

NATIONAL HUMAN GENOME RESEARCH INSTITUTE Division of Intramural Research



*Current Topics in Genome Analysis*  
*Spring 2008*

*Week 4: Mining Genomic Sequence Data*

*Tyra G. Wolfsberg, Ph.D.*

U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES | NATIONAL INSTITUTES OF HEALTH | genome.gov DIR



## Accessing public genome sequence data

UCSC's Genome Browser ("Golden Path")  
<http://genome.ucsc.edu>

NCBI's Map Viewer  
<http://www.ncbi.nlm.nih.gov/mapview/>

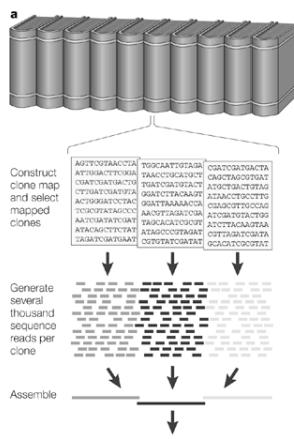
Ensembl  
<http://www.ensembl.org>

## Types of data integrated in genome browsers

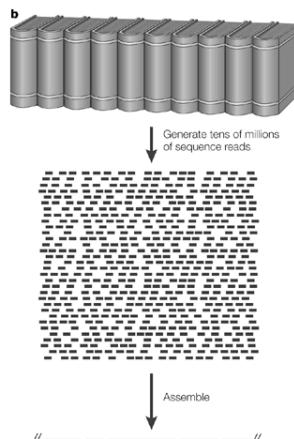
- Same starting material for all genome browsers: genomic sequence
- Annotations calculated independently by each genome browser
  - Genes
    - RefSeq mRNAs (non-redundant)
    - GenBank mRNAs (redundant)
    - Gene predictions
  - SNPs
  - Homologous sequences from other organisms
  - STSs

## Overview of genome sequencing strategies

Clone-by-clone shotgun sequencing



Whole-genome shotgun sequencing



Nature Reviews | Genetics  
Green ED. Strategies for the systematic sequencing of complex genomes.  
Nat Rev Genet. 2001; 2:573-83.

## Genome Sequence Assemblies

- Complex algorithms needed to incorporate all sequence data
- Assemblies updated periodically as new sequence becomes available
  - Mouse and human genomes assembled by NCBI
  - Other genomes assembled by sequencing centers or consortia
- Assemblies not updated concurrently by the three Genome Browsers
  - “Pre-release” assemblies and annotations available at
    - UCSC: <http://genome-test.cse.ucsc.edu/>
    - pre!Ensembl: <http://pre.ensembl.org/>
  - UCSC and Ensembl provide archive of all genome assemblies and annotations; NCBI provides only limited archive
- IF YOU ARE COMPARING DATA FROM DIFFERENT GENOME BROWSERS, MAKE SURE YOU ARE LOOKING AT THE SAME VERSION OF THE ASSEMBLY

## Genome Assembly Versions

	Same assembly?	UCSC	NCBI	Ensembl
Human	Yes	Mar 2006/hg18/Build 36.1	Build 36.2	Build 36
Mouse	YES	July 2007/mm9/Build 37	Build 37.1	Build 37
Dog		May 2005 /canFam 2.0	Build 2.1/CanFam 2.1	CanFam 2.0
Zebrafish	NO	July 2007/danRer5/Zv7	Zv6/build 2.1	Zv7
Rhesus	YES	Jan 2006/rheMac2/v.1.0, Mmul_051212	Build 1.1/v.1.0, Mmul_051212	Mmul_1

## NCBI Reference Sequences (RefSeqs)

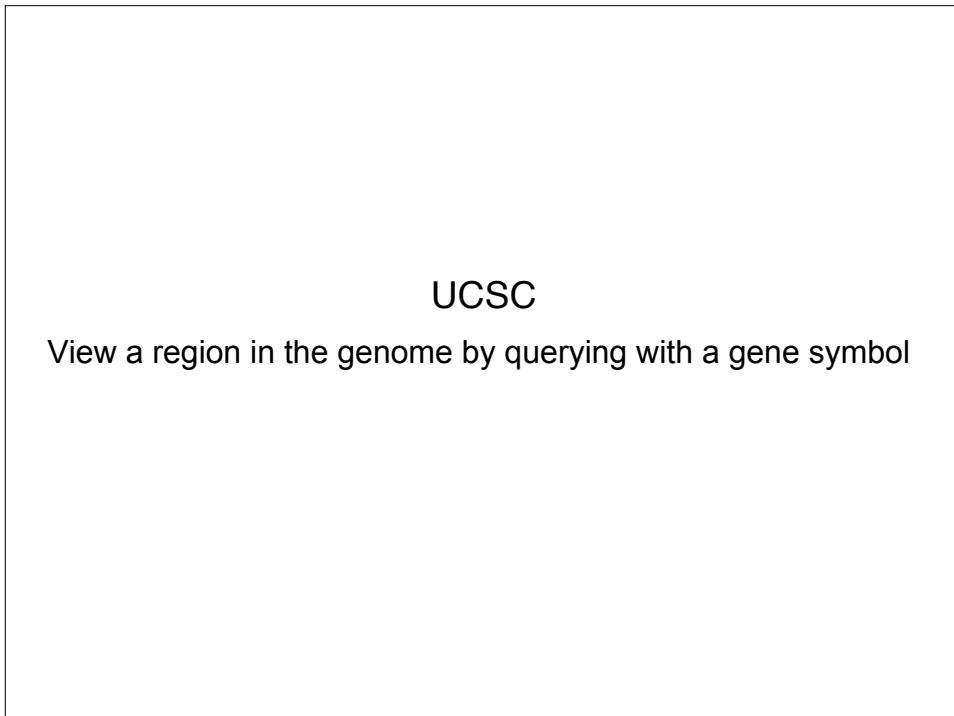
- Non-redundant collection of richly annotated DNA, RNA, and protein sequences from diverse taxa.
- Each RefSeq represents a single, naturally occurring molecule from one organism.

	derived from GenBank submissions	model reference sequences produced by NCBI's Genome Annotation project
mRNA	NM_123456	XM_123456
protein	NP_123456	XP_123456
non-coding transcripts	NR_123456	XR_123456

