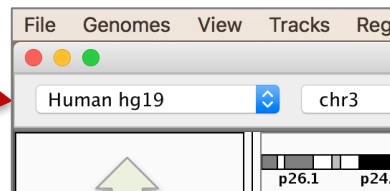


## 1. Launch IGV

## 2. Select reference genome.



- Click on *Human hg19* in the genome drop-down menu in the upper left corner.

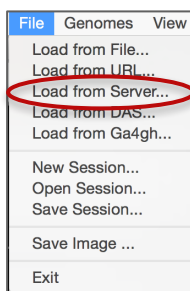
If you only see *Human hg18* in the menu, it's ok to select that instead



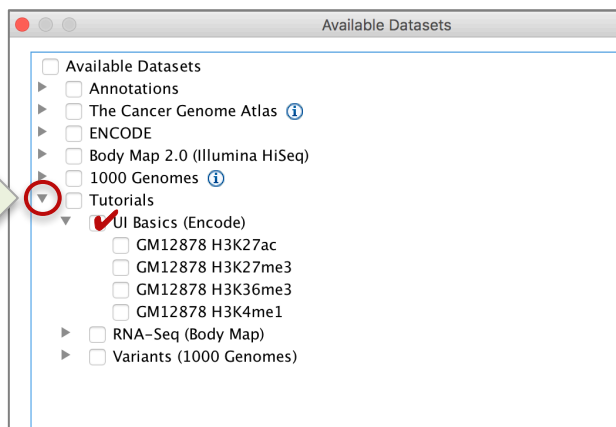
If this is the first time you run IGV, there may be **only one** entry in the menu. More about that later...

## 3. Load data from the IGV hosted server.

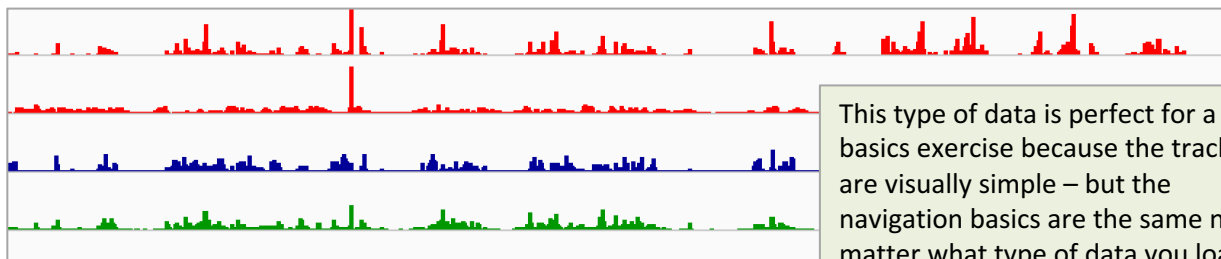
- Select *File > Load from Server...*
- Open the *Tutorials* menu (Use  on Mac, and  on Windows) and click on the *UI Basics* checkbox.



Make sure you only **open** the *Tutorials* menu. Do **not** check the box next to *Tutorials*. That will select everything under *Tutorials*, but we only want *UI Basics* for this exercise.



Four tracks are loaded: ENCODE project ChIP-seq data representing histone modifications. Each track is displayed as a bar chart of signal intensities.

