

Genome Informatics Course: UCSC Genome Browser

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Introduction

main sections:

1. UCSC Genome Browser
2. BLAT
3. Custom tracks, Sessions and Track Hubs
4. Table Browser
5. Other UCSC tools
 - what does it do?
 - How do I use it?
 - What problems does it help me solve?

Genome
Browser

Ebola

Blat

Table
Browser

Gene Sorter

In Silico
PCRGenome
Graphs

Galaxy

VisiGene

Utilities

Downloads

Release
LogCustom
TracksCancer
BrowserMicrobial
Genomes

ENCODE

Neanderthal

Mirrors

Training

Blog

Credits

About the UCSC Genome Bioinformatics Site

Welcome to the UCSC Genome Browser website. This site contains the reference sequence and working draft assemblies for a large collection of genomes. It also provides portals to [ENCODE](#) data at UCSC (2003 to 2012) and to the [Neanderthal](#) project. Download or purchase the Genome Browser source code, or the Genome Browser in a Box ([GBiB](#)) at our [online store](#).

We encourage you to explore these sequences with our tools. The [Genome Browser](#) zooms and scrolls over chromosomes, showing the work of annotators worldwide. The [Gene Sorter](#) shows expression, homology and other information on groups of genes that can be related in many ways. [Blat](#) quickly maps your sequence to the genome. The [Table Browser](#) provides convenient access to the underlying database. [VisiGene](#) lets you browse through a large collection of *in situ* mouse and frog images to examine expression patterns. [Genome Graphs](#) allows you to upload and display genome-wide data sets.

The UCSC Genome Browser is developed and maintained by the Genome Bioinformatics Group, a cross-departmental team within the [UC Santa Cruz Genomics Institute](#) and the Center for Biomolecular Science and Engineering ([CBSE](#)) at the University of California Santa Cruz ([UCSC](#)). If you have feedback or questions concerning the tools or data on this website, feel free to contact us on our [public mailing list](#).

The Genome Browser project team relies on public funding to support our work. Donations are welcome -- we have many more ideas than our funding supports! If you have ideas, drop a comment in our [suggestion box](#).

DONATE NOW

News

News Archives ►

To receive announcements of new genome assembly releases, new software features, updates and training seminars by email, subscribe to the [genome-announce](#) mailing list. Please see our [blog](#) for posts about Genome Browser tools, features, projects and more.

29 October 2014 - Genome Browser in a Box (GBiB)

Sometimes you just want to keep your genomics data to yourself. Have you ever hesitated when uploading your data set into the UCSC Genome Browser? If so, you'll be happy to know that we have created a stand-alone personal version: Genome Browser in a Box (GBiB). If you have sensitive genomics data that you would like to view securely on your own laptop in the context of the UCSC Genome Browser, GBiB is for you.

GBiB is an easy-to-install personal copy of the Genome Browser that comes preloaded with the most popular annotation tracks for human. It is highly configurable — you can access or download other annotation tracks of interest, or view any of the other 90+ organisms featured in the public Genome Browser. GBiB runs inside of Oracle's free VirtualBox virtual machine. It has the same core functionality as the UCSC Genome Browser, but keeps your data private and local to your own computer.

GBiB is free for non-commercial use by non-profit organizations, academic institutions, and for personal use. Commercial use requires purchase of a license with setup fee and annual payment. Download or purchase GBiB in our secure online [store](#).

You can read more about GBiB on our [blog](#), or in the [help doc](#).

20 October 2014 - dbSNP 141 Available for hg19 and hg38

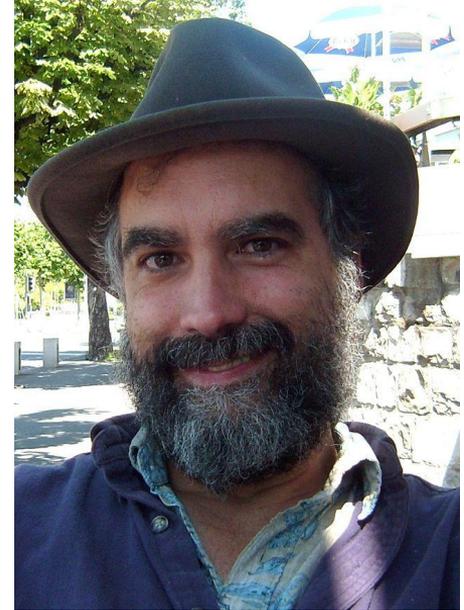
We are pleased to announce the release of four tracks derived from NCBI [dbSNP](#) Build 141 data, available on the two most recent human assemblies GRCh37/hg19 and GRCh38/hg38. The new tracks contain additional annotation data not included in previous dbSNP tracks, with corresponding coloring and filtering options in the Genome Browser.

There are three SNP tracks available for the GRCh37/hg19 assembly. One is a track containing all mappings of reference SNPs to the human assembly, labeled "All SNPs (141)". The other two tracks are subsets of this track and show interesting and easily defined subsets of dbSNP:

- Common SNPs (141): uniquely mapped variants that appear in at least 1% of the population or are 100% non-reference
- Flagged SNPs (141): uniquely mapped variants, excluding Common SNPs, that have been flagged by dbSNP as "clinically associated"

UCSC Genome Bioinformatics

David Haussler



Jim Kent

1. UCSC Browser

- Understanding the browser interface
- Basic searches
- Viewing tracks
- Configuring the display
- Navigating
- Printing images
- Retrieving DNA sequences and annotation

Graphical view of genes, gene structure and annotation

Genome viewer

UCSC Genome Browser on Human Feb. 2009 (GRCh37/hg19) Assembly

move <<< << < > >> >>> zoom in 1.5x 3x 10x base zoom out 1.5x 3x 10x 100x

chr9:21,076,124-21,078,923 2,800 bp. enter position, gene symbol or search terms go

chr9 (p21.3) 21.3 9p24.1 9p23 22q3 9p21.3 21q2 9p21.1 21q3 13.1 9p12 9p11.2 9q12 9q13 21q11 9p21.19 21.31 9p1.33 22.33 9q31.1 23.1 29q31.3 9q32 9q33.1 23.2 9q33.3 34.11 9q34.3

UCSC Genes (RefSeq, GenBank, CCDS, Rfam, tRNAs & Comparative Genomics)

move start < 2.0 > Click on a feature for details. Click or drag in the base position track to zoom in. Click side bars for track options. Drag side bars or labels up or down to reorder tracks. Drag tracks left or right to new position. move end < 2.0 >

track search default tracks default order hide all add custom tracks track hubs configure reverse resize refresh

collapse all Use drop-down controls below and press refresh to alter tracks displayed. expand all

Annotation

- + Mapping and Sequencing refresh
- + Genes and Gene Predictions refresh
- + Phenotype and Literature refresh
- + mRNA and EST refresh
- + Expression refresh
- + Regulation refresh
- + Comparative Genomics refresh
- + Neandertal Assembly and Analysis refresh
- + Denisova Assembly and Analysis refresh
- + Variation refresh
- + Repeats refresh