<http://www.ncbi.nlm.nih.gov/RefSeq/key.html>

### Beta actin mRNA RefSeq

REF ID: NM\_001101  
 DEFINITION Homo sapiens actin, beta (ACTB), mRNA. linear PRI 14-JAN-2008  
 ACCESSION NM\_001101  
 VERSION NM\_001101.2 GI:5016089  
 NUMBER OF RECORDS 1  
 SOURCE Homo sapiens (human)  
 ORGANISM Homo sapiens  
 TAXONOMY Metazoa; Chordata; Craniate; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Homorhini; Catarrhini; Hominidae; Homo.  
 (bases 1 to 1793)  
 REFERENCES (1)  
 AUTHORS Vilim,B.F., Roparski,S.B., Lindgren,M., Hammarstrom,P. and Jonsson,B.H.  
 TITLE Domain-specific chaperone-induced expansion is required for beta-actin folding: a comparison of beta-actin conformations upon interactions with GroEL and tail-less complex polypeptide I ring complex (TRIC)  
 JOURNAL Biochemistry 46 (44), 12639-12647 (2007)  
 PUBLISHER 1793480  
 ....  
 COMMENT REVISED REFSEQ. This record has been curated by NCBI staff. The reference sequence was derived from NM\_001101 and NC\_013727.  
 On Jun 8, 1999 this sequence version replaced gi|4501884.  
 Summary: This gene encodes one of six different actin proteins. Actins are highly conserved proteins that are involved in cell motility, structure, and integrity. This actin is a major constituent of the contractile apparatus and one of the two main muscle cytoskeletal actins.  
 Publication Note: This RefSeq record includes a subset of the publications available for this gene. Please see the Entrez Gene record to access additional publications.  
 COMPLETENESS: complete on the 3' end.  
 ....  
 CDS  
 74..1201  
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 /GO component="actin filament; cytoskeleton; TIP60 histone acetyltransferase complex [PMID:10966108]"  
 /GO process="ATP binding; nucleotide binding; protein binding [PMID:15527411]; structural constituent of cytoskeleton"  
 /function="beta cytoskeletal actin; F51TP5-binding protein 1; actin, tropomyosin 1; beta-actin"  
 /codon\_start=1  
 /product="beta actin"  
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 /db\_xref="MGI:1026332.1"  
 /db\_xref="G1:4501885"  
 /db\_xref="CCDS:CCDS5341.1"  
 /db\_xref="GeneID:160"  
 /db\_xref="MIM:138500"  
 /db\_xref="HPRD:00032"  
 /db\_xref="MIM:1026334"  
 ....  
 ORIGIN 1 cccgcacggcc cccggagcac agggcgcccp ctggccgtt ccggccggccg tccacacccg  
 61 cccgcacgtt accatggatc atggatatccg cccggcttcg gtcgacaaacg gtcggcgat  
 121 gtgcggccg ggtttcgccg gagcgatgc ccccgccgc gttttccctt ccatcggtt



UCSC Genome Bioinformatics

Genomes - Blat - Tables - Gene Sorter - PCR - VisiGene - Proteome - Session - FAQ - Help

**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 a portal to the ENCODE project.

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 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](#). To view the results of the Genome Browser users' survey we conducted in May 2007, click [here](#).

**News**

To receive announcements of new genome assembly releases, new software features, updates and training seminars by email, subscribe to the [genome-announce mailing list](#).

**8 Jan. 2008 - Additional Job Opening with UCSC Genome Browser Project**

In addition to the openings listed in the 12 Dec. 2007 announcement (see below), the UCSC Genome Browser project is accepting applications for Research Software Architect, a position in the UC Project Scientist academic series. For the job description, qualifications and application information, please see the Center for Biomolecular Science and Engineering [website](#). To ensure full consideration, applications must be received by 22 Jan. 2008.

**12 Dec. 2007 - Job Openings with UCSC Genome Browser Project**

The UCSC Genome Browser project is currently accepting applications for two positions on our development team: Software Development Engineer (Programmer/Analyst 3) and Biological Database Testing/User Support Technician (Programmer/Analyst 1). We are looking for talented self-motivated individuals who would like to use their skills in computer science, biology, and bioinformatics on a fast-paced project featuring the work of top genomics scientists worldwide.

For a summary of the position details and qualifications, see the entries for [Software Developer](#) and [Biological Database Testing/User Support Technician](#) on the Center for Biomolecular Science and Engineering (CBSE) website. For detailed job descriptions and application information, go to the [UCSC Staff Employment](#) website, click the "Search Postings" link on the sidebar, and type in job #0701419

**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: Vertebrate; genome: Human; assembly: Mar. 2006; position or search term: ADAM2; image width: 820; submit; click

**About the Human Mar. 2006 (hg18)**  
 The March 2006 human reference sequence (hg18) was produced by the International Human Genome Sequencing Consortium.

**Sample position queries**  
 A genome position can be specified by its chromosomal coordinate range, or keyword. For example, "Tetradon" displays all chromosomes of Tetradon, "fugu" displays the genome of fugu, "zebrafish" displays the genome of zebrafish, "Medaka" displays the genome of Medaka.

**Request:** Genome Browser Response:

- chr7 Displays all of chromosome 7
- 20p13 Displays region for band p13 on chr 20
- chr3:1-1000000 Displays first million bases of chr 3, counting from p arm telomere
- chr3:1000000+2000 Displays a region of chr3 that spans 2000 bases, starting with position 1000000
- D16S3046 Displays region around STS marker D16S3046 from the Genethon/Marshfield maps. Includes 100,000 bases on each side as well.
- RH18061;RH80175 Displays region between STS markers RH18061;RH80175. This syntax may also be used for other range queries, such as between cytobands and uniquely-determined ESTs, mRNAs, refSeqs, etc.
- AA205474 Displays region of EST with GenBank accession AA205474 in BRCA1 cancer gene on chr 17
- AC008101 Displays region of clone with GenBank accession AC008101
- AF083811 Displays region of mRNA with GenBank accession number AF083811
- PRNP Displays region of genome with HUGO Gene Nomenclature Committee identifier PRNP
- NM\_017414 Displays the region of genome with RefSeq identifier NM\_017414
- NP\_059110 Displays the region of genome with protein accession number NP\_059110
- pseudogene mRNA Lists transcribed pseudogenes, but not cDNAs
- homeobox caudal Lists mRNAs for caudal homeobox genes

