

Use The UCSC Genome Browser To Visualize And Analyze Your Genomic Data



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Introduction

The UCSC Genome Browser (http://genome.ucsc.edu) is a free, web-based tool that integrates and displays genomic data from a wide variety of sources, including GenBank, ENCODE, UCSC and many others. We provide several tools to help users upload their own data and view it alongside this genomic information or export the data for analysis with other applications. Large genomic data sets and custom genome assemblies can be uploaded and displayed using the browser's data hub tools. If you need to view private data, such as protected patient data from a clinical trial, Genome Browser in a Box (GBiB) allows you to run your own private copy of the Genome Browser on your own computer. The new Data Integrator tool lets you quickly combine input from up to five genome-wide data sets, including your own data uploaded through custom tracks or track hubs, and then export a customized output set based on intersections with a primary track. We are continually working to extend our toolset to allow users to explore their data in new and unique ways.



The UCSC Genome Browser allows you to view your annotations alongside those from a variety of sources, including ENCODE, dbSNP, ClinVar, Ensembl and many others. Here, we can see a combination of custom tracks, including a VCF file of patient SNPs,

Use The Data Integrator To Intersect Multiple Data Sets At Once

The Data integrator allows you to quickly and effective combine data from multiple sources. For instance, you could use it to find the genes, variants and conserved genomic regions that overlap with peaks from a recent RNA-seq experiment. Here's how you can build a query to get that data:

Start by choosing your				
assembly from one of the				
hundreds hosted by UCSC,				
or upload your own with an				
assembly hub.				

	Select Genome Assembly and Region			
	group	genome	assembly	
ノ, n	region to a	nnotate		~

position or search term **\$** chr21:33,031,597-33,041,570

Next, choose a region to query. This can be a single or set of regions,

Load Your Data Into The Genome Browser Using **Data Hubs**

Tracks hubs - package and upload any number of tracks for existing assemblies.

Assembly hubs - upload a custom assembly and related annotations to view in the Genome Browser.

Both support bigBed, bigWig, bigGenePred, BAM, VCF, and HAL formats.

Find more information and examples at: http://genome.ucsc.edu/goldenPath/help/ hgTrackHubHelp.html

JCSC Genome Browser on Human Feb. 2009 (GRCh37/hg19) Assembly chr21:33031597-33041570 9,974 bp. enter position, gene symbol or search te

Use The Genome Browser In A Box To View Your Private Data



GBiB is a virtual machine that runs on your computer. It supports custom tracks and data hubs. Any personal data loaded into GBiB never leaves your computer.

genome-wide, or a position found by search term.



The hgMirror tool allows for easy downloading of genomic data for quicker access.

Runs on Windows, Mac, or Linux. Download is 7GB, which expands to 20GB.

Get it today!

- 1. Download GBiB from Genome Browser Store: https://genome-store.ucsc.edu/
- 2. Download and install VirtualBox: https:// www.virtualbox.org/

3. Unzip GBiB file and load it into VirtualBox

For more information on using GBiB, see: http://genome.ucsc.edu/goldenPath/help/ gbib.html

More information:

Send us a message on our

Find us on:

Lastly, configure your output.

33032152

33032152

chr21 33032015

chr21 33032015

View your output in your web browser or save it to a file, optionally compressed with gzip.

By default, all fields from selected tracks are output. Click "Choose fields" to customize the fields in your output.

myPeak.1

myPeak.1

	Iname	Name of gene
	chrom	Reference sequence chromosome or scaffold
	strand	+ or - for strand
	✓ txStart	Transcription start position
	✓ txEnd	Transcription end position
	✓ cdsStart	Coding region start
	cdsEnd	Coding region end
	exonCount	Number of exons
	exonStarts	Exon start positions
	exonEnds	Exon end positions
	proteinID	UniProt display ID for Known Genes, UniProt accession or RefSeg protein ID for UCSC Genes
		Unique identifier for each (known gene, alignment position) pair
-	geneSymbol	HGNC gene symbol
	5	

My Patient Variant

Set all Clear all

- Set all Clear all chrom An identifier from the reference genome
- The reference position, with the 1st base having position ' pos
- Semi-colon separated list of unique identifiers where available
- 🖌 alt Comma separated list of alternate non-reference alleles called on at least one of the sample
- 🗹 qual Phred-scaled quality score for the assertion made in ALT. i.e. give -10log_10 prob(call in ALT is wrong
- ✓ filter nas passed all filters. Otherwise, a semicolon-separated list of codes for filters that fail
- 🕑 info Additional information encoded as a semicolon-separated series of short keys with optional comma-separated values format If genotype columns are specified in header, a semicolon-separated list of of short keys starting with GT
- If genotype columns are specified in header, a tab-separated set of genotype column values; each value is a colon-separated list of values corresponding to keys ir genotypes the format colum

100 Vert. El

lod=37 351

lod=132 477

- Set all Clear all chrom Reference sequence chromosome or scaffold chromStart Start position in chromosom chromEnd End position in chromosom
- Name of item. name score
 - Score (0-1000)

#ct RNAseqPeaks 3524.chrom ct_RNAseqPeaks_3524.chromStart ct_RNAseqPeaks_3524.chromEnd knownGene.name ct_MyPatientVariants_2846.id ct_MyPatientVariants_2846.ref ct_MyPatientVarian Done phastConsElements100way.score phastConsElements100way.name chr21 33032015 33032152 lod=39 356 myPeak.1 uc002ypa.3 rs142752986 chr21 33032015 lod=49 379 33032152 myPeak.1 rs200447364 33032015 33032152 rs199766524 lod=144 485 chr21 myPeak.1 33032015 33032152 lod=21 295 chr21 myPeak.1 33032015 33032152 chr21 myPeak.1 lod=19 285

For more information on using the Data Integrator, see: http://genome.ucsc.edu/goldenPath/help/hgIntegratorHelp.html View online by scanning the code or going to http://genomewiki.ucsc.edu/index.php/ BioengSymp2015Poster

public mailing list: genome@soe.ucsc.edu	GenomeBrowser
View training info, videos, and user's quides:	SenomeBrowser
http://genome.ucsc.edu/training/	UCSC Genome Browser
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Reference:

The UCSC Genome Browser database: 2015 update. Rosenbloom KR, Armstrong J, Barber GP, Casper J, Clawson H, Diekhans M, Dreszer TR, Fujita PA, Guruvadoo L, Haeussler M, Harte RA, Heitner S, Hickey G, Hinrichs AS, Hubley R, Karolchik D, Learned K, Lee BT, Li CH, Miga KH, Nguyen N, Paten B, Raney BJ, Smit AF, Speir ML, Zweig AS, Haussler D, Kuhn RM, Kent WJ. Nucleic Acids Res. 2015 Jan;43(Database issue):D670-81.

Navigating protected genomics data with UCSC Genome Browser in a Box. Haeussler M, Raney BJ, Hinrichs AS, Clawson H, Zweig AS, Karolchik D, Casper J, Speir ML, Haussler D, Kent WJ. Bioinformatics. 2015 Mar 1;31(5):764-6.

