

Upgraded genome browsers at the Rat Genome Database support comparative and translational studies

<u>Jennifer R Smith</u>¹, Stanley JF Laulederkind¹, G Thomas Hayman¹, Shur-Jen Wang¹, Monika Tutaj¹, Mahima Vedi¹, Wendy M Demos¹, Marek A Tutaj¹, Jyothi Thota¹, Logan Lamers¹, Adam C Gibson¹, Akhilanand Kundurthi¹, Varun Reddy Gollapally¹, Kent C Brodie², Stacy Zacher³, Jeffrey L De Pons¹, Melinda R Dwinell¹, Anne E Kwitek¹

¹Rat Genome Database, Department of Physiology, ²Clinical and Translational Science Institute, ³Finance and Administration, Medical College of Wisconsin, Milwaukee, WI, 53226, USA



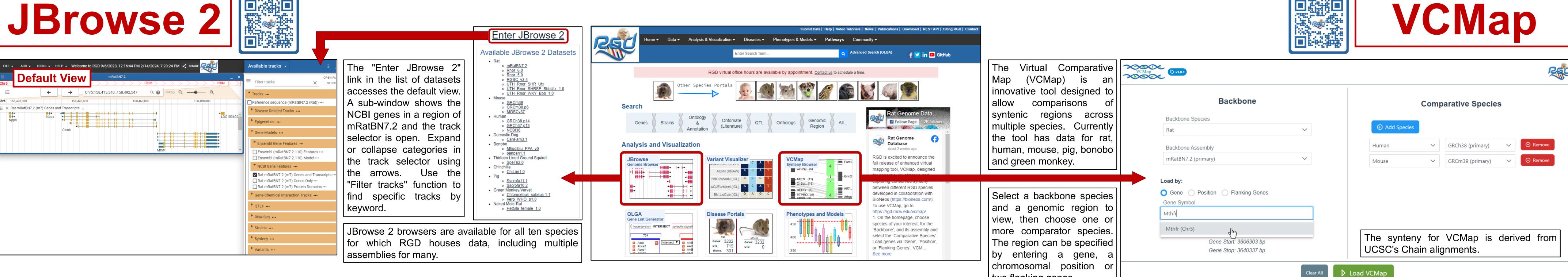
Rat Chr5:158.346.742 - 158.603.505

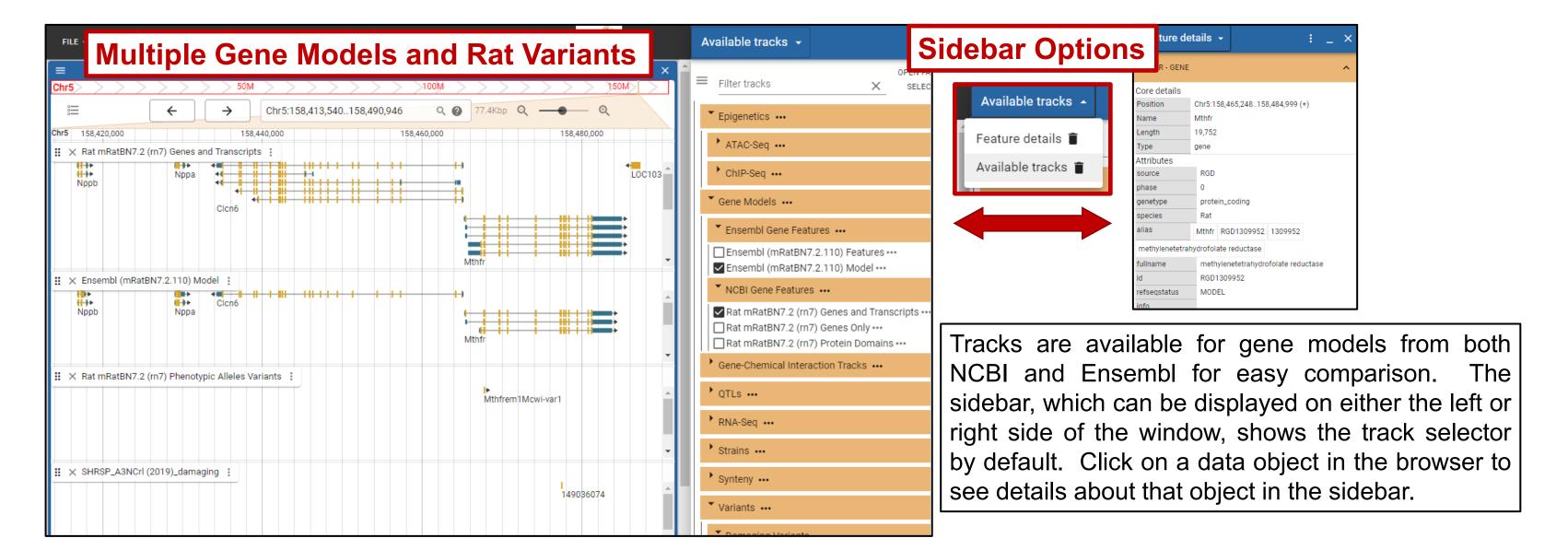
human compared to both rat and mouse, it is shown in the opposite