**UCSC Genes**

Human ADAM2 – UCSC Genome Browser v174

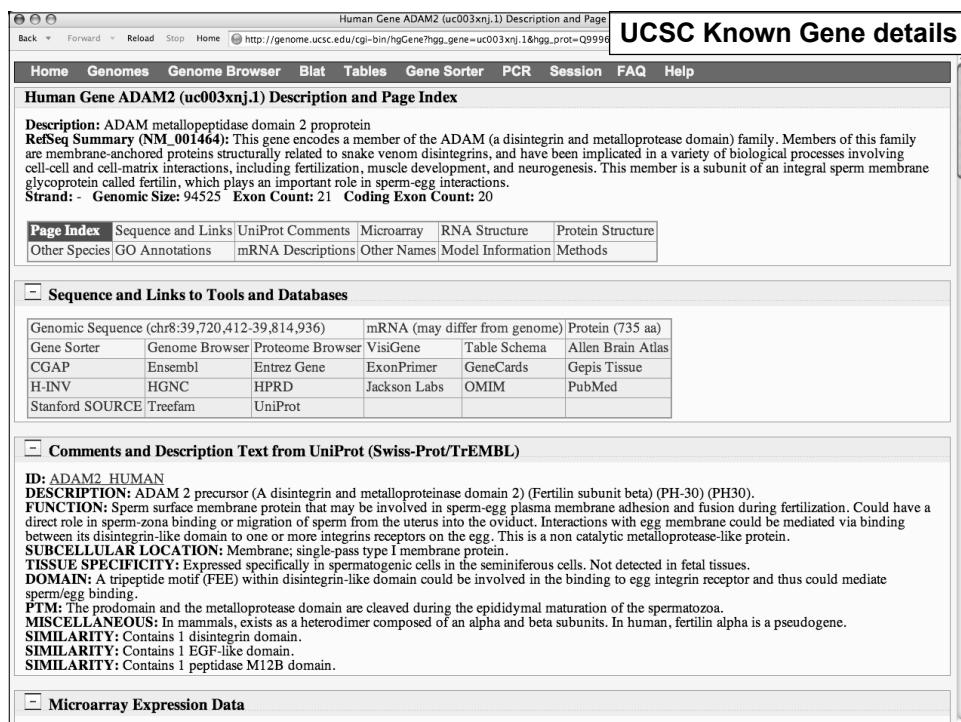
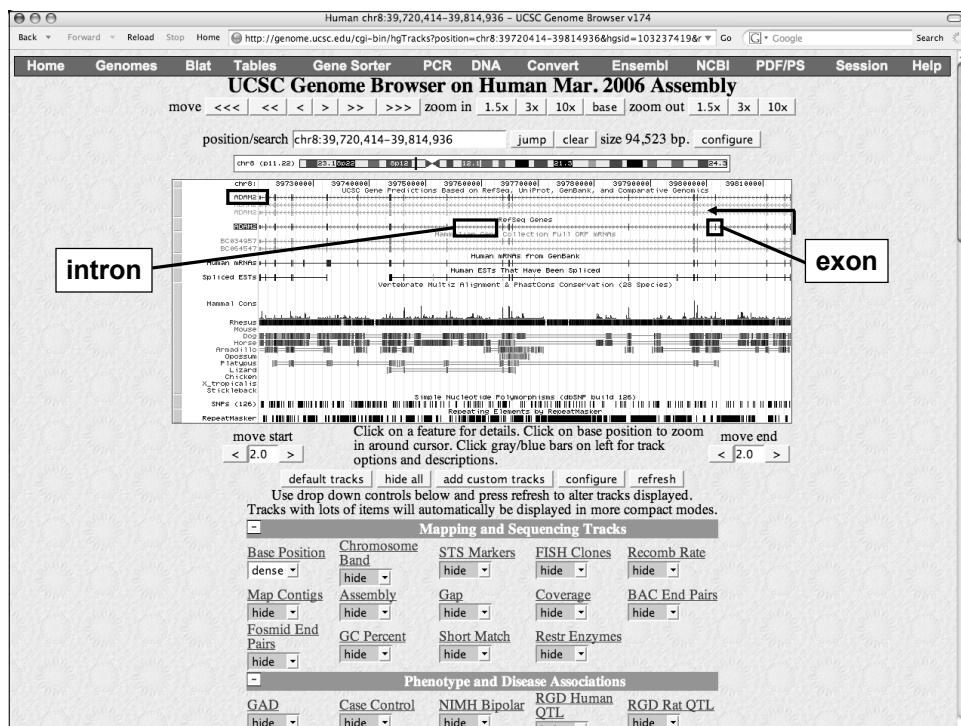
ADAM2 (uc003xnl\_1) at chr8:39720412-39814936 - ADAM metallopeptidase domain 2 proprotein  
 ADAM2 (uc003xnl\_1) at chr8:39720414-39814936 - ADAM metallopeptidase domain 2 proprotein  
 ADAM2 (uc003xnl\_1) at chr8:39720414-39814936 - ADAM metallopeptidase domain 2 proprotein  
 ADAM28 (uc003xdy\_1) at chr8:24207525-24268671 - ADAM metallopeptidase domain 28 isoform 1  
 ADAM28 (uc003xdy\_1) at chr8:24207525-24268671 - ADAM metallopeptidase domain 28 isoform 3  
 ADAM22 (uc003ujp\_1) at chr7:87401638-87664385 - ADAM metallopeptidase domain 22 isoform 4  
 ADAM22 (uc003ujp\_1) at chr7:87401638-87664385 - ADAM metallopeptidase domain 22 isoform 3  
 ADAM22 (uc003ujp\_1) at chr7:87401638-87664385 - ADAM metallopeptidase domain 22 isoform 1  
 ADAM22 (uc003ujp\_1) at chr7:87401638-87664385 - ADAM metallopeptidase domain 22 isoform 2  
 ADAM22 (uc003ujp\_1) at chr7:87401638-87664385 - ADAM metallopeptidase domain 22 isoform 5  
 ADAM22 (uc003ujp\_1) at chr7:87401638-87664386 - ADAM metallopeptidase domain 22 isoform 4  
 ADAM22 (uc003ujp\_1) at chr7:87401638-87664386 - ADAM metallopeptidase domain 22 isoform 5  
 ADAM22 (uc003ujp\_1) at chr7:87401638-87664386 - ADAM metallopeptidase domain 22 isoform 6  
 ADAM22 (uc003ujp\_1) at chr7:87401638-87664386 - ADAM metallopeptidase domain 22 isoform 5  
 ADAM29 (uc003lue\_1) at chr4:1760988112-176135906 - ADAM metallopeptidase domain 29 preproprotein  
 ADAM29 (uc003lue\_1) at chr4:176076134-176135906 - ADAM metallopeptidase domain 29 preproprotein  
 ADAM29 (uc003lue\_1) at chr4:176076134-176135906 - ADAM metallopeptidase domain 29 preproprotein  
 ADAM29 (uc002vbg\_1) at chr2:207016613-207190924 - ADAM metallopeptidase domain 23 preproprotein  
 ADAM20 (uc001xme\_1) at chr4:700598831-70071485 - ADAM metallopeptidase domain 20 preproprotein  
 ADAM21 (uc001xmd\_1) at chr4:69993970-69994375 - ADAM metallopeptidase domain 21 preproprotein

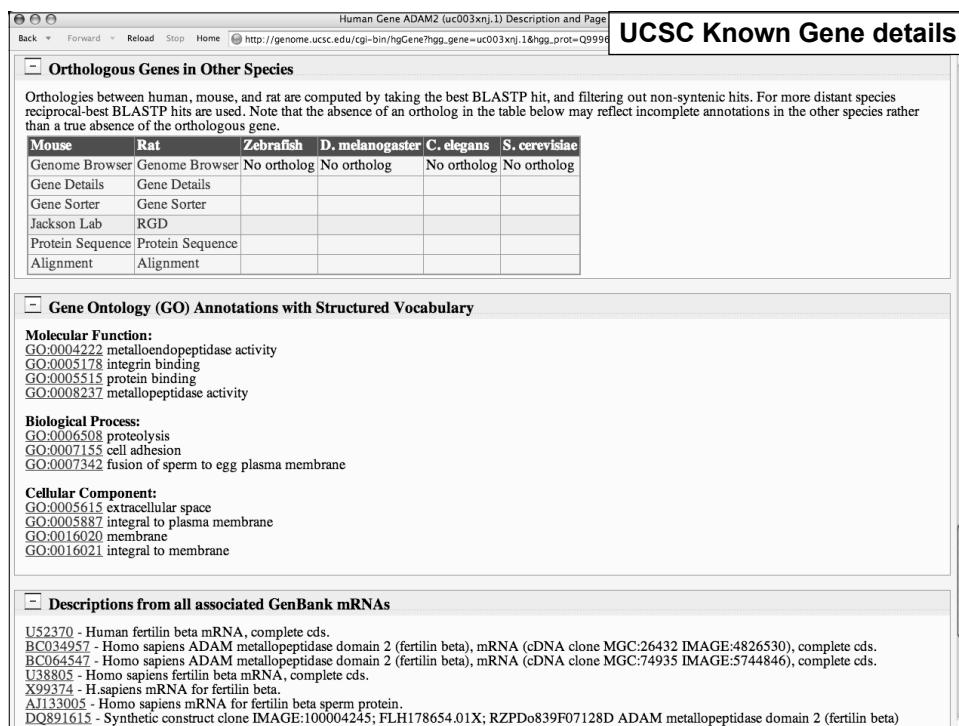
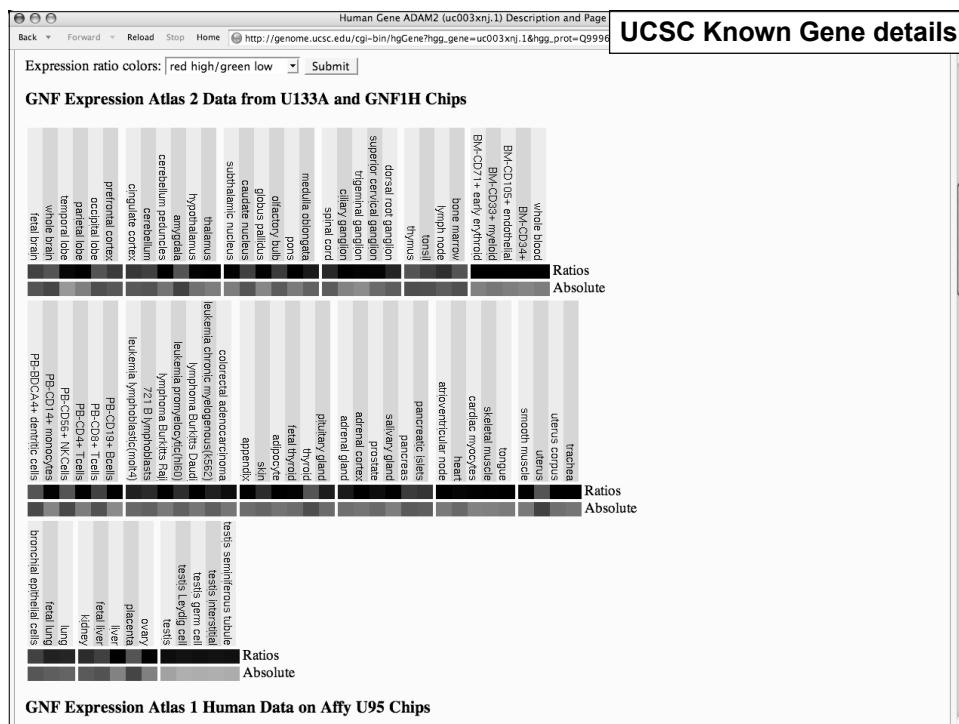
**RefSeq Genes**

