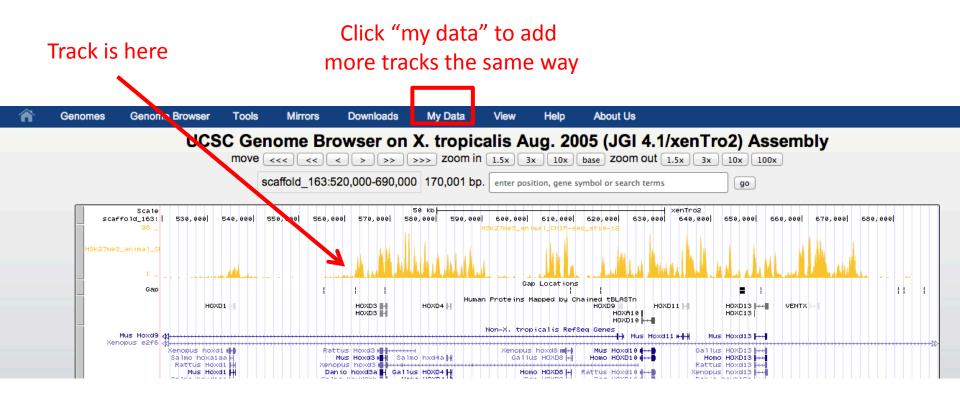


You'll see this now

Manage Custom Tracks

genome: X. tropicalis assembly: Aug. 2005 (JGI 4.1/xenTro2) [xenTro2]





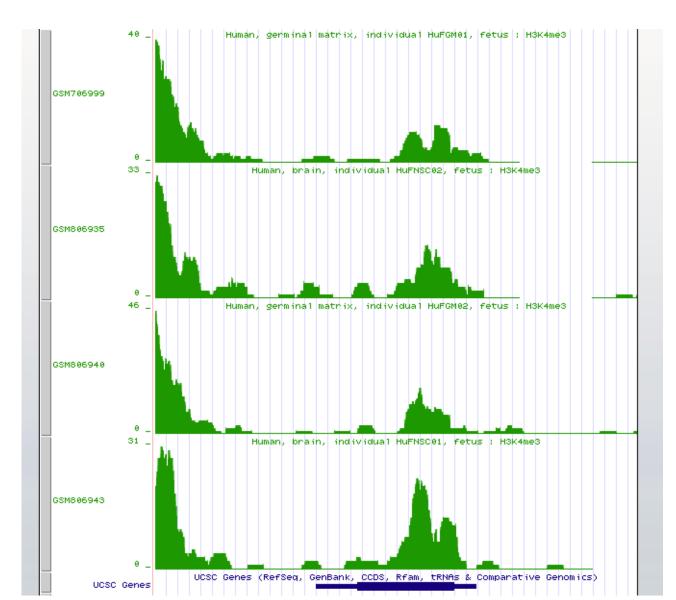
Also in 'my data' you can save sessions to keep the same tracks for later so you don't have to keep reloading them. Very useful.

- You pretty much follow this for everything.
- After the course, you know how to map with bowtie2 and can download fastqs and map those yourself if you want, or just use the wigs/bedgraphs from somebody else

- NCBI geo often has wig or bedgraph files. Can also get fastq and align yourself
- UCSC has some nice collections:
- <u>http://genome.ucsc.edu/cgi-bin/hgHubConnect?db=hg19</u>
- Also encode project:
- <u>http://genome.ucsc.edu/ENCODE/dataMatrix/</u> <u>encodeDataMatrixMouse.html</u>

- Epigenetics roadmap:
- <u>http://www.ncbi.nlm.nih.gov/geo/roadmap/e</u> <u>pigenomics/</u>
- Yeah, there sure is a lot there. Click on experimental matrix and you can click on samples that take you straight to browser.
- Each one, right-click track, get url, paste into UCSC custom tracks. Some have direct links too.

Human fetal brain from roadmap



- Just remember the labs making these data are real factories. They're pretty good at it but don't have questions in mind, conditions and samples might be suboptimal. They also like cell lines because they can get lots of material.
- Have fun, there is a lot to discover. Good luck!