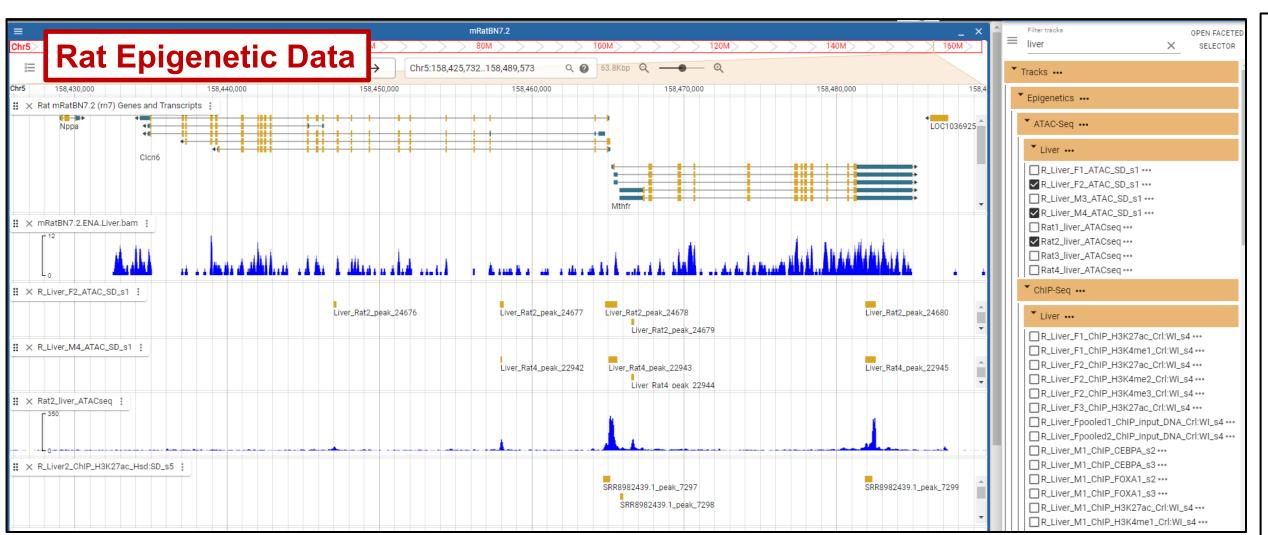
direction as demonstrated by the higher chromosomal position at the top

of the block and the lower number at the bottom in the overview and the









RNA-Seq BAM alignments and epigenetic data from a collaborative project at MCW are incorporated into RGD's JBrowse 2. Tracks include ATAC-Seq peaks for 25 tissues and BAM alignments for liver and ChIP-Seq peaks for three tissues. In some cases, the lists of available tracks can be long, so functionality is provided to filter the tracks on keywords, for example to only give options for "liver", as shown.