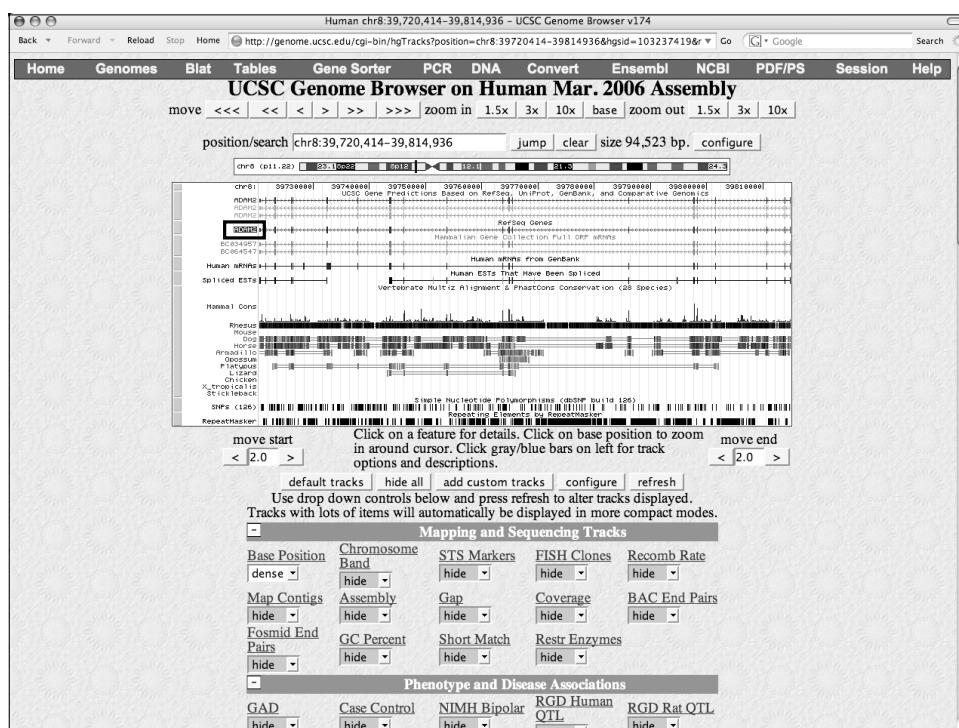
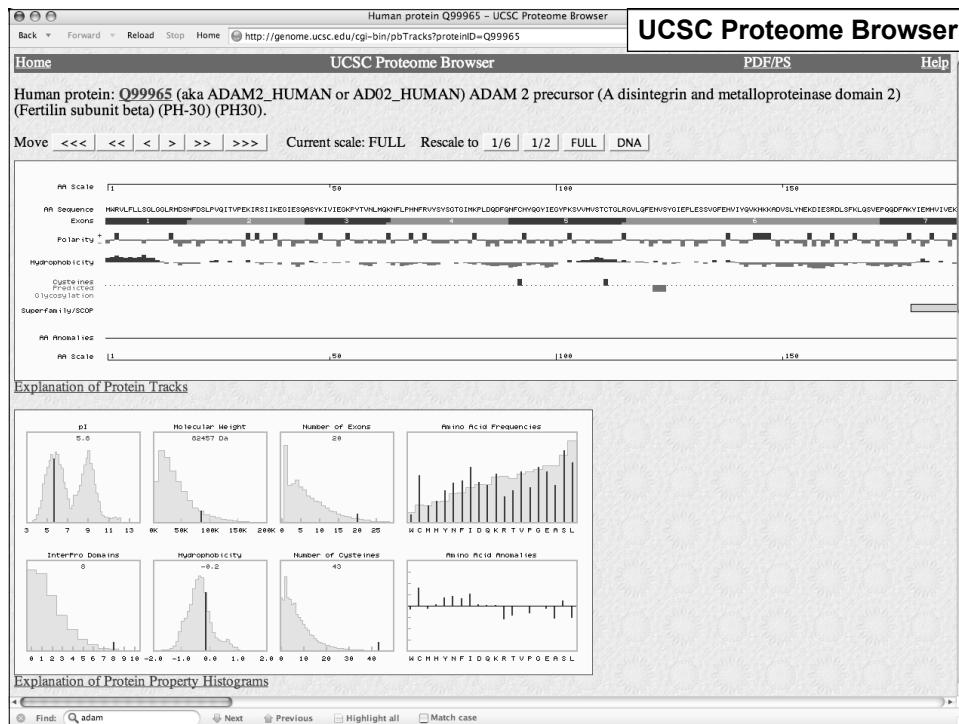
ADAM2 at chr8:39720414-39814936 - (NM\_001464) ADAM metallopeptidase domain 2 proprotein  
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 ADAM21 at chr14:69993970-69996374 - (NM\_003813) ADAM metallopeptidase domain 21 preprotein  
 ADAM22 at chr7:87401638-87664386 - (NM\_001944) ADAM metallopeptidase domain 22 isoform 4  
 ADAM22 at chr7:87401638-87664386 - (NM\_021720) ADAM metallopeptidase domain 22 isoform 2  
 ADAM22 at chr7:87401638-87664383 - (NM\_021722) ADAM metallopeptidase domain 22 isoform 2  
 ADAM22 at chr7:87401638-87664383 - (NM\_021723) ADAM metallopeptidase domain 22 isoform 1  
 ADAM22 at chr7:87401638-87664383 - (NM\_016351) ADAM metallopeptidase domain 22 isoform 3  
 ADAM23 at chr2:207016613-207190922 - (NM\_003812) ADAM metallopeptidase domain 23 preproprotein  
 ADAM23 at chr2:207016613-207190922 - (NM\_021721) ADAM metallopeptidase domain 23 preproprotein  
 ADAM28 at chr8:24207525-24268670 - (NM\_014265) ADAM metallopeptidase domain 28 isoform 1  
 ADAM29 at chr4:176076134-176135905 - (NM\_014269) ADAM metallopeptidase domain 29 preproprotein