refresh

Browser Interface

UCSC Genome Browser on Human Feb. 2009 (GRCh37/hg19) Assembly

move <<< << < > >> >>> zoom in 1.5x 3x 10x base zoom out 1.5x 3x 10x 100x

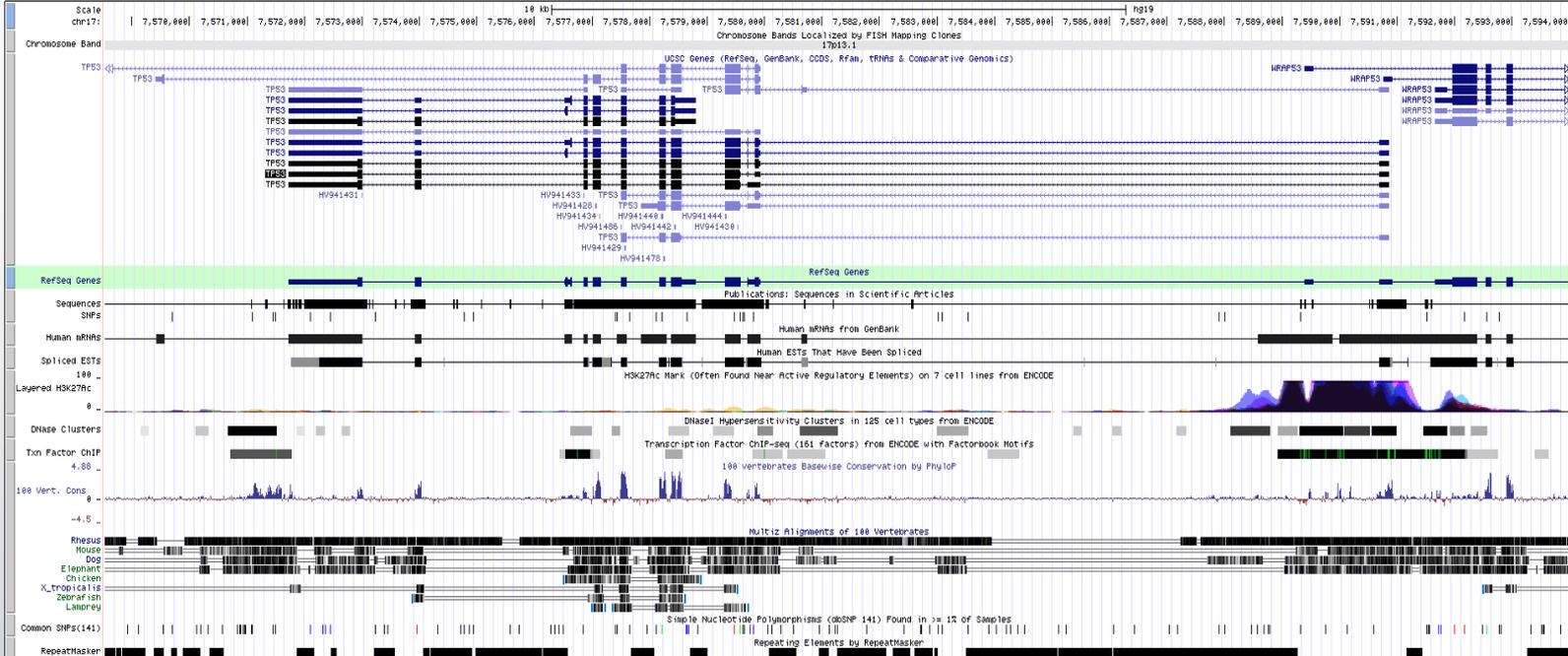
chr17:7568528-7594059 25,532 bp. enter position, gene symbol or search terms

Display Navigation

Search and Configure



chromosome ideogram



Annotation tracks

move start < 2.0 > Click on a feature for details. Click or drag in the base position track to zoom in. Click side bars for track options. Drag side bars or labels up or down to reorder tracks. Drag tracks left or right to new position. move end < 2.0 >

Display Navigation

track search default tracks default order hide all add custom tracks track hubs configure reverse resize refresh

Configuration

Track Configuration

- Track configuration depends on track type and enables you to;
 - Set data thresholds
 - Include or exclude data from a specific source
 - Choose data labels
 - Choose graph type, height, range and scale
- Track and element descriptions contain additional information

Configuring the genome browser display

track search default tracks default order hide all add custom tracks track hubs configure reverse resize refresh

Search **Advanced**

search clear cancel

return to browser (0 of 10 selected)

+ -	Visibility	Track Name	Sort: <input checked="" type="radio"/> by Relevance <input type="radio"/> Alphabetically <input type="radio"/> by Hierarchy
<input type="checkbox"/>	hide ▾	HeLa E2F1 Std	HeLa-S3 E2F1 Standard ChIP-seq Signal from ENCODE/SYDH ▾
<input type="checkbox"/>	hide ▾	HeLa E2F1 Std	HeLa-S3 E2F1 Standard ChIP-seq Peaks from ENCODE/SYDH ▾
<input type="checkbox"/>	hide ▾	MCF-7 E2F1	MCF-7 TFBS Uniform Peaks of HA-E2F1 from ENCODE/USC/Analysis ▾
<input type="checkbox"/>	hide ▾	HeLa-S3 E2F1 c2	HeLa-S3 TFBS Uniform Peaks of HA-E2F1 from ENCODE/USC/Analysis ▾
<input type="checkbox"/>	hide ▾	HeLa-S3 E2F1 c1	HeLa-S3 TFBS Uniform Peaks of E2F1 from ENCODE/USC/Analysis ▾
<input type="checkbox"/>	hide ▾	MCF7 HAE2 UCD	MCF-7 HA-E2F1 UC Davis ChIP-seq Signal from ENCODE/SYDH ▾
<input type="checkbox"/>	hide ▾	MCF7 HAE2 UCD	MCF-7 HA-E2F1 UC Davis ChIP-seq Peaks from ENCODE/SYDH ▾
<input type="checkbox"/>	hide ▾	HeLa HAE2 Std	HeLa-S3 HA-E2F1 Standard ChIP-seq Signal from ENCODE/SYDH ▾
<input type="checkbox"/>	hide ▾	HeLa HAE2 Std	HeLa-S3 HA-E2F1 Standard ChIP-seq Peaks from ENCODE/SYDH ▾
<input type="checkbox"/>	hide ▾ 	SYDH TFBS	Transcription Factor Binding Sites by ChIP-seq from ENCODE/Stanford/Yale/USC/Harvard ▾

Return to Browser (0 of 10 selected)

Search for data types

 Tracks so marked are containers which group related data tracks. Containers may need additional configuration (by clicking on the  icon) before they can be viewed in the browser.

Visual cues

Visual Cues on the Genome Browser



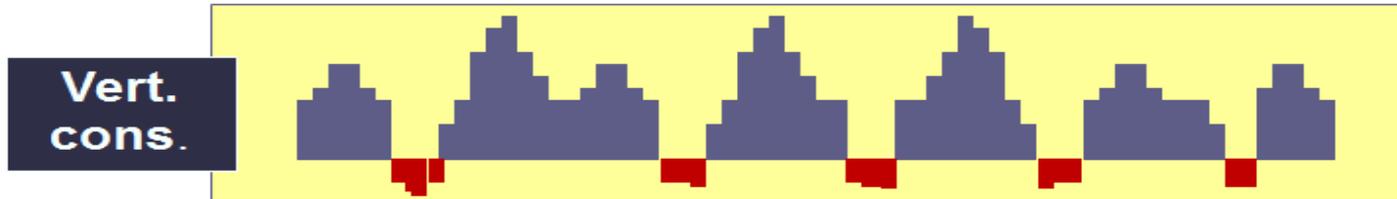
Tick marks; a single location (STS, SNP)



Intron and direction of transcription <<< or >>>

Track colors may have meaning—for example, UCSC Gene track:

- If there is a corresponding PDB entry = black
- If there is a corresponding reviewed/validated seq = dark blue
- If there is a non-RefSeq seq = lightest blue



Vert.
cons.