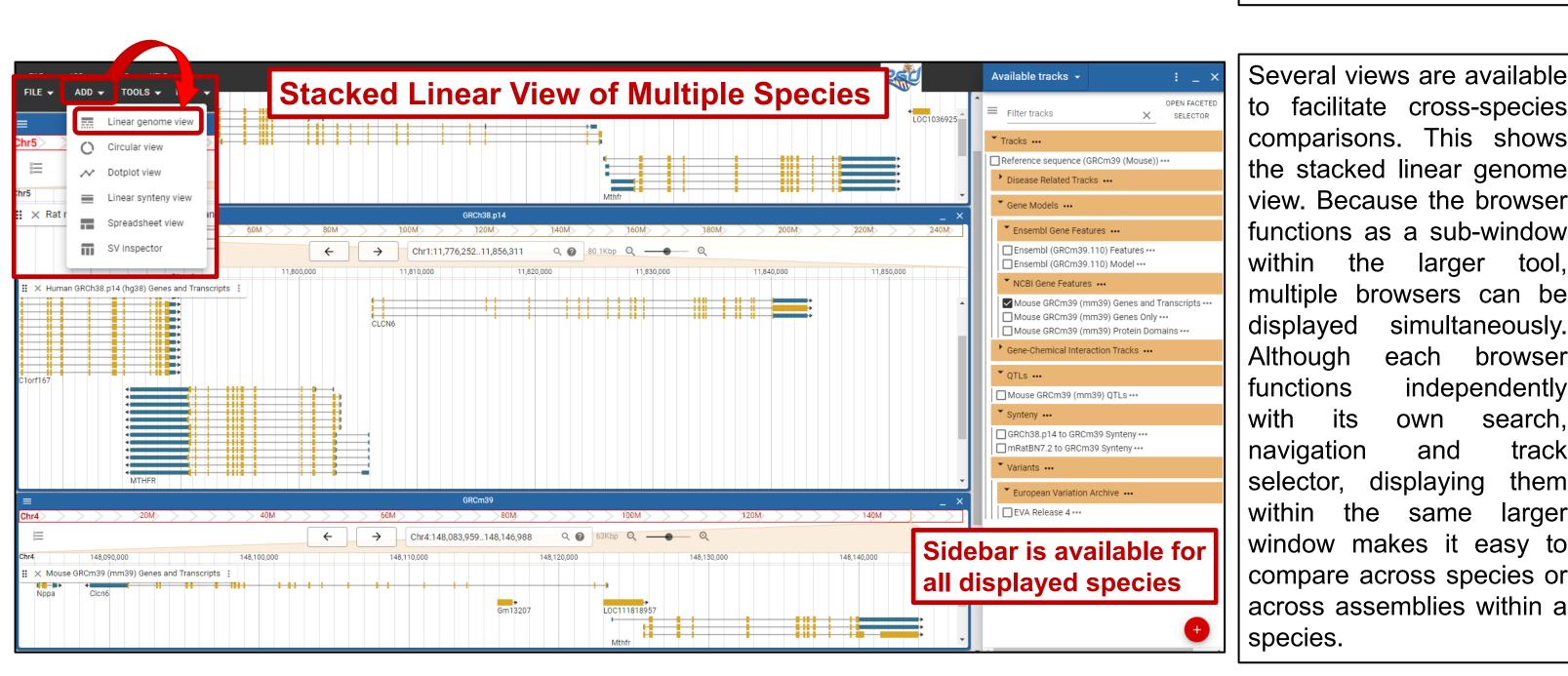
the larger tool,

simultaneously.

each browser

independently

track



Abstract:

The Rat Genome Database (RGD, https://rgd.mcw.edu), a cross-species knowledgebase and the premier online resource for rat genetic and physiologic data, has recently added two advanced comparative genome browsers to our suite of innovative analysis tools—JBrowse 2 and VCMap. JBrowse 2 (https://rgd.mcw.edu/jbrowse2/) is a newly developed genome browser with improved functionality for structural variant and comparative genomics visualization. JBrowse 2 shares many features with its predecessors GBrowse and JBrowse 1, such as the ability to select and view multiple data types in a single view, easy zooming and navigation across a chromosome, and the ability to search for the name or ID of a genome feature, for example a gene or QTL, and go directly to that region of the genome. In addition, it provides expanded functionality designed to facilitate comparative studies, including the linear synteny viewer for crossspecies analyses, the breakpoint split view and circular plot for structural variants, and a dot plot viewer for whole genome alignments.