**Non-Human RefSeq Genes**

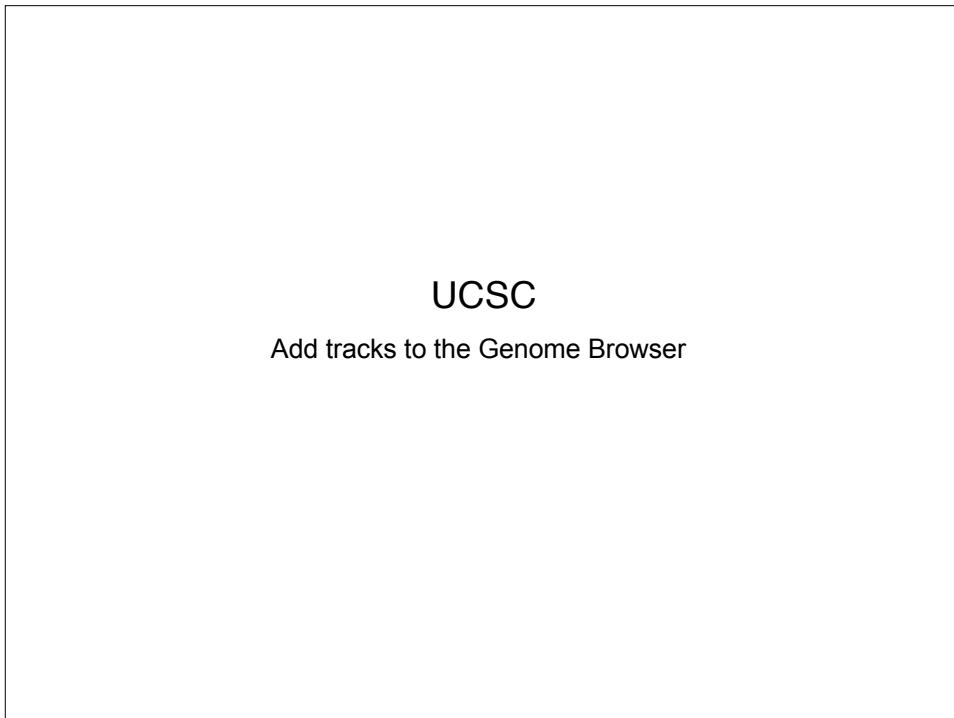
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 ADAM2 at chr8:39723146-39814802 - (NM\_213957) ADAM metallopeptidase domain 2  
 Adam2 at chr8:39723147-3981569 - (NM\_009618) a disintegrin and metalloprotease domain 2  
 Adam2 at chr8:39723147-39813877 - (NM\_020077) a disintegrin and metalloprotease domain 2  
 Adam21 at chr14:69988547-69996354 - (NM\_020330) a disintegrin and metalloprotease domain 21  
 Adam21 at chr14:69781958-7061197 - (NM\_020330) a disintegrin and metalloprotease domain 21  
 Adam22 at chr7:87401747-87642822 - (NM\_001007221) a disintegrin and metalloprotease domain 22  
 Adam22 at chr7:87401747-87670128 - (NM\_001098225) a disintegrin and metalloprotease domain 22  
 Adam23 at chr2:207016503-20719450 - (NM\_011780) a disintegrin and metalloprotease domain 23  
 Adam23 at chr2:207016759-207190724 - (NM\_001029899) a disintegrin and metalloprotease domain 23  
 Adam25 at chr4:18890467-18890635 - (NM\_010086) a disintegrin and metalloproteinase domain 24  
 Adam25 at chr4:18890441-188906248 - (NM\_011781) a disintegrin and metalloprotease domain 25