Wiggle

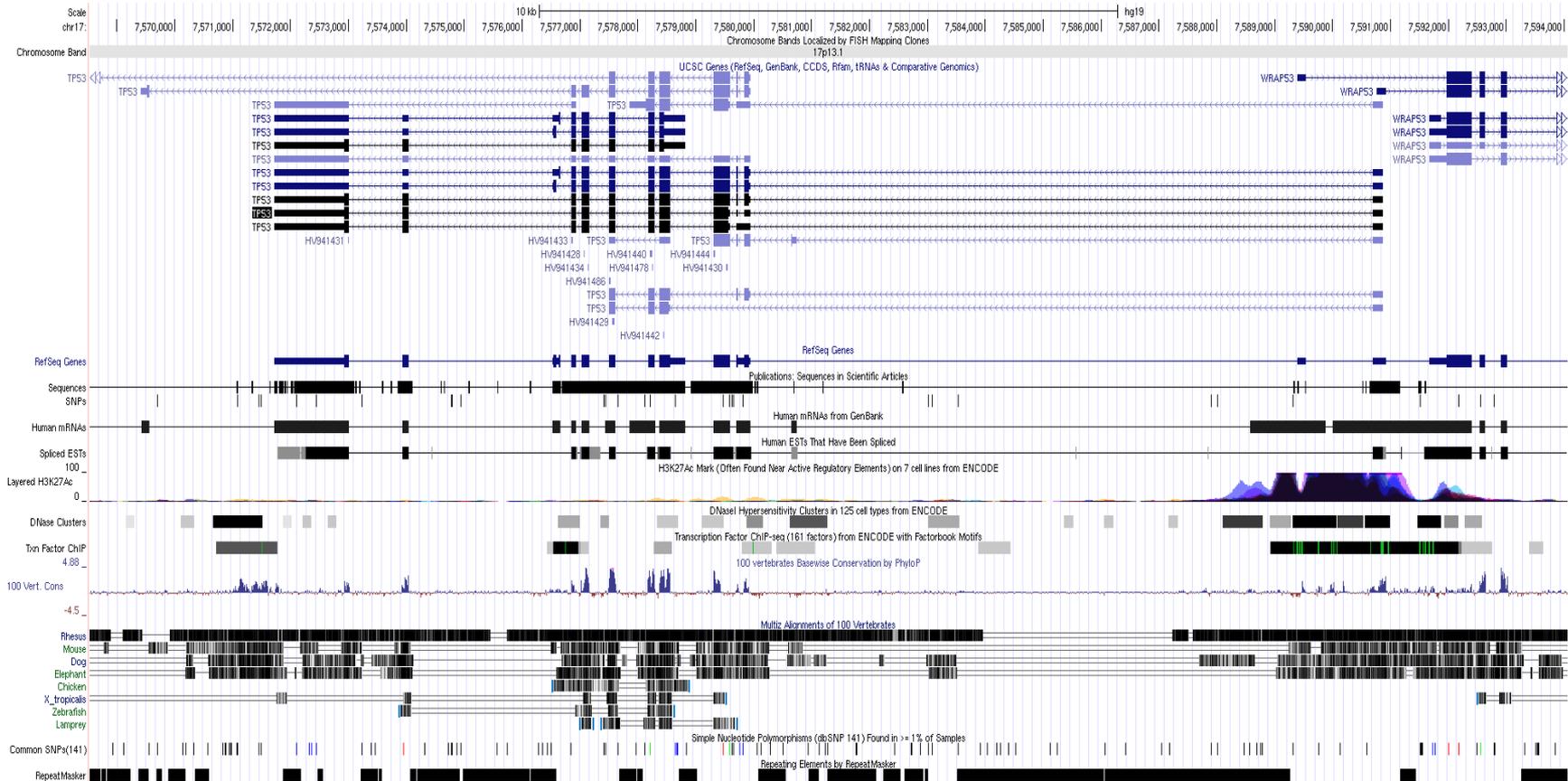
height of a blue bar is increased likelihood of conservation,
red indicates a likelihood of faster-evolving regions

Alignment indications (Conservation pairs: “chain” or “net” style)

- Alignments = boxes, Gaps = lines



Example search for human TP53



Spliced ESTs

hide
dense
dense
squish
pack
full

Annotation Track menu options

Hide: removes a track from view



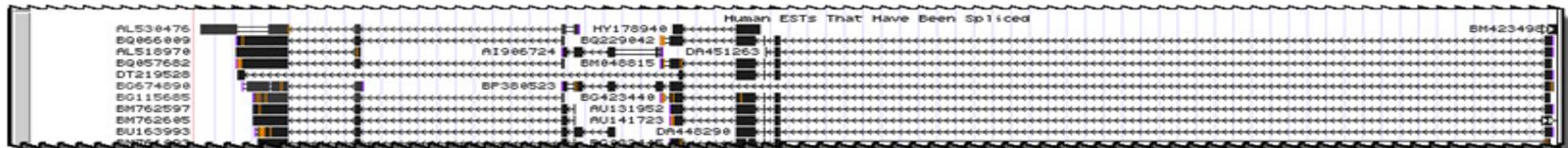
Dense: all items collapsed into a single line



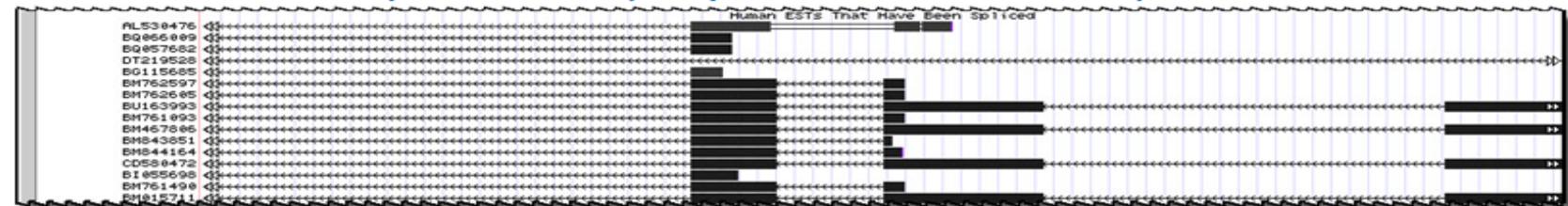
Squish: each item = separate line, but 50% height + packed



Pack: each item separate, but efficiently stacked (full height)



Full: each item on separate line (may need to zoom to fit)



Additional Options: Filters, Supertracks ...

Human ESTs Track Settings

Human ESTs Including Unspliced (All mRNA and EST Tr

Display mode: pack

Filter: red green blue exclude include **Combination Logic:** and or

accession: _____ author: _____ library: _____ **tissue:** neuroblastoma

cell: _____ keyword: _____ gene: _____ product: _____

description: _____

Color track by bases: OFF

Alignment Gap/Insertion Display Options

Draw double horizontal lines when both genome and query have an insertion

Draw a vertical purple line for an insertion at the beginning or end of the query, orange for insertion in the middle of the query

Draw a vertical green line where

Regulation

CpG Islands...

ENC TF Binding...

ENC TF Binding Super-track Settings

Display mode: show

All

dense Uniform TFBS Transcription Factor ChIP-seq Uniform

HAIB TFBS Transcription Factor Binding Sites by C

SYDH TFBS Transcription Factor Binding Sites by C

UChicago TFBS Transcription Factor Binding Sites by F

UTA TFBS Open Chromatin TFBS by ChIP-seq fr

UW CTCF Binding CTCF Binding Sites by ChIP-seq from

Select subtracks by cell line and factor: (help)

Factor	GM12878 (Tier 1)	H1-hESC (Tier 1)	K562 (Tier 1)	HeLa-S3 (Tier 1)	HepG2 (Tier 2)	HUVEC (Tier 2)	IMR90 (Tier 2)	A549 (Tier 2)	MCF-7 (Tier 2)	SK-N-SH (Tier 2)	AG0449	AG0449
ARID3A	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>				
ATF1	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>				
ATF2	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>				
ATF3	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
BACH1	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>				
BATF	<input type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>									
BCL11A	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>				
BCL3	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>				
BCLAF1	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>				

Filter

Supertrack

On Off

- Some tracks have filters (*ESTs shown; SNPs other good example*)
- Super-tracks may have multiple components, various settings
- Some tracks may have un-displayed data

Mid page options to change settings

move start move end

Click on a feature for details. Click or drag in the base position track to zoom in. Click side bars for track options. Drag side bars or labels up or down to reorder tracks. Drag tracks left or right to new position.

track search default tracks default order hide all add custom tracks track hubs configure reverse resize refresh

collapse all expand all

Use drop-down controls below and press refresh to alter tracks displayed. Tracks with lots of items will automatically be displayed.

Mapping and Sequencing

Configure Image

submit

image width: pixels

label area width: characters

text size:

Display chromosome ideogram above main graphic

Show light blue vertical guidelines

Display labels to the left of items in tracks

- Search for data types
- Reset to defaults
- Configure options page
- You control the views with numerous features

Configure Tracks on UCSC Genome Browser: Human Feb. 2009 (GRCh37/hg19)

Tracks: track search hide all show all default Groups: collapse all expand all

Control track and group visibility more selectively below.