JBrowse 2 browsers have been set up for all ten RGD species, including browsers for multiple assemblies for each species where those are available. For rat in particular, a substantial set of tracks is available. In addition to gene, QTL and strain tracks, RGD's JBrowse 2 provides tracks for RNA-Seq BAM alignments, strain-specific variants, and ATAC-Seq and ChIP-Seq epigenetics data aligned onto the mRatBN7.2 assembly. Tracks for consolidated variants from the European Variant Archive are available for all applicable species, including rat, mouse, dog, pig and green monkey, allowing comparisons with human variants from ClinVar and the GWAS Catalog for a gene or region of interest.

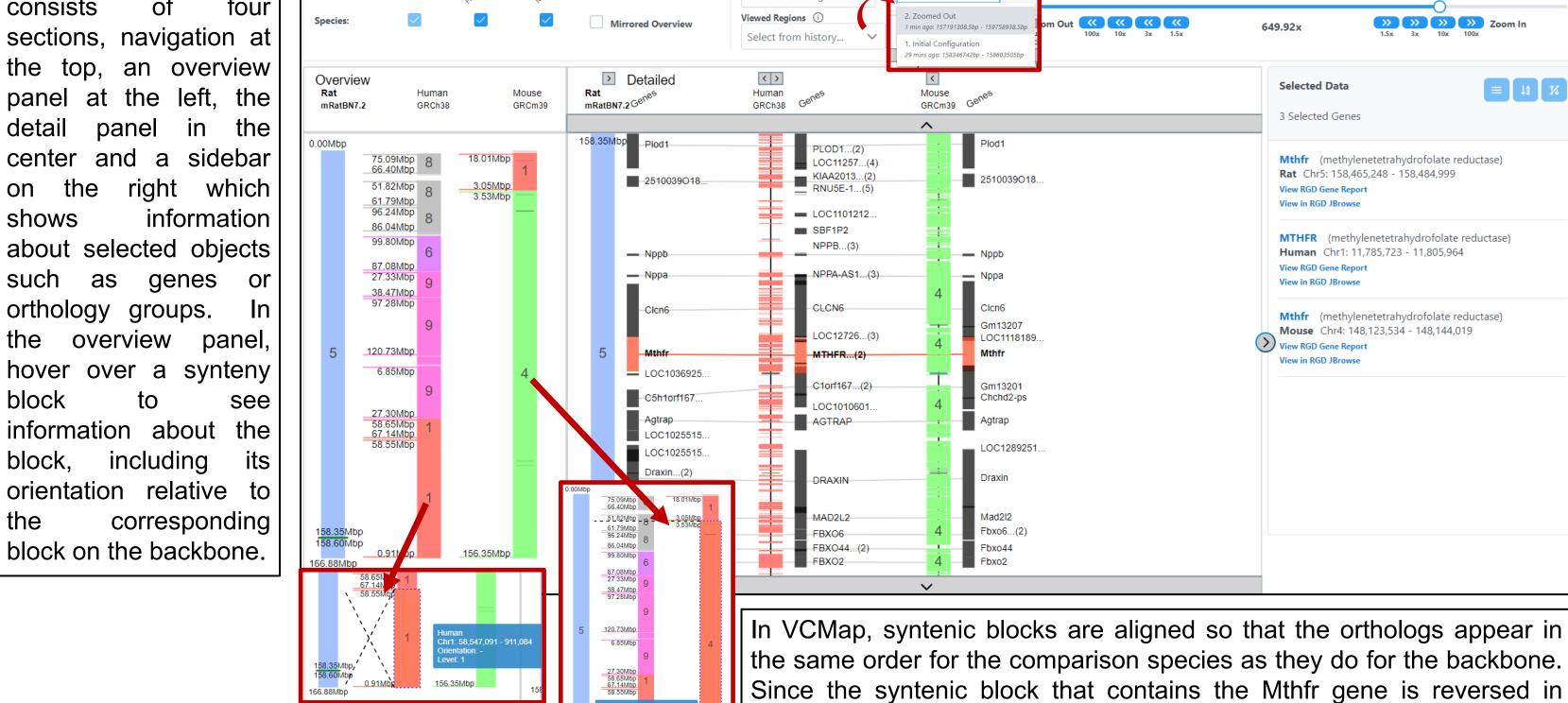
For those interested in synteny across more than two species, RGD is Comparative tool https://rgd.mcw.edu/vcmap/). VCMap, currently released as a beta version, provides a bird's eye view of synteny between two or more species/assemblies, whereas the synteny viewer in JBrowse 2 is limited to only two. Since the original version of VCMap, substantial improvements have been made to the performance and navigation for comparisons of syntenic regions across species. In addition to genes, tracks for data such as genomic variant densities can be added to the display. In conclusion, RGD's JBrowse 2 and VCMap provide valuable functionality for researchers engaging in comparative genomics and translational medicine.