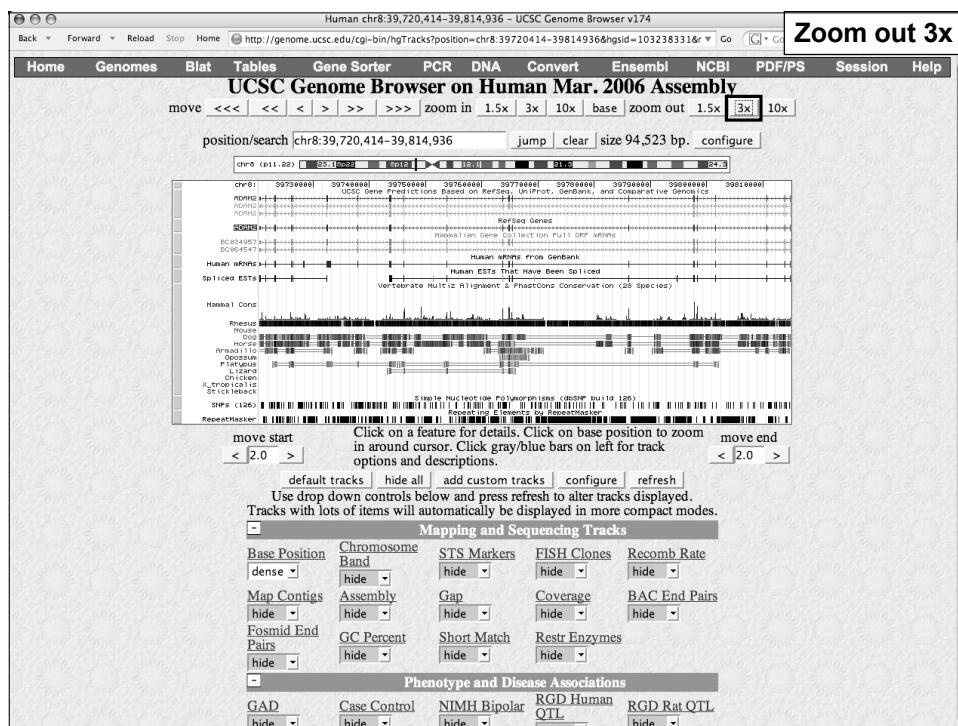


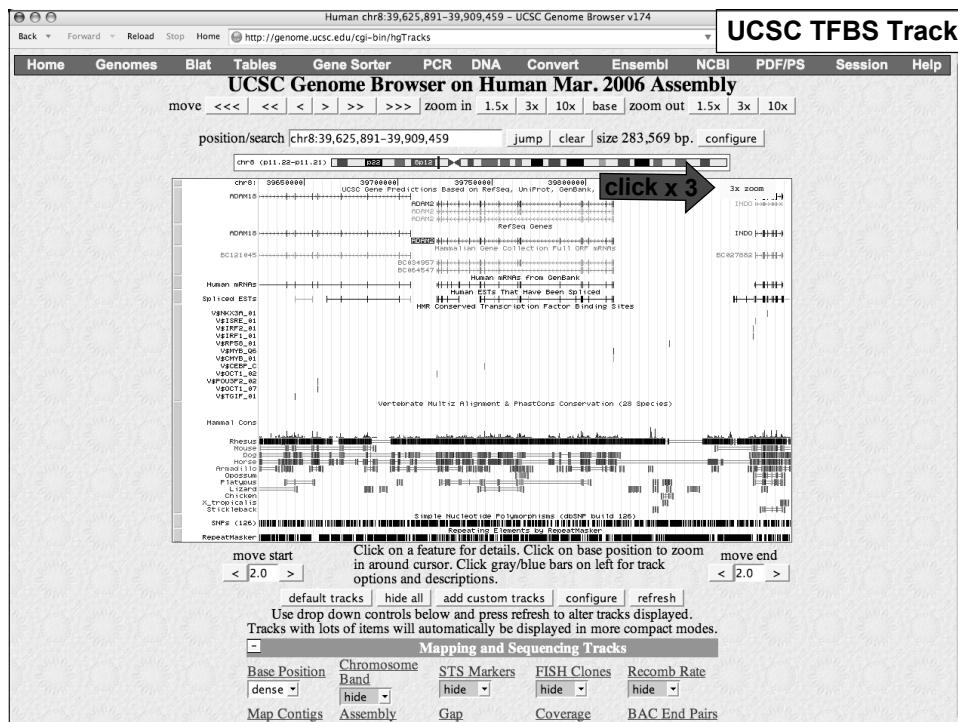
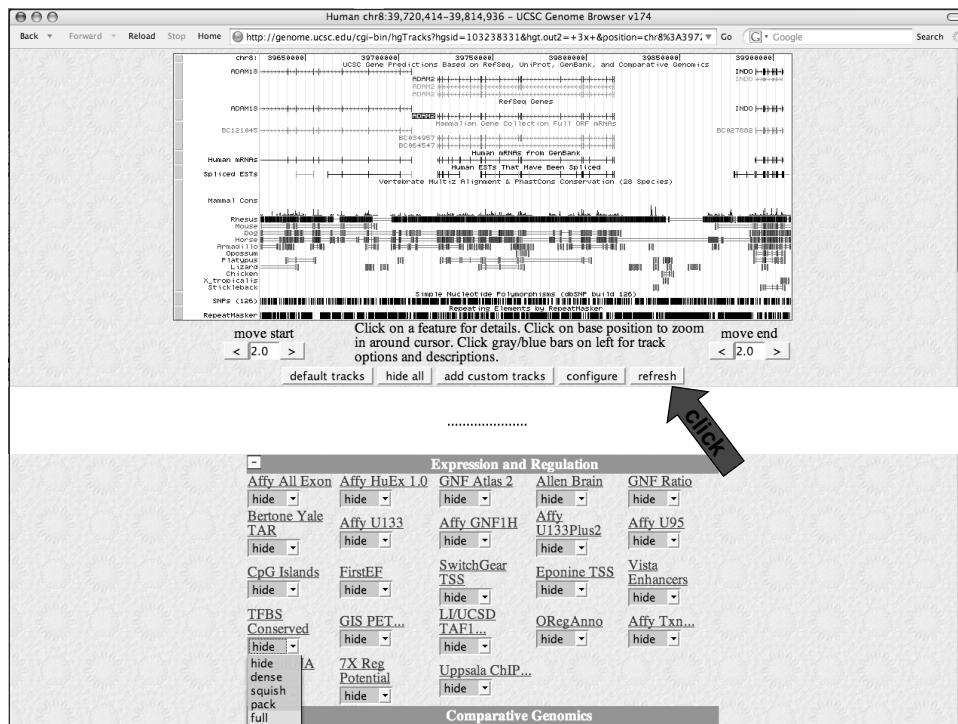


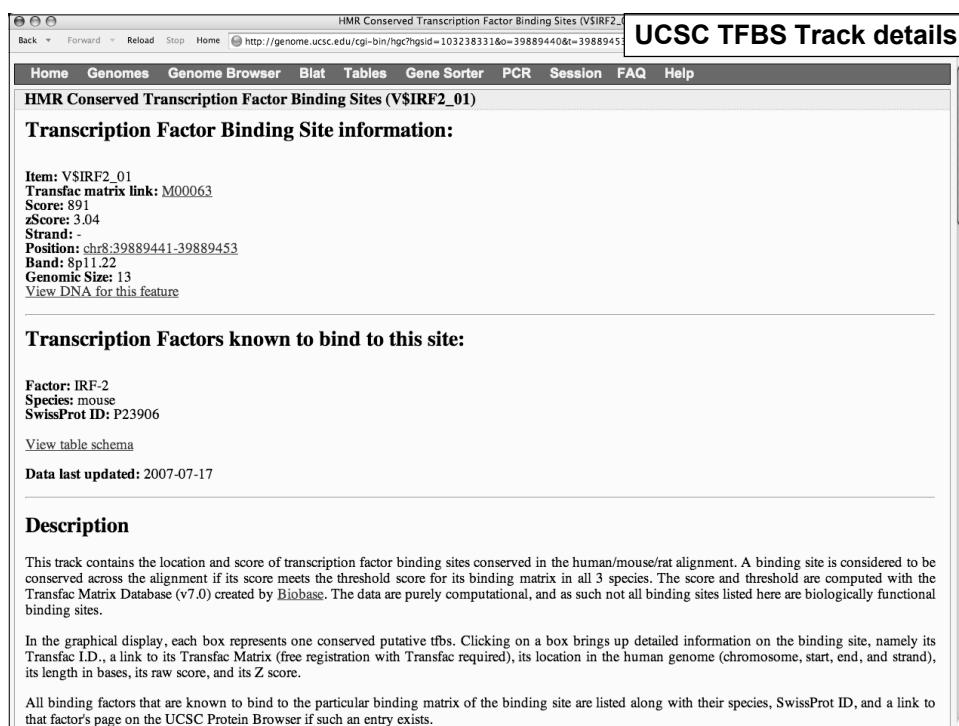
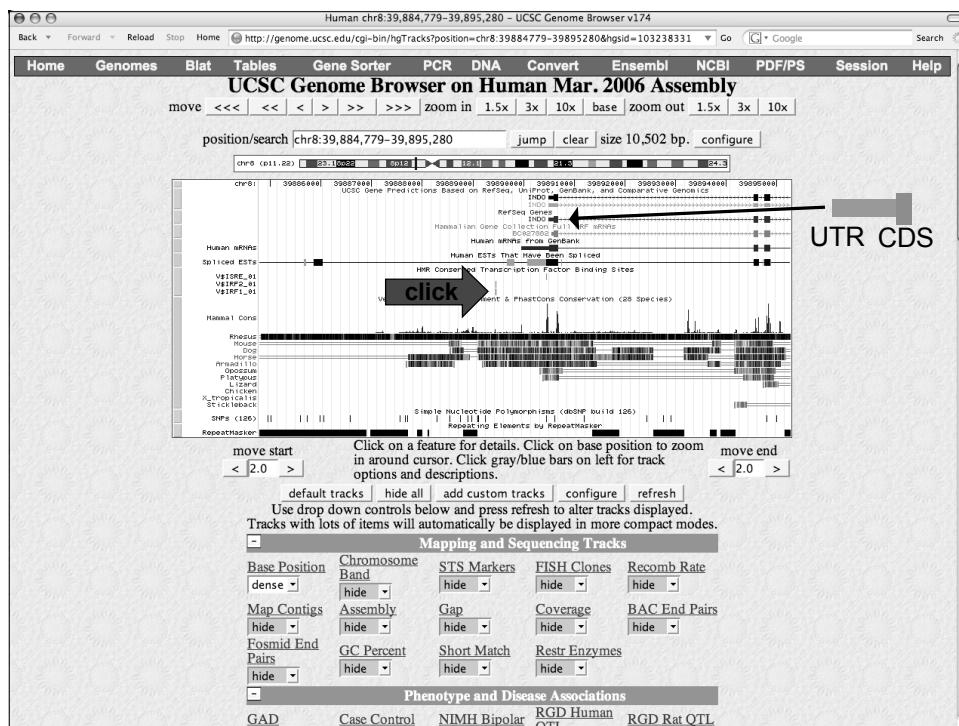