Mapping and Sequencing Tracks			
Base Position	dense	-	Chromosome position in bases. (Clicks here zoom in 3x)
Chromosome Band	hide	-	Chromosome Bands Localized by FISH Mapping Clones
STS Markers	hide	-	STS Markers on Genetic (blue) and Radiation Hybrid (black) Maps
FISH Clones	hide	-	Clones Placed on Cytogenetic Map Using FISH
Recomb Rate	hide	-	Recombination Rate from deCODE, Marshfield, or Genethon Maps (deCODE default)
deCODE Recomb	hide	-	deCODE Recombination maps, 10Kb bin size, October 2010

Printing track figures

- Customize track
- Add title
- consider showing only one transcript per gene by turning off splice variants
- Increase the font size and remove the light blue vertical guide lines in the image configuration menu
- Change image size
- Click on blue navigation menu-> view -> **PDF/PS** link

Retrieve DNA sequence

[Genomes](#)[Genome Browser](#)[Tools](#)[Mirrors](#)[Downloads](#)[My Data](#)[Help](#)[About Us](#)

Get DNA in Window (hg19/Human)

Get DNA for

blue navigation menu -> view-> DNA

Position

Note: This page retrieves genomic DNA for a single region. If you would prefer to get DNA for many items in a particular track, or get DNA with formatting options based on gene structure (introns, exons, UTRs, etc.), try using the [Table Browser](#) with the "sequence" output format.

Sequence Retrieval Region Options:

Add extra bases upstream (5') and extra downstream (3')

Note: if a feature is close to the beginning or end of a chromosome and upstream/downstream bases are added, they may be truncated in order to avoid extending past the edge of the chromosome.

Sequence Formatting Options:

- All upper case.
- All lower case.
- Mask repeats: to lower case to N
- Reverse complement (get '-' strand sequence)

Note: The "Mask repeats" option applies only to "get DNA" not to "extended case/color options"

2. BLAT (Blast Like Alignment Tool)

- Rapid sequence search by indexing entire genome
- Useful for finding high similarity matches
- 95% and greater similarity of length 25 bases or more OR sequences of 80% and greater similarity of length 20 amino acids or more
- Limits: DNA (25000 bp), Protein (10000 aa) or 25 sequences
- Can be installed and run locally

Human BLAT Search

BLAT Search Genome

Genome: Assembly: Query type: Sort output: Output type:

BLAT results

Human BLAT Results												
BLAT Search Results												
ACTIONS	QUERY	SCORE	START	END	QSIZE	IDENTITY	CHRO	STRAND	START	END	SPAN	
browser	details	uc002gij.3	2581	1	2591	2591	100.0%	17	-	7571720	7590868	19149
browser	details	uc002gij.3	177	2158	2436	2591	83.1%	1	+	45290354	45290634	281
browser	details	uc002gij.3	176	2134	2433	2591	85.6%	10	-	27408468	27408791	324
browser	details	uc002gij.3	174	2141	2437	2591	83.7%	2	+	27384674	27384975	302
browser	details	uc002gij.3	174	2134	2436	2591	87.6%	10	+	67312526	67312836	311
browser	details	uc002gij.3	173	2148	2431	2591	87.4%	10	+	71133346	71133631	286
browser	details	uc002gij.3	173	2149	2504	2591	84.0%	10	+	65420577	65421000	424
browser	details	uc002gij.3	172	2153	2433	2591	83.4%	3	+	27600067	27600347	281
browser	details	uc002gij.3	165	2160	2444	2591	88.4%	X	-	122127686	122127972	287
browser	details	uc002gij.3	162	2152	2435	2591	83.2%	2	-	109493652	109493934	283
browser	details	uc002gij.3	162	2137	2434	2591	84.0%	1	-	225930110	225930396	287
browser	details	uc002gij.3	162	2144	2437	2591	83.5%	10	+	15559328	15559614	287
browser	details	uc002gij.3	160	2138	2552	2591	82.9%	9	-	131379044	131379531	488
browser	details	uc002gij.3	160	2158	2435	2591	82.2%	4	-	139925816	139926096	281
browser	details	uc002gij.3	160	2134	2414	2591	84.3%	10	-	12095247	12095528	282
browser	details	uc002gij.3	160	2127	2434	2591	86.0%	2	+	170700494	170700797	304
browser	details	uc002gij.3	159	2189	2435	2591	85.4%	16	-	106084386	106084386	806
browser	details	uc002gij.3	26	2128	2154	2591	100.0%	3	-	27607611	27607638	28
browser	details	uc002gij.3	26	2408	2437	2591	93.4%	X	+	47169213	47169242	30
browser	details	uc002gij.3	26	2273	2304	2591	90.7%	5	+	7460469	7460500	32
browser	details	uc002gij.3	25	2358	2389	2591	82.8%	2	+	124842060	124842089	30
browser	details	uc002gij.3	23	2353	2379	2591	92.6%	X	-	100332288	100332314	27
browser	details	uc002gij.3	23	2323	2345	2591	100.0%	X	+	47169722	47169744	23
browser	details	uc002gij.3	22	2369	2404	2591	80.6%	20	-	33243008	33243043	36
browser	details	uc002gij.3	21	2182	2202	2591	100.0%	2	+	38998603	38998623	21
browser	details	uc002gij.3	21	2347	2367	2591	100.0%	1	+	199938363	199938383	21

- Results with demo sequences, settings default; sort = Query, Score
 - **Score is a count of matches—higher number, better match**
- Click [browser](#) to go to Genome Browser image location (next slide)
- Click [details](#) to see the alignment to genomic sequence (2nd slide)

3. Custom tracks, session and track Hubs

Sessions

- **Signing in** enables you to save current settings into a named session, and then restore settings from the session later.
- **lifespan: 4 months**
- **If you wish, you can share named sessions with other users.**
- **Individual sessions may be designated as either *shared* or *non-shared* to protect the privacy of confidential data.**

The screenshot shows the UCSC Genome Bioinformatics website. The navigation bar includes 'Genomes', 'Genome Browser', 'Tools', 'Mirrors', 'Downloads', 'My Data', 'Help', and 'About Us'. The 'My Data' menu is open, showing 'Sessions', 'Track Hubs', and 'Custom Tracks'. Below the navigation bar, there is a 'Sign in to UCSC Genome Bioinformatics' section with links for 'Login' and 'Create an account'. A text block explains that signing in allows saving current settings into a named session and restoring them later. The 'Session Management' section provides a link to the 'Sessions User's Guide' and a link to 'reset' browser settings. It also explains that signing in allows saving named sessions. The 'Save Settings' section has a form to save current settings to a local file, with a 'file' input, a 'file type returned' dropdown set to 'plain text', and a 'submit' button. The 'Restore Settings' section has three forms: one to use settings from another user's saved session (with 'user' and 'session name' inputs and a 'submit' button), one to use settings from a local file (with a 'Choose file' button and a 'submit' button), and one to use settings from a URL (with a text input and a 'submit' button'). The 'Sharing Sessions' section explains that there are several ways to share saved sessions, including saving named sessions for display with Browser and Email links, and sending email with a local file.

Sign in to UCSC Genome Bioinformatics

[Login](#)
[Create an account](#)

Signing in enables you to save current settings into a named session, and then restore settings from the session later. If you wish, you can share named sessions with other users.

Session Management

See the [Sessions User's Guide](#) for more information about this tool.

[Click here to reset](#) the browser user interface settings to their defaults.

If you [sign in](#), you will also have the option to save named sessions.

Save Settings

Save current settings to a local file:

file: file type returned:

(leave file blank to get output in browser window)

Restore Settings

Use settings from another user's saved session:

user: session name:

Use settings from a local file: No file chosen

Use settings from a URL (http://..., ftp://...):

Sharing Sessions

There are several ways to share saved sessions with others.

- If you [sign in](#), you will be able to save named sessions which will be displayed with Browser and Email links.
- If you have saved your settings to a local file, you can send email to others with the file as an

Custom tracks

it is possible for users to upload their own annotation data for temporary display in the browser. These custom annotation tracks are viewable only on the machine from which they were uploaded and are automatically discarded 48 hours after the last time they are accessed, unless they are saved in a [Session](#). Optionally, users can make custom annotations viewable by others as well.