The browser window consists sections, navigation at the top, an overview panel at the left, center and a sidebar right which on the information shows 86.04Mbp about selected objects such as genes or orthology groups. panel, the overview hover over a synteny

two flanking genes.

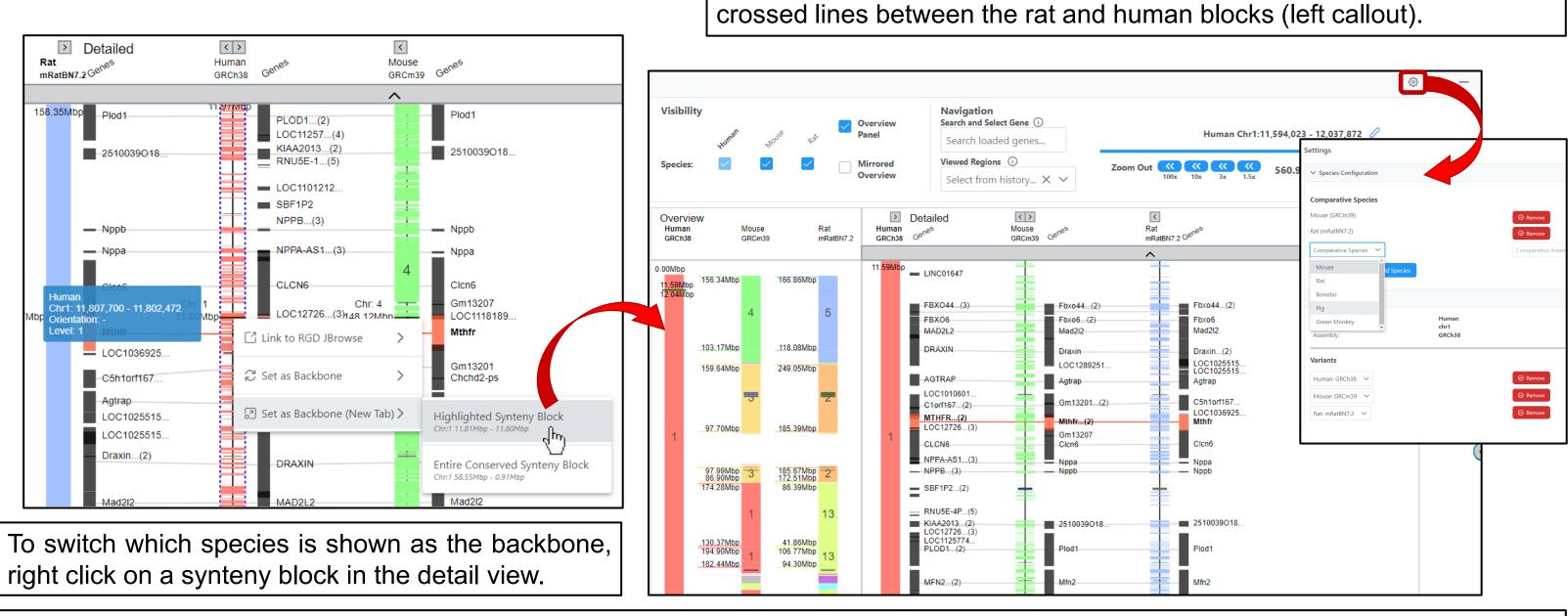
block, including



Show/hide species, change the order of the species, search for a gene within the current view, access previous