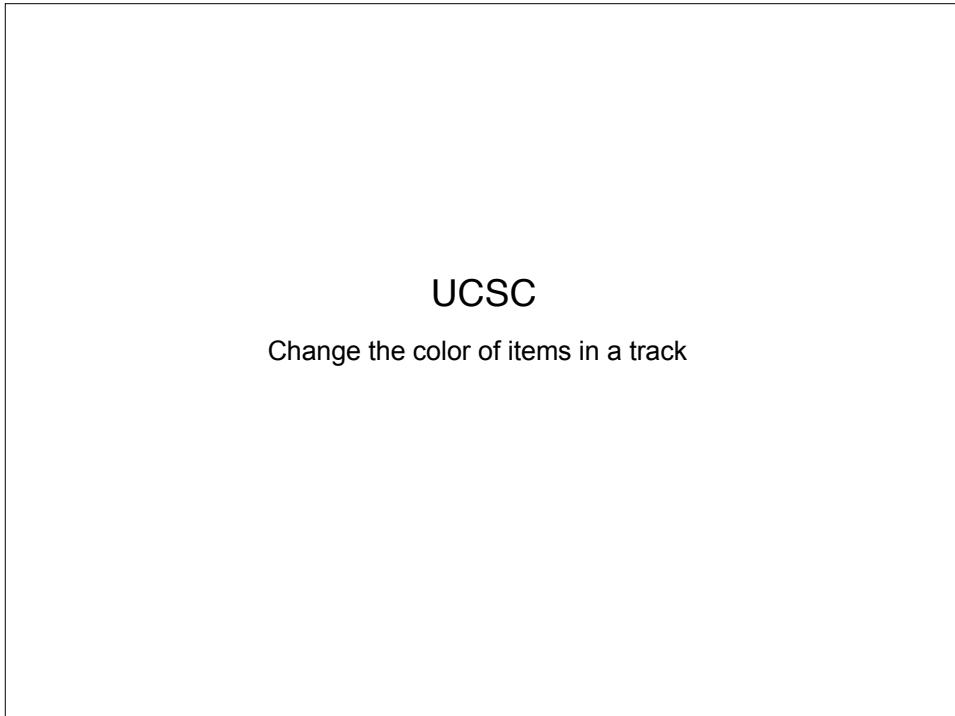
## UCSC

Add tracks to the Genome Browser









## UCSC

Change the color of items in a track

Human chr8:39,720,414-39,814,936 – UCSC Genome Browser v174

Back Forward Reload Stop Home http://genome.ucsc.edu/cgi-bin/hgTracks?position=chr8:39720414-39814936&hgSID=103238331&r Go Google Search

Home Genomes Blat Tables Gene Sorter PCR DNA Convert Ensembl NCBI PDF/PS Session Help

UCSC Genome Browser on Human Mar. 2006 Assembly

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

position/search chr8:39,720,414-39,814,936 jump clear size 94,523 bp. configure

chr8 (p11.22) 39720414 39814936 39720414 39814936 39720414 39814936 39720414 39814936

ORF6: 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39814936 Gene Predictions Based on RefSeq, Uniprot, GenBank, and Comparative Genomics

ADM2: 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39814936 RefSeq Genes

PCMV: 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39814936 Human mRNAs from GenBank

BC041957: 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39814936 Human ESTs That Have Been Spliced

Human refSeq: 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39814936 Human mRNAs from GenBank

SilicoESTs: 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39814936 Human ESTs That Have Been Spliced

Vertebrate Multiz Alignment & PhastCons Conservation (28 Species)

Human Cons: 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39814936 Human Conservation Scores

Chimpanzee: 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39814936 Chimp Conservation Scores

Rhesus: 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39814936 Rhesus Conservation Scores

Mouse: 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39814936 Mouse Conservation Scores

House Sparrow: 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39814936 House Sparrow Conservation Scores

Opossum: 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39814936 Opossum Conservation Scores

Lizard: 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39814936 Lizard Conservation Scores

Chicken: 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39814936 Chicken Conservation Scores

X-Trop-Callid: 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39814936 X-Trop-Callid Conservation Scores

Strewnius: 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39720414 39814936 Strewnius Conservation Scores

SNPs (126): hide dense move start < 2.0 > Click on a feature for details. Click on base position to zoom in around cursor. Click gray/blue bars on left for track options and descriptions. move end < 2.0 >

Variation and Repeats

SNP Arrays: hide move start < 2.0 > Click on a feature for details. Click on base position to zoom in around cursor. Click gray/blue bars on left for track options and descriptions. move end < 2.0 >

HapMap SNPs: hide move start < 2.0 > Click on a feature for details. Click on base position to zoom in around cursor. Click gray/blue bars on left for track options and descriptions. move end < 2.0 >

HapMap LD Unph.: hide move start < 2.0 > Click on a feature for details. Click on base position to zoom in around cursor. Click gray/blue bars on left for track options and descriptions. move end < 2.0 >

Structural Var: hide move start < 2.0 > Click on a feature for details. Click on base position to zoom in around cursor. Click gray/blue bars on left for track options and descriptions. move end < 2.0 >

Segmental Dups: hide move start < 2.0 > Click on a feature for details. Click on base position to zoom in around cursor. Click gray/blue bars on left for track options and descriptions. move end < 2.0 >

Exapted Repeats: hide move start < 2.0 > Click on a feature for details. Click on base position to zoom in around cursor. Click gray/blue bars on left for track options and descriptions. move end < 2.0 >

RepeatMasker: dense move start < 2.0 > Click on a feature for details. Click on base position to zoom in around cursor. Click gray/blue bars on left for track options and descriptions. move end < 2.0 >

Interrupted Rpts: hide move start < 2.0 > Click on a feature for details. Click on base position to zoom in around cursor. Click gray/blue bars on left for track options and descriptions. move end < 2.0 >

Simple Repeats: hide move start < 2.0 > Click on a feature for details. Click on base position to zoom in around cursor. Click gray/blue bars on left for track options and descriptions. move end < 2.0 >

Microsatellite: hide move start < 2.0 > Click on a feature for details. Click on base position to zoom in around cursor. Click gray/blue bars on left for track options and descriptions. move end < 2.0 >

Self Chain: hide move start < 2.0 > Click on a feature for details. Click on base position to zoom in around cursor. Click gray/blue bars on left for track options and descriptions. move end < 2.0 >

Structural Var: hide move start < 2.0 > Click on a feature for details. Click on base position to zoom in around cursor. Click gray/blue bars on left for track options and descriptions. move end < 2.0 >

Variation and Repeats

**click**

**UCSC SNP Track details**

**SNPs (126) Track Settings**

**Simple Nucleotide Polymorphisms (dbSNP build 126)**

**Display mode:** full | hide | Submit

**Include Chimp dense observed human alleles in name:**  (If enabled, chimp & squish is displayed first, then '>', then human alleles).

**Minimum Ave\_Zygosity:** 0

**Maximum Weight:** 3 *SNPs with higher weights are less reliable*

Any type of data can be excluded from view by deselecting the checkbox below. Not all assemblies include values in all categories.