- Format your data
- Define browser characteristics
- Define track characteristics
- Upload and view your track
- Add URL for annotation details (option)

Track Hubs

Home Genomes Blat Tables Gene Sorter PCR Session FAQ Help

Human (*Homo sapiens*) Genome Browser Gateway

The UCSC Genome Browser was created by the [Genome Bioinformatics Group of UC Santa Cruz](#).
Software Copyright (c) The Regents of the University of California. All rights reserved.

clade: Mammal genome: Human assembly: Feb. 2009 (GRCh37/hg19) position or search term: chr21:33,031,597-33,041,570 gene: image width: 1134 submit

Track Data Hubs

Track data hubs are collections of tracks from outside of UCSC that can be imported into the Genome Browser. To import a public hub check the box next to the hub will show up as a group of tracks with its own blue bar and label underneath the main browser graphic, and in the configure page. For more information see the [Track Hubs Guide](#).

NOTE: Because Track Hubs are created and maintained by external sources, UCSC is not responsible for their content.

Public Hubs	My Hubs	Display	Hub Name	Description	Assemblies	URL
<input type="checkbox"/>		<input type="checkbox"/>	SDSU NAT	Sense/antisense gene/exon expression using Affymetrix exon array from South Dakota State University, USA	mm4, mm9, hg19	http://bioinformatics.sdsstate.edu
<input checked="" type="checkbox"/>		<input checked="" type="checkbox"/>	DNA Methylation	DNA Methylation	rheMac3, mm9, hg18, hg19	http://smithlab.usc.edu/trackdata
<input type="checkbox"/>		<input type="checkbox"/>	Translation Initiation Sites (TIS)	Translation Initiation Sites (TIS) track	hg19	http://genagastro.1.med.uni-kl.de/
<input type="checkbox"/>		<input type="checkbox"/>	ENCODE Analysis Hub	ENCODE Integrative Analysis Data Hub	hg19	http://ftp.ebi.ac.uk/pub/databases/ENCODE/
<input type="checkbox"/>		<input type="checkbox"/>	miRcode microRNA sites	Predicted microRNA target sites in GENCODE transcripts	hg19	http://www.mircode.org/ucsc/
<input type="checkbox"/>		<input type="checkbox"/>	Roadmap Epigenomics Data Complete Collection at Wash U VizHub	Roadmap Epigenomics Data Complete Collection at Wash U VizHub	hg19	http://vizhub.wustl.edu/VizHub/

Use Selected Hubs **Load** [@soe.ucsc.edu](#)

UCSC Genome Browser on Human Mar. 2006 (NCBI36/hg18) Assembly

chr17:38,449,840-38,530,994 81,155 bp. enter position, gene symbol or search terms go

chr17 (q21.31) p13.3 p13.2 p13.1 17p12 17p11.2 17q11.2 17q12 17q21 17q22 17q23 17q24 17q25 17q25.3

Scale chr17: 38,460,000| 38,470,000| 38,480,000| 38,490,000| 38,500,000| 38,510,000| 38,520,000| 38,530,000| hg18

CD133HSC
HSPC
HSPC
Neut
Neut

Changes in Human Hematopoietic Stem Cells, Hodges 2011
Human_HSPC_Meth
Human_HSPC_Meth
Human_Neut_Meth

DNA Methylation refresh

Acute Myeloid Leukemia hide	B Cells hide	Blood Cells from Different Ages hide	Brains hide	Breast Cancer hide	Chronic Lymphocytic Leukemia hide
Colon Cancer hide	Colorectal Cancer and Adenomatous Polyp hide	Developing human brain hide	Fetal Lung Fibroblasts hide	Fibroblasts hide	Hematopoietic Stem Cells full
Induced Pluripotent Stem Cells hide	Leukocytes hide	Lymphoblastoid hide	Neuroepithelium Cells hide	Neuronal Cells hide	Peripheral Blood Mononuclear Cells hide
Placenta, kidney, etc hide	Sperm full				

Track Hubs

Table Browser

Use this program to retrieve the data associated with a track in text format, to calculate intersections between tracks, and to retrieve DNA sequence covered by a track. For help in using this application see [Using the Table Browser](#) for a description of the controls in this form, the [User's Guide](#) for general information and sample queries, and the [OpenHelix Table Browser tutorial](#) for a narrated presentation of the software features and usage. For more complex queries, you may want to use [Galaxy](#) or our [public MySQL server](#). To examine the biological function of your set through annotation enrichments, send the data to [GREAT](#). Refer to the [Credits](#) page for the list of contributors and usage restrictions associated with these data. All tables can be downloaded in their entirety from the [Sequence and Annotation Downloads](#) page.

clade: genome: assembly:

group: track: ← **Track Hubs**

table:

region: genome ENCODE Pilot regions position

identifiers (names/accessions):

filter:

intersection with knownGene:

correlation:

output format:

output file:

file type returned: plain text

Note: The all fields and selected fields options are only available when using the custom track output format.

To reset all user cart settings (including custom tracks), click [here](#).

Table Browser

Use this program to retrieve the data associated with a track in text format, to calculate intersections between tracks, and to retrieve DNA sequence covered by a track. For help in using this application see [Using the Table Browser](#) for a description of the controls in this form, the [User's Guide](#) for general information and sample queries, and the [OpenHelix Table Browser tutorial](#) for a narrated presentation of the software features and usage. For more complex queries, you may want to use [Galaxy](#) or our [public MySQL server](#). To examine the biological function of your set through annotation enrichments, send the data to [GREAT](#). Refer to the [Credits](#) page for the list of contributors and usage restrictions associated with these data.

clade: genome: assembly:

group: track:

table:

engine:

filter:

intersect:

output:

output:

output:

file type:

Note: to return more than 100,000 lines, change the filter setting (above). The entire data set may be available for download as a very large file that contains the original data values (not compressed into the wiggle format) -- see the Downloads page.

To reset all user cart settings (including custom tracks), click [here](#).

4. UCSC Table Browser

- Search for genes and annotation
- Setup and filters
- Join tables
- Retrieve sequences
- Intersecting tracks
- Export to external resources

Table browser interface

clade: Mammal ▼ **genome:** Human ▼ **assembly:** Feb. 2009 (GRCh37/hg19) ▼

group: Mapping and Sequencing ▼ **track:** Assembly ▼

table: gold ▼

region: genome ENCODE Pilot regions position

identifiers (names/accessions):

filter:

intersection:

correlation:

output format: all fields from selected table ▼ Send output to [Galaxy](#) [GREAT](#) [GenomeSpace](#)

output file: (leave blank to keep output in browser)

file type returned: plain text gzip compressed

To reset **all** user cart settings (including custom tracks), [click here](#).

Table browser usage

- Retrieve the DNA [sequence data or annotation data](#) underlying Genome Browser tracks for the entire genome, a specified coordinate range, or a set of accessions
- Apply a [filter](#) to set constraints on field values included in the output
- Generate a [custom track](#) and automatically add it to your session so that it can be graphically displayed in the Genome Browser
- Conduct both structured and free-form SQL queries on the data
- Combine queries on multiple tables or custom tracks through an [intersection or union](#) and generate a single set of output data
- Display [basic statistics](#) calculated over a selected data set
- Display the schema for table and list all other tables in the database connected to the table
- Organize the [output data](#) into several different formats for use in other applications, spreadsheets, or databases

Table Browser driven discovery

Task: Search entire genome for “CAG” trinucleotide repeats from USCS tables.

- Choose genome [hg19]
- Choose table [Repeats->Simple Repeats]
- Describe table -find correct data fields
- Choose region [genome]
- Upload locations
- Data summary - approx. 1 million simple repeats

Features of trinucleotide expansion in humans

Disease	Sequence	Location	Parent of origin of expansion	Repeat number (normal)	Repeat number (pre-mutation)	Repeat number (disease)	Somatic instability
<i>Diseases with coding TNRs</i>							
DRPLA	CAG	<i>ATN1</i> (exon 5)	P	6–35	35–48	49–88	Yes
HD	CAG	<i>HTT</i> (exon 1)	P	6–29	29–37	38–180	Yes
OPMD	GCN	<i>PABPN1</i> (exon 1)	P and M	10	12–17	>11	None found in tissue tested (hypothalamus)
SCA1	CAG	<i>ATXN1</i> (exon 8)	P	6–39	40	41–83	Yes
SCA2	CAG	<i>ATXN2</i> (exon 1)	P	<31	31–32	32–200	Unknown
SCA3 (Machado–Joseph disease)	CAG	<i>ATXN3</i> (exon 8)	P	12–40	41–85	52–86	Unknown
SCA6	CAG	<i>CACNA1A</i> (exon 47)	P	<18	19	20–33	None found
SCA7	CAG	<i>ATXN7</i> (exon 3)	P	4–17	28–33	>36 to >460	Yes
SCA17	CAG	<i>TBP</i> (exon 3)	P > M	25–42	43–48	45–66	Yes
SMBA	CAG	<i>AR</i> (exon 1)	P	13–31	32–39	40	None found

McMurray CT. Mechanisms of trinucleotide repeat instability during human development. Nat Rev Genet. 2010 Nov;11(11):786-99.

Table Browser:Filtering

clade: Mammal
genome: Human
group: All Tracks **track:** Com
table: snp141Common
region: genome ENCODE Pilot regions position
identifiers (names/accessions):
filter:
intersection:
correlation:
output format: all fields from selected table
output file: (leave blank)
file type returned: plain text gzip compressed

search for simple repeats in the entire genome with "CAG" sequence and extract data table.

Filter on Fields:

bin	is			
chrom				
chromStart	is			
chromEnd	is			
name		does	match	^
period	is	ignored		0
copyNum	is	ignored		0
consensusSize	is	ignored		0
perMatch	is	ignored		0
perIndel	is	ignored		0
score	is	ignored		0
A	is	ignored		0
C	is	ignored		0
G	is	ignored		0
T	is	ignored		0
entropy	is	ignored		0
sequence		does	match	CAG

AND Free-form query:

item count	80
item bases	3,222 (0.00%)
item total	3,222 (0.00%)
smallest item	25
average item	40
biggest item	93
smallest score	50
average score	67
biggest score	130

Results

Table Browser: Intersections

- Combines the output of two queries into a single set of data based on specific join criteria.
- For example, this can be used to find all SNPs that intersect with RefSeq coding regions. The intersection can be configured to retain the existing alignment structure of the table with a specified amount of overlap, or discard the structure in favor of a simple list of position ranges using a base-pair intersection or union of the two data sets.
- The button functionalities are similar to those of the *filter* option.

Other tools

- Gene sorter
- *In silico* PCR
- VisiGene browser
- Cancer Browser and Encode portal
- Genome graphs
- Other tools:
 - liftOver
 - Dusters
 - Tree maker

Search for related genes

UCSC Human Gene Sorter

genome	Human	assembly	Mar. 2006 (NCBI36/hg18)	search	tp53	Go!		
sort by	Expression (GNF Atlas2)	configure	filter (now off)	display	50	output	sequence	text

About the Gene Sorter

This program displays a sorted table of genes that are related to one another. The relationship can be one of several types, including protein-level homology, similarity of gene expression profiles, or genomic proximity.

To display a gene and its relatives:

1. Select a genome and assembly from the corresponding pull-down menus.
2. Type a word or phrase into the *search* text box to specify which gene should be displayed in the Gene Sorter. Examples of search terms include FOXA2, HOXA9, and MAP kinase.
3. Choose the gene relationship with which you would like to sort the list by selecting an option from the *sort by* pull-down menu.
4. Press the Go! button to display your results.

Following a successful search, the Gene Sorter displays a table containing the specified gene -- highlighted in light green -- and its relatives, each on a separate line. To adjust the number of rows shown, select an option from the *display* pull-down menu.

The default set of table columns -- which can be expanded, reduced, and rearranged via the *configure* button -- shows additional information about the genes. Some of the column data, such as those in the *BLAST E-value* and *%ID* columns, are calculated relative to the highlighted gene. To select a different gene in the list, click on its name. Clicking on a gene's *Genome Position* will open the UCSC Genome Browser to the location of that gene. Similarly, clicking on a gene's *Description* will open a page showing detailed information about the gene.

One of the most powerful features of the Gene Sorter is its filtering capabilities, accessed via the *filter* button. Use the filter to fine-tune the list of displayed genes to a subset based on a selection of detailed and flexible criteria. For example, the filter may be used to select all human genes over-expressed in the cerebellum that have GO-annotated G-protein coupled receptor activity.

The Gene Sorter offers two options for displaying and downloading sequence associated with the genes in the table. Clicking on the *sequence* button will fetch associated protein, mRNA, promoter, or genomic sequence. To dump the table into a simple tab-delimited format suitable for import into a spreadsheet or relational database, click the *text* button.

The UCSC Gene Sorter was designed and implemented by Jim Kent, Fan Hsu, Donna Karolchik, David Haussler, and the UCSC Genome Bioinformatics Group. This work is supported by a grant from the National Human Genome Research Institute and by the Howard Hughes Medical Institute.

Gene Sorter

UCSC Human Gene Sorter

genome assembly search

sort by display output

#	Name	VisiGene	fetal brain	whole brain	amygdala	thymus	bone marrow	PB-CD4+ Tcells	skin	adipocyte	pancreatic islets	heart	lung	kidney	liver	ovary	testis	BLASTP E-Value	Genome Position	Description
1	TP53	n/a																0	chr17 7,522,016	tumor protein p53 isoform a
2	RPS20	n/a																n/a	chr8 57,148,895	ribosomal protein S20
3	H2AFV	n/a																n/a	chr7 44,846,994	H2A histone family, member V isoform 1
4	RPL7A	187765																n/a	chr9 135,206,495	ribosomal protein L7a
5	RPS13	n/a																n/a	chr11 17,054,155	ribosomal protein S13
6	SNRPG	181122																n/a	chr2 70,368,191	small nuclear ribonucleoprotein polypeptide G
7	EIF4A1	176036																n/a	chr17 7,419,687	eukaryotic translation initiation factor 4A
8	ADSL	77625																n/a	chr22 39,082,485	adenylosuccinate lyase isoform a
9	CR601950	n/a																n/a	chr17 72,069,204	Homo sapiens primary hepatoblastoma cDNA, clone:HKMT0728, full insert sequence.
10	UBE2A	182203																n/a	chrX 118,597,467	ubiquitin-conjugating enzyme E2A isoform 1
11	GMPS	176663																n/a	chr3 157,104,616	guanine monophosphate synthetase
12	G3BP1	176455																n/a	chr5 151,148,388	Ras-GTPase-activating protein SH3-domain-binding
13	NUP37	187198																n/a	chr12 101,014,297	nucleoporin 37kDa
14	QARS	180161																n/a	chr3 49,112,772	glutamyl-tRNA synthetase
15	ZNF207	26352																n/a	chr17 27,711,425	zinc finger protein 207 isoform c
16	XRCC5	n/a																n/a	chr2 216,730,812	ATP-dependent DNA helicase II
17	LOC647099	n/a																n/a	chr17 24,073,314	similar to ribosomal protein L23A
18	PABPC4	36799																n/a	chr1 39,807,039	poly A binding protein, cytoplasmic 4 isoform 2
19	RPS18	180521																n/a	chr6 33,350,044	ribosomal protein S18
20	RPS18	n/a																n/a	chr6_cox_hap1 4,622,203	ribosomal protein S18
21	RPS18	n/a																n/a	chr6_gbl_hap2 4,428,251	ribosomal protein S18
22	PSMA5	180067																n/a	chr1 109,758,277	proteasome alpha 5 subunit
23	LOC441743	n/a																n/a	chr16 376,999	Uncharacterized protein ENSP00000332117.
24	PHF10	27218																n/a	chr6 169,855,917	PHD finger protein 10 isoform a
25	RPS27	59894																n/a	chr1 152,230,551	ribosomal protein S27

Configure

Configure Gene Sorter

Columns:
 Settings:
 Expression ratio colors:
 Show all splicing variants:

Name	On	Position	Description	Configuration
#	<input checked="" type="checkbox"/>	▼	Item Number in Displayed List/Select Gene	n/a
Name	<input checked="" type="checkbox"/>	▲▲▼	Gene Name/Select Gene	n/a
UniProtKB	<input type="checkbox"/>	▲▲▼	UniProtKB Protein Display ID	n/a
UniProtKB Acc	<input type="checkbox"/>	▲▲▼	UniProtKB Protein Accession	n/a
RefSeq	<input type="checkbox"/>	▲▲▼	NCBI RefSeq Gene Accession	n/a
Entrez Gene	<input type="checkbox"/>	▲▲▼	NCBI Entrez Gene/LocusLink ID	n/a
UCSC ID	<input type="checkbox"/>	▲▲▼	UCSC Transcript ID	n/a
GenBank	<input type="checkbox"/>	▲▲▼	GenBank mRNA Accession	n/a
Ensembl	<input type="checkbox"/>	▲▲▼	Ensembl Transcript ID	n/a
KEGG	<input type="checkbox"/>	▲▲▼	KEGG Pathway ID	n/a
GNF Atlas 2 ID	<input type="checkbox"/>	▲▲▼	ID of Associated GNF Atlas 2 Expression Data	n/a
Gene Category	<input type="checkbox"/>	▲▲▼	High Level Gene Category - Coding, Antisense, etc.	n/a
CDS Score	<input type="checkbox"/>	▲▲▼	Coding potential score from tCdsPredict	n/a
VisiGene	<input checked="" type="checkbox"/>	▲▲▼	UCSC VisiGene In Situ Image Browser	n/a
Allen Brain	<input type="checkbox"/>	▲▲▼	Allen Brain Atlas In Situ Images of Adult Mouse Brains	n/a
U133 ID	<input type="checkbox"/>	▲▲▼	ID of Associated Affymetrix U133 Expression Data	n/a
U133Plus2 ID	<input type="checkbox"/>	▲▲▼	ID of Associated Affymetrix U133 Plus 2.0 Expression Data	n/a
U95 ID	<input type="checkbox"/>	▲▲▼	ID of Associated Affymetrix U95 Expression Data	n/a
GNF Atlas 2	<input checked="" type="checkbox"/>	▲▲▼	GNF Expression Atlas 2 Data from U133A and GNF1H Chips	brightness: <input type="text" value="1.0"/> tissues: <input type="button" value="selected"/> values: <input type="button" value="ratio"/>
H-Inv	<input type="checkbox"/>	▲▲▼	H-Invitational Gene Database	n/a
Max GNF Atlas 2	<input type="checkbox"/>	▲▲▼	Maximum Expression Value of GNF Expression Atlas 2	n/a
GNF Atlas 2 Delta	<input type="checkbox"/>	▲▲▼	Normalized Difference in GNF Expression Atlas 2 from Selected Gene	n/a
GNF U95	<input type="checkbox"/>	▲▲▼	GNF Expression Atlas 1 Human Data on Affy U95 Chips	brightness: <input type="text" value="1.0"/> tissues: <input type="button" value="selected"/> values: <input type="button" value="ratio"/>
Max GNF U95	<input type="checkbox"/>	▲▲▼	Maximum Expression Value of GNF Expression Atlas 1	n/a
GNF Atlas1 Delta	<input type="checkbox"/>	▲▲▼	Normalized Difference in GNF Atlas 1 Expression from Selected Gene	n/a
Affy Exons	<input type="checkbox"/>	▲▲▼	Affymetrix All Exon Microarrays	brightness: <input type="text" value="1.0"/>
Affy Exon Dst	<input type="checkbox"/>	▲▲▼	Affymetrix All Exon Microarrays Distance	n/a
BLASTP Bits	<input type="checkbox"/>	▲▲▼	NCBI BLASTP Bit Score	n/a
BLASTP	<input type="checkbox"/>	▲▲▼	NCBI BLASTP E-Value	n/a

In silico PCR

UCSC In-Silico PCR

Genome: Human	Assembly: Mar. 2006 (NCBI36/hg18)	Target: genome assembly	Forward Primer: <input type="text"/>	Reverse Primer: <input type="text"/>	<input type="submit" value="submit"/>
Max Product Size: <input type="text" value="4000"/>	Min Perfect Match: <input type="text" value="15"/>	Min Good Match: <input type="text" value="15"/>	Flip Reverse Primer: <input type="checkbox"/>		

About In-Silico PCR

In-Silico PCR searches a sequence database with a pair of PCR primers, using an indexing strategy for fast performance.

Configuration Options

Genome and Assembly - The sequence database to search.

Target - If available, choose to query transcribed sequences.

Forward Primer - Must be at least 15 bases in length.

Reverse Primer - On the opposite strand from the forward primer. Minimum length of 15 bases.

Max Product Size - Maximum size of amplified region.

Min Perfect Match - Number of bases that match exactly on 3' end of primers. Minimum match size is 15.

Min Good Match - Number of bases on 3' end of primers where at least 2 out of 3 bases match.

Flip Reverse Primer - Invert the sequence order of the reverse primer and complement it.

Output

When successful, the search returns a sequence output file in fasta format containing all sequence in the database that lie between and include the primer pair. The fasta header describes the region in the database and the primers. The fasta body is capitalized in areas where the primer sequence matches the database sequence and in lower-case elsewhere. Here is an example from human:

```
>chr22:31000551+31001000 TAACAGATTGATGATGCATGAAATGGG CCCATGAGTGGCTCCTAAAGCAGCTCG
TtACAGATTGATGATGCATGAAATGGGgggtggccaggggtgggggggtga
gactgcagagaaagcagggctggttcataacaagcttggcgctcccaa
tatgcagctgaagtttccaggggctgattgagccagtgagggtaag
tacacagaacatcctagagaaccctcttccctaaagattaaaaataaa
gactgtgctgtaagggattggatttccatttgagaattctgtta
tccagaaaggcttaccaccacaatgctgaaaagtgtgtaccgtaactcaa
agcaagctcctccagacagagaaacaccagcctcacaggaagcaaa
aaattggcttcacttttaagtgaaatccagaaccagatgtcagagctcc
aagcacttgcctcagctccacGCAGCTGCTTTAGGAGCCACTCATGg
```

The + between the coordinates in the fasta header indicates this is on the positive strand.

Author

In-Silico PCR was written by [Jim Kent](#). Interactive use on this web server is free to all. Sources and executables to run batch jobs on your own server are available free for academic, personal, and non-profit purposes. Non-exclusive commercial licenses are also available. Contact Jim for details.

In silico PCR usage

- Select genome
- Genomic or transcript?
- Enter primers
- Set configuration options

About In-Silico PCR

In-Silico PCR searches a sequence database with a pair of PCR primers, using an indexing strategy for fast performance.

Configuration Options

Genome and Assembly - The sequence database to search.

Target - If available, choose to query transcribed sequences.

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```
-chr22:31000551+31001000 TAACAGATTGATGATGCATGAAATGGG CCCATGAGTGGCTCCTAAAGCAGCTGC
TtACAGATTGATGATGCATGAAATGGGggtgcccaggggtggggggtga
gactgcagagaaagccagggctgggttcataacaagctttgtgctcccaa
tatgacgctgaagtttccaggggctgatggtgagccagtgagggtaa
tacacagaacatcctagagaaaccctatccttaagattaaaaataaa
gacttgcctgctgtaagggattggattatcctatttgagaattctgtta
tccagaatggcttaccaccaaatgctgaaaagtgtaccgtaactcaaa
agcaagctcctcctcagacagagaaaaccagcgcacagaaagcaaa
aaattggcttcaacttttaaggtgaatccagaaccagatgacagagctcc
aagcaacttgcctcaagctccacGAGCTGCTTTAGGAGCCACTCATGg
```

The + between the coordinates in the fasta header indicates this is on the positive strand.

Visigene

VisiGene Image Browser

VisiGene is a virtual microscope for viewing *in situ* images. These images show where a gene is used in an organism, sometimes down to cellular resolution. With VisiGene users can retrieve images that meet specific search criteria, then interactively zoom and scroll across the collection.

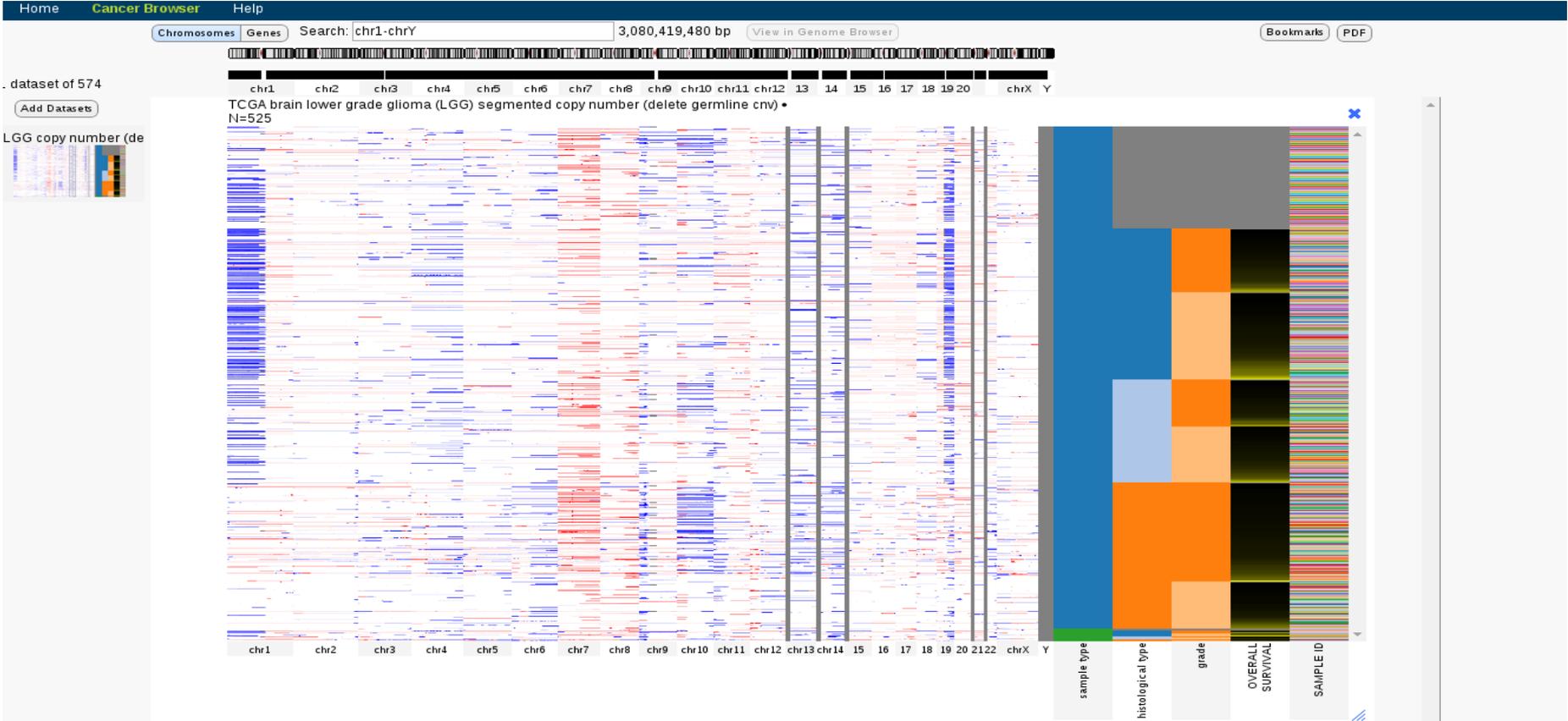
Images Available

The following image collections are currently available for browsing:

- High-quality high-resolution images of eight-week-old male mouse sagittal brain slices with reverse-complemented mRNA hybridization probes from the [Allen Brain Atlas](#), courtesy of the [Allen Institute for Brain Science](#)
- Mouse *in situ* images from the [Jackson Lab Gene Expression Database](#) (GXD) at MGI
- Transcription factors in mouse embryos from the Mahoney Center for Neuro-Oncology
- Mouse head and brain *in situ* images from NCBI's [Gene Expression Nervous System Atlas](#) (GENSAT) database
- *Xenopus laevis in situ* images from the [National Institute for Basic Biology](#) (NIBB) XDB project



Cancer Browser



Encode



Encyclopedia of DNA Elements at UCSC 2003 - 2012

Human Data at UCSC

Downloads

Experiment Matrix

Search

Genome Browser (hg19)

Experiment List

Cell Types

Mouse Data at UCSC

Downloads

Experiment Matrix

Search

Genome Browser (mm9)

Experiment List

Cell Types

Metadata Terms

Registered Variables

Antibodies

Other Resources

News Archive

First Production (2007-2012)

Pilot (2003-2007)

Contacts

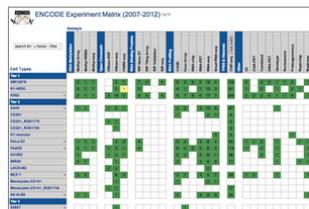
About

The [Encyclopedia of DNA Elements](#) (ENCODE) Consortium is an international collaboration of research groups funded by the National Human Genome Research Institute ([NHGRI](#)). The goal of ENCODE is to build a comprehensive parts list of functional elements in the human genome, including elements that act at the protein and RNA levels, and regulatory elements that control cells and circumstances in which a gene is active.

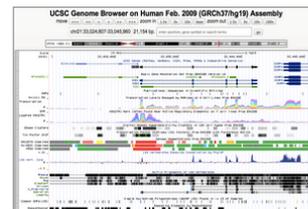
UCSC coordinated data for the ENCODE Consortium from its inception in 2003 (Pilot phase) to the end of the first 5 year phase of whole-genome data production in 2012. All data produced by ENCODE investigators and the results of ENCODE analysis projects from this period are hosted in the UCSC Genome browser and database. Explore ENCODE data using the image links below or via the left menu bar. **All ENCODE data at UCSC are freely available for download and analysis.**

ENCODE results from 2013 and later are available from the ENCODE Project Portal, [encodeproject.org](#). The ENCODE Project Portal also hosts ENCODE data from the first production phase, additional ENCODE access tools, and ENCODE project pages including up-to-date information about data releases, publications, and upcoming tutorials.

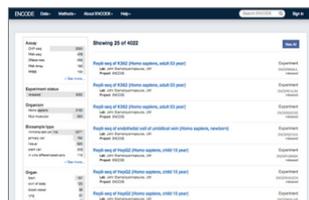
Explore ENCODE data at UCSC



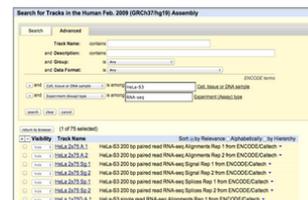
View ENCODE data in the UCSC Genome Browser



Search for data at the ENCODE Portal



Search for ENCODE tracks in the UCSC Browser



Other utilities

UCSC Genome Bioinformatics

[Home](#) · [Genomes](#) · [Blat](#) · [Tables](#) · [Gene Sorter](#) · [PCR](#) · [Session](#) · [FAQ](#) · [Help](#)

UCSC Genome Browser Utilities

This page contains links to tools and utilities created by the UCSC Genome Bioinformatics Group.

- [Batch Coordinate Conversion \(liftOver\)](#) - converts genome coordinates and genome annotation files between assemblies. The current version supports both forward and reverse conversions, as well as conversions between selected species.
- [DNA Duster](#) - removes formatting characters and other non-sequence-related characters from an input sequence. Offers several configuration options for the output format, including translated protein.
- [Protein Duster](#) - removes formatting characters and other non-sequence-related characters from an input sequence. Offers several configuration options for the output format.
- [Phylogenetic Tree Gif Maker](#) - creates a gif image from the phylogenetic tree specification given. Offers several configuration options for branch lengths, normalized lengths, branch labels, legend etc.
- [Executable and Source Code Downloads](#) - executable and source code downloads of the Genome Browser, Blat and liftOver.

Acknowledgements

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RESEARCH
UK

CAMBRIDGE
INSTITUTE