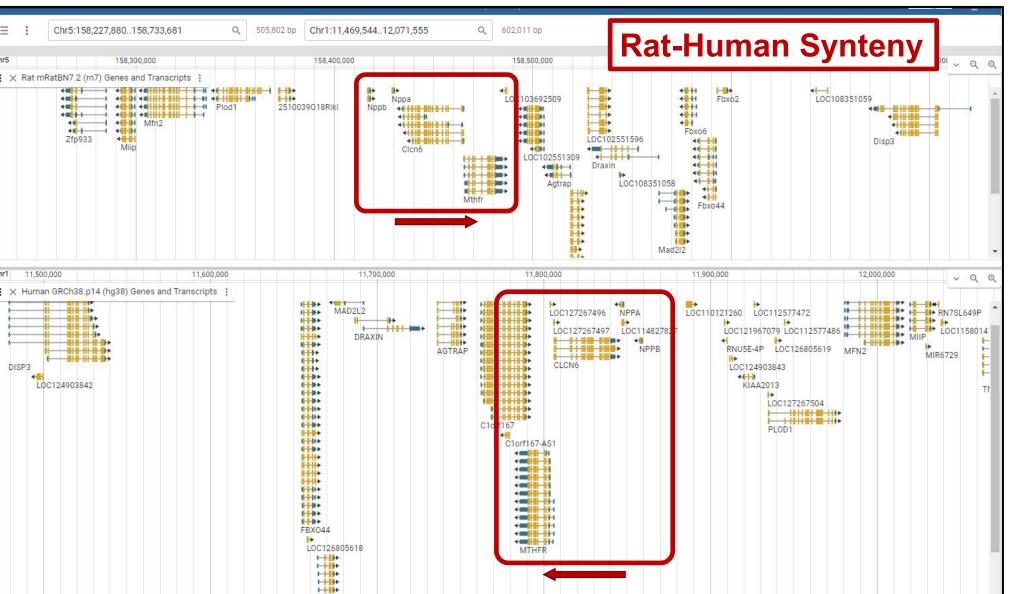
views from the current session, and zoom the detail display using functions in the top section of the page.

Navigation Search and Select Gene (i)