**Location Type:**  Unknown  Range  Exact  Between  RangeInsertion  RangeSubstitution  RangeDeletion

**Class:**  Unknown  Single Nucleotide Polymorphism  In/Del  Heterozygous  Microsatellite  Named  No Variation  Mixed  Mnp  Insertion  Deletion

**Validation:**  Unknown  By Cluster  By Frequency  By Submitter  By 2 Hit / 2 Allele  By HapMap

**Function:**  Unknown  Locus  Coding - Synonymous  Coding - Non-Synonymous  Untranslated  Intron  Splice Site  Reference (coding)

**Molecule Type:**  Unknown  Genomic  cDNA

**SNP Feature for Color Specification:** Function

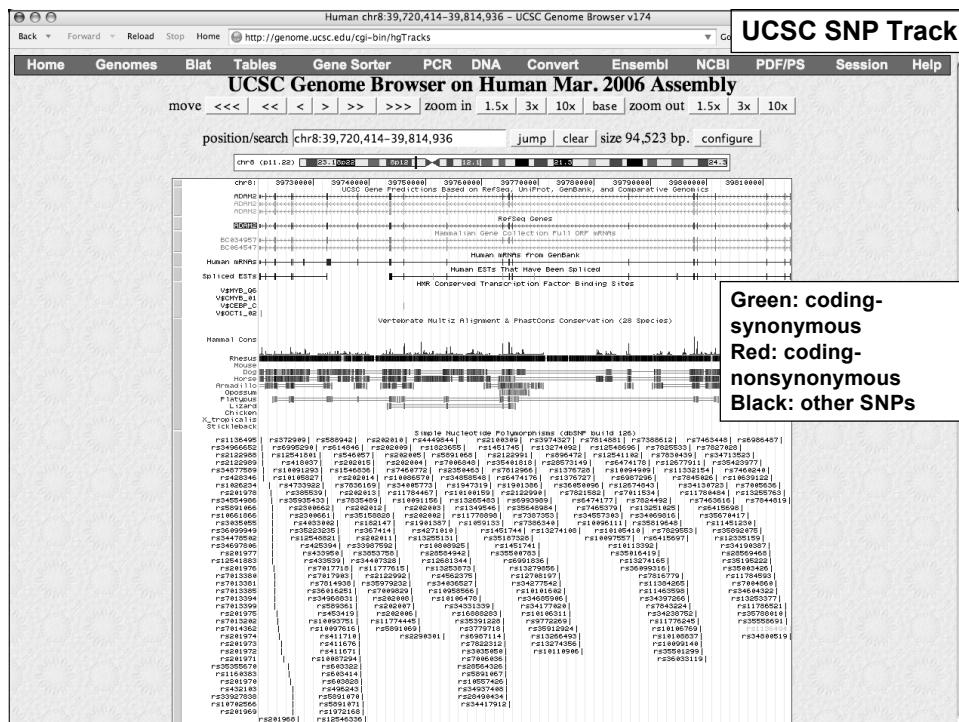
The selected feature above has the following values below. For each value, a selection of colors is available.

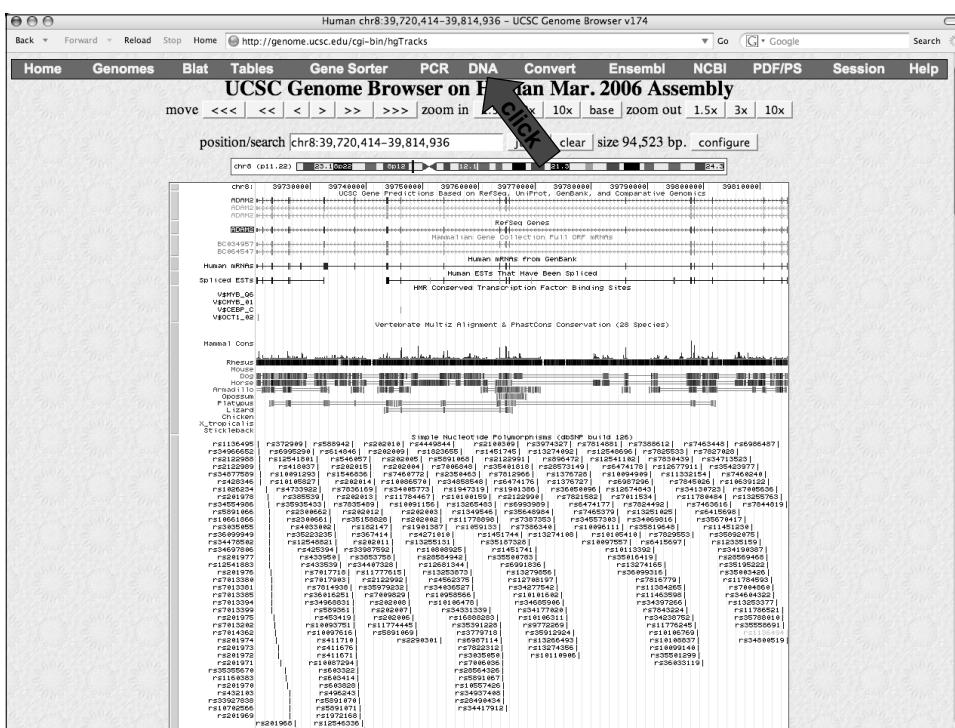
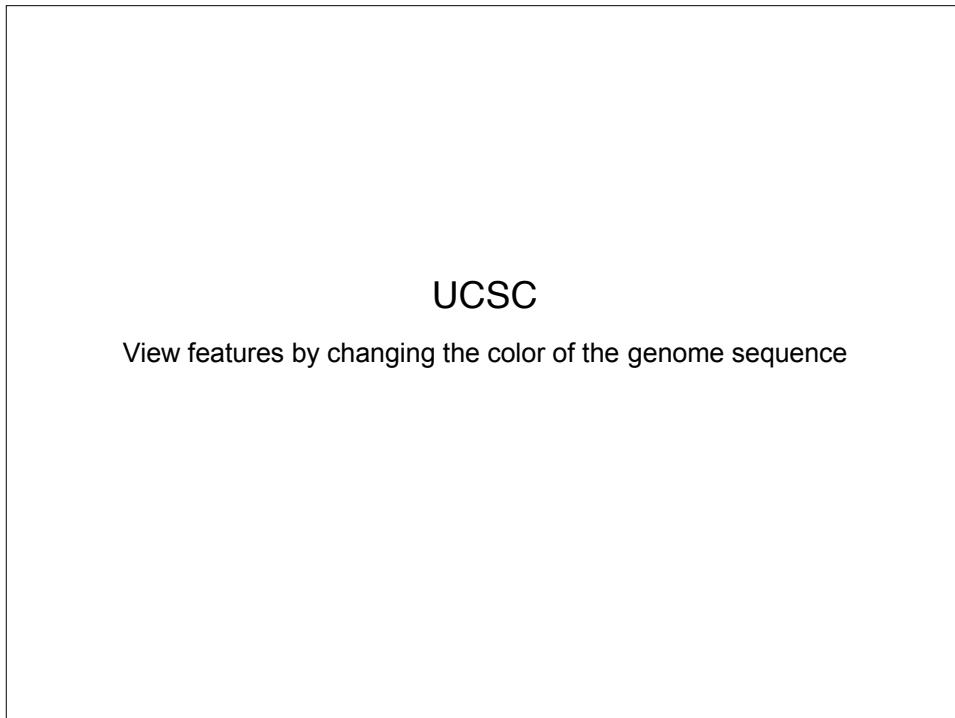
Unknown: black	Locus: black	Coding - Synonymous: green	