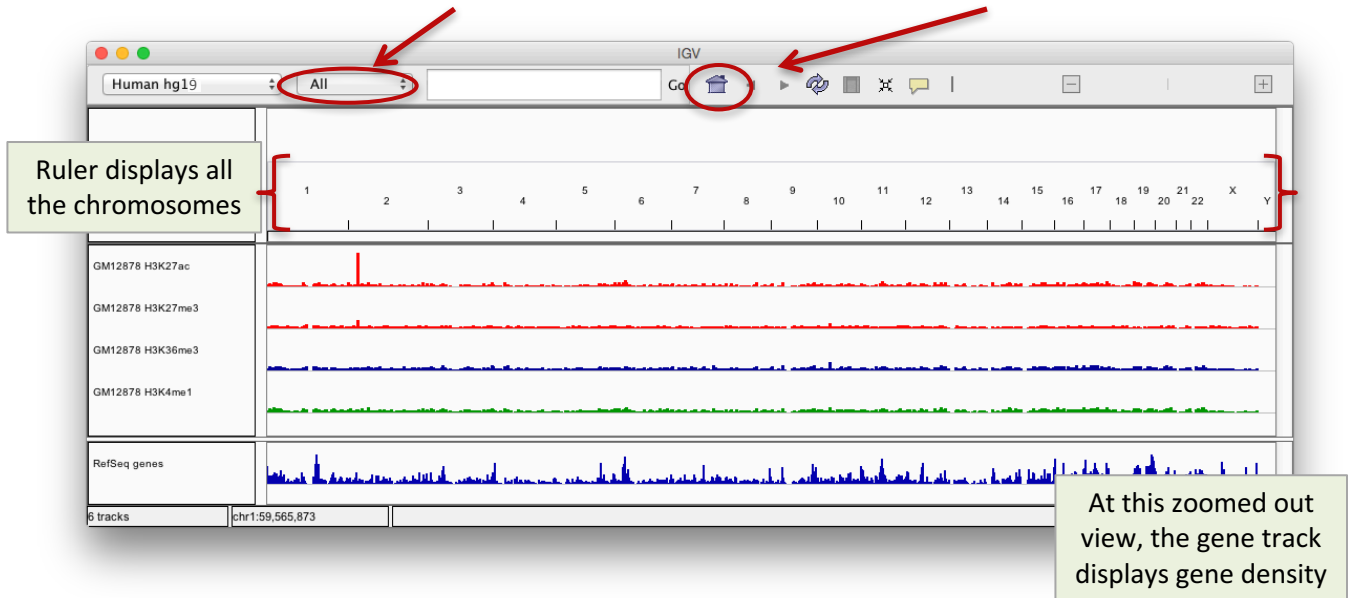


This type of data is perfect for a UI basics exercise because the tracks are visually simple – but the navigation basics are the same no matter what type of data you load.

4. **Navigate** across different genomic loci and at different zoom levels, from whole genome view and down to base-pair resolution.

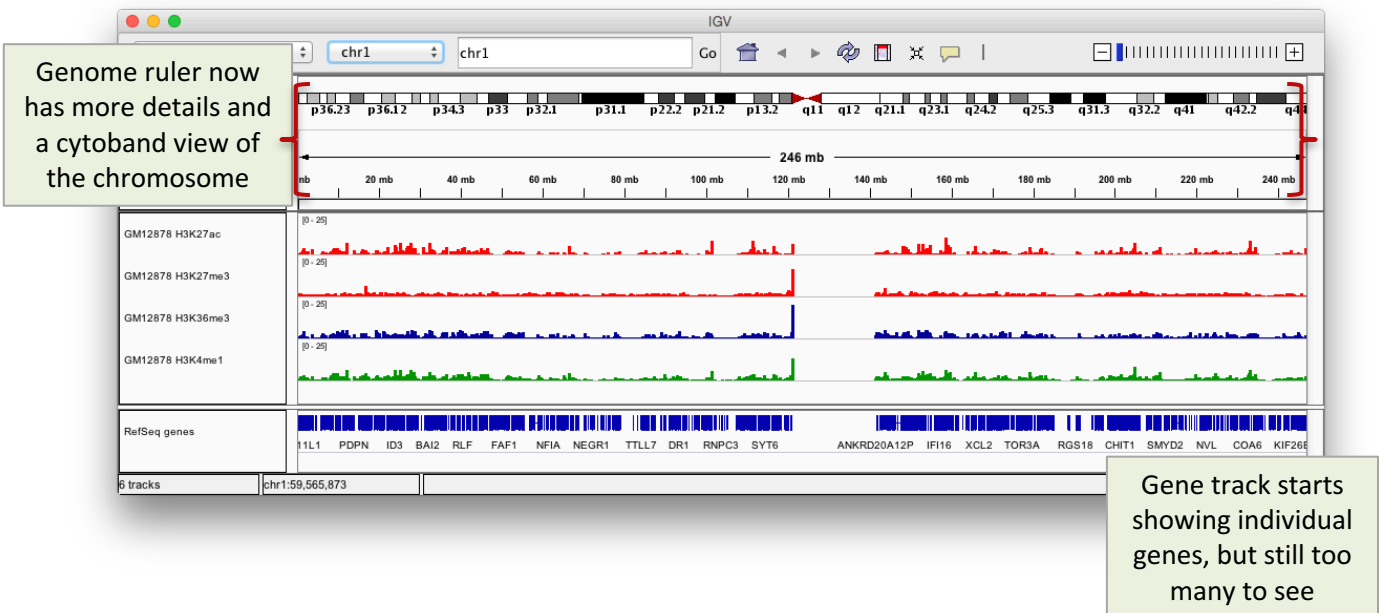
4a. Start at **whole genome view**:

- Select *All* from the chromosome drop-down menu –OR– Click the *Home* button.



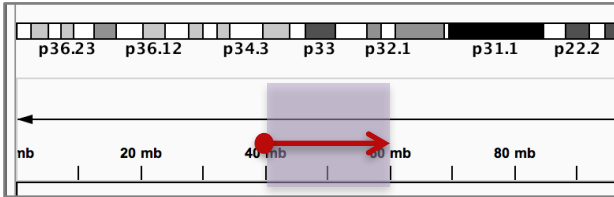
4b. Zoom in to **view one whole chromosome**:

- Select *Chr1* from the chromosome drop-down menu –OR– Click the *1* in the genome ruler.

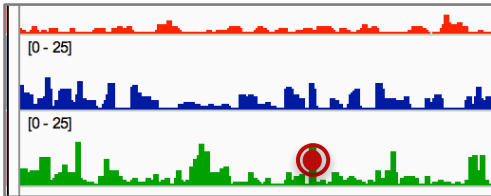


#### 4c. Zoom in further:

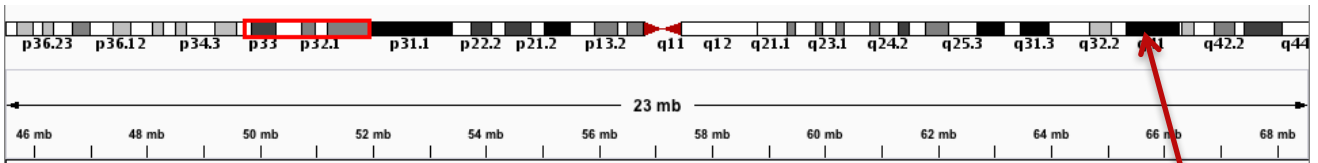
- Click and drag to zoom in on a region swept out in the ruler



- Double-click in the data track to zoom in on a point of interest. [Alt-click to zoom out]

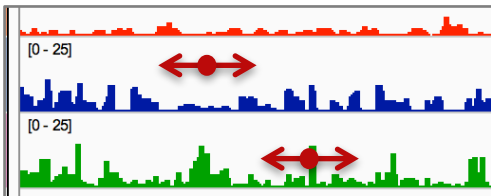


Ruler measurements and a red box on the cytoband diagram show where you are in the chromosome



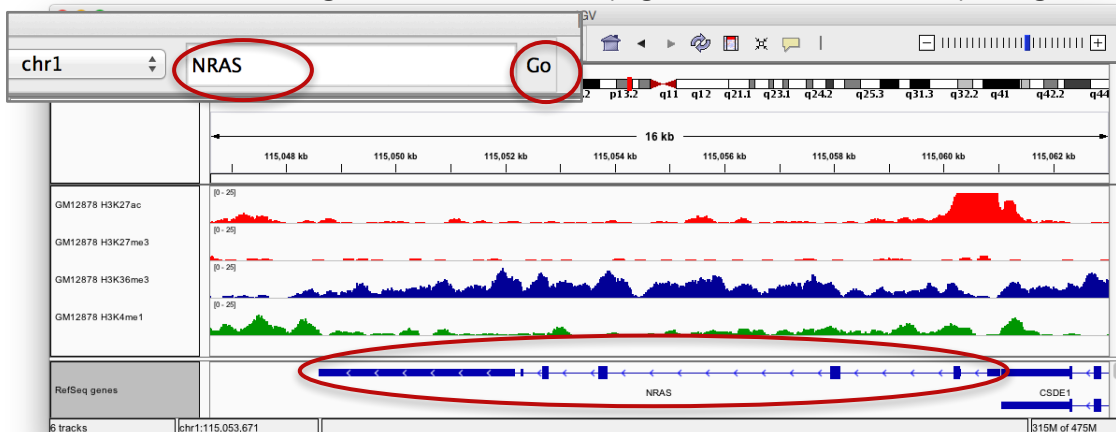
#### 4d. Move around within the chromosome:

- Jump** to another region in the same chromosome (no change in zoom level): Click anywhere in the cytoband diagram.
- Scroll** across genome coordinates: Click anywhere in the data panel and drag left & right.



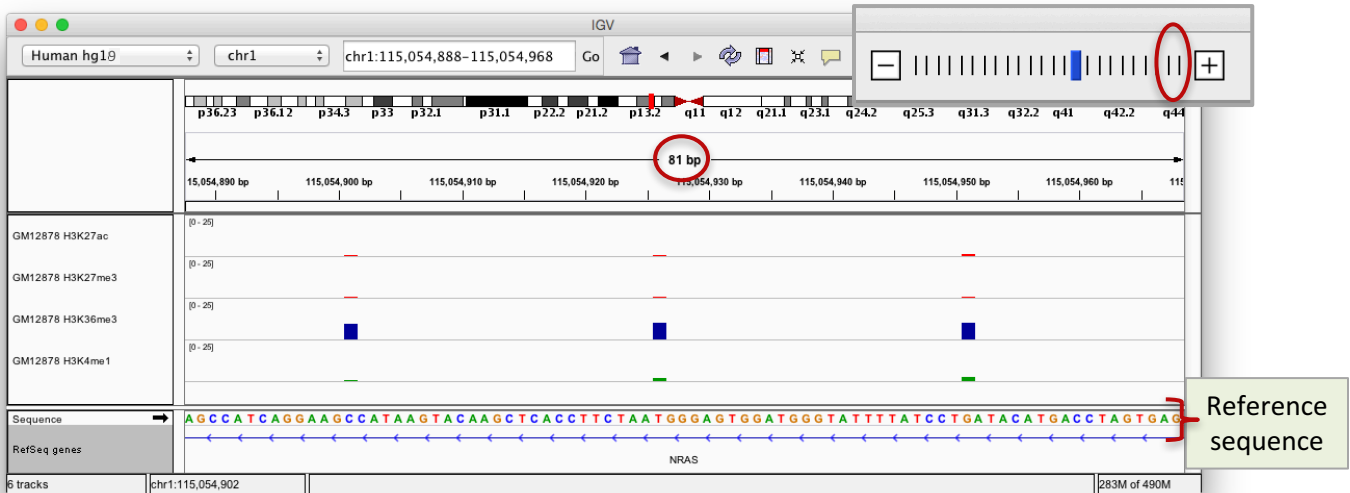
#### 4e. Navigate to specific locus or gene on any chromosome

- Type into the search box in the IGV toolbar and click **Go**:  
either a locus in **genomic coordinates** (e.g. chr1:144,874-969,268) or a **gene name** (e.g. NRAS)



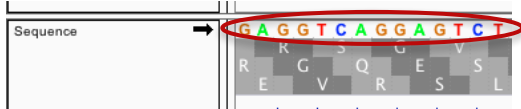
#### 4f. Zoom in to base-pair resolution:

- Keep zooming in as before, or click on one of the rightmost ticks on the “railroad track” zoom widget in the upper right corner.

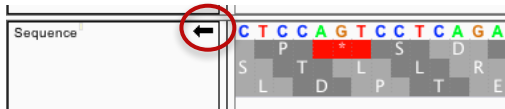


#### 5. Options for viewing the reference sequence track

- Click anywhere on the sequence to show/hide a 3-frame translation

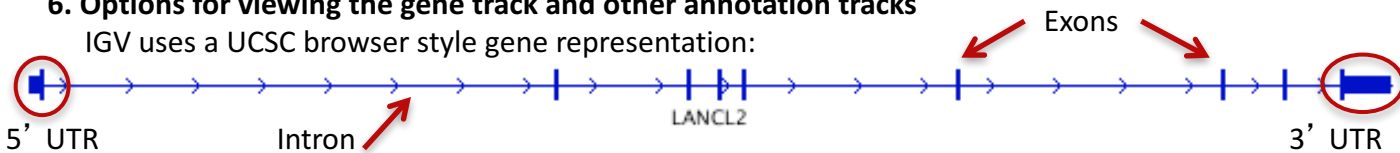


- By default, the sequence for the forward strand is shown. Click on the arrow to reverse the strand.



#### 6. Options for viewing the gene track and other annotation tracks

IGV uses a UCSC browser style gene representation:

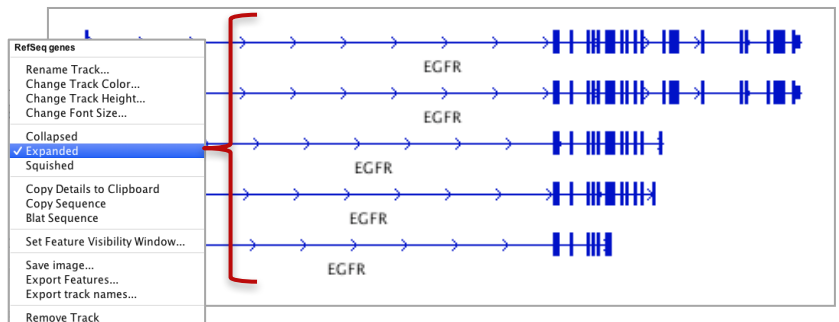


Features are drawn in a single line, by default



- Expand the track using the right-click popup menu

Use *Squished* for an even more compact view



END OF EXERCISE