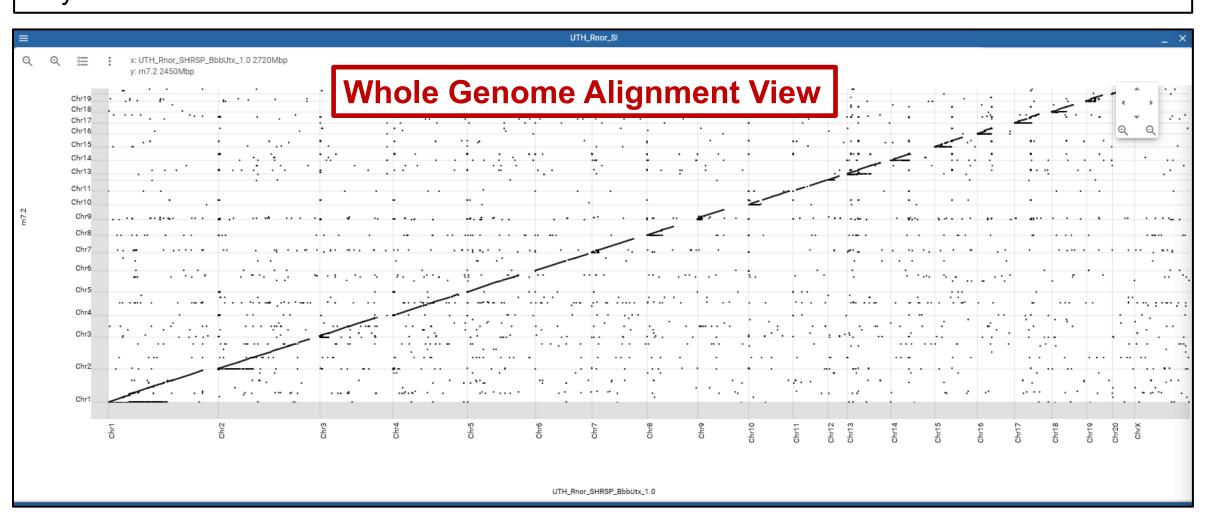


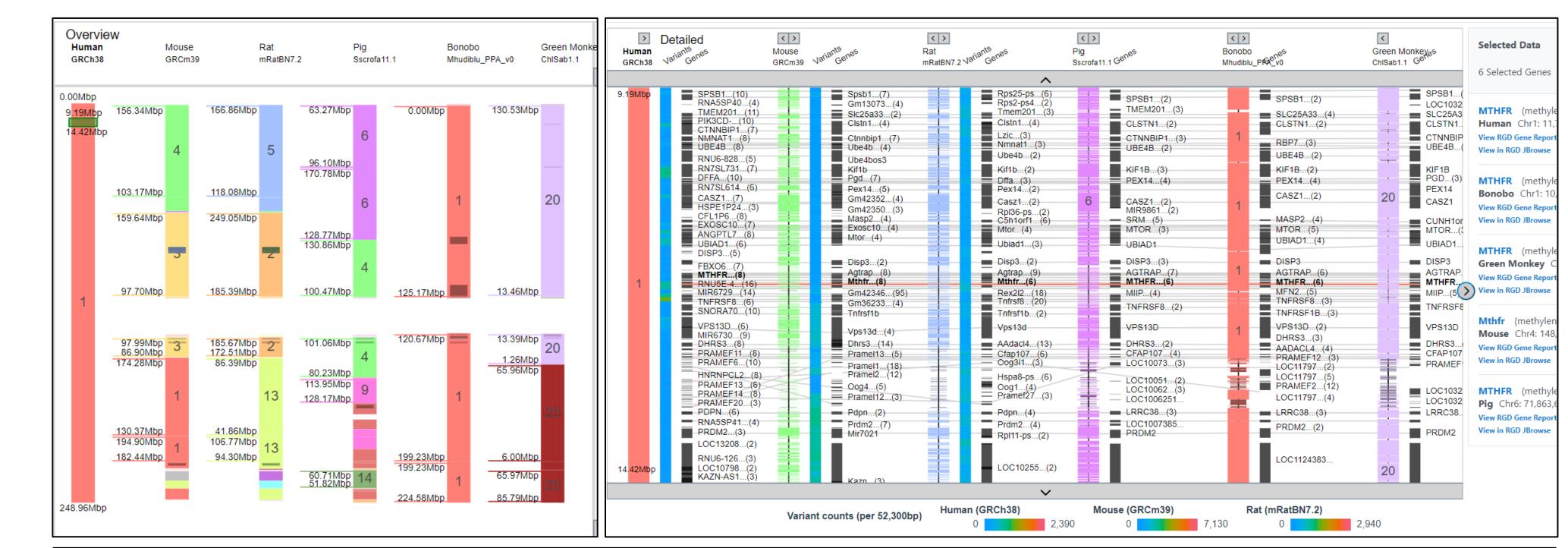
Changing the backbone species from rat to human makes it possible to add additional species to the display based on the synteny data available. As shown in the callout, the "Settings" icon provides access to a popup page where species can be added or removed and additional data tracks, currently variant density tracks, can be added to the display.

The linear synteny view shows syntenic regions in two species or two assemblies. The view below clearly shows that the regions around the Mthfr gene in mRatBN7.2 and human GRCh38 are reversed.



The whole genome alignment view shown below gives an overview of the correspondence between two assemblies. Here the reference assembly mRatBN7.2 is aligned to a de novo assembly of another rat strain, the SHRSP/BbbUtx. The dot plot view demonstrates that there is good correspondence between the two assemblies with no major structural variations. The JBrowse 2 software also provides a breakpoint split viewer and a circular view for visualization of structural variations (not shown). These functions have not yet been implemented at RGD but may be added in the future.





With human as the backbone, all five additional species can be compared showing MTHFR orthologs in all of the species. For human, mouse and rat, heatmaps show variant density. Human variants are imported from ClinVar and the GWAS Catalog and mouse variants are imported from the European Variant Archive. Rat variants are derived from whole genome sequencing of multiple rat strains as part of Dr. Mindy Dwinell's Hybrid Rat Diversity Project at the Medical College of Wisconsin. Mouse over a bin to see the number of variants in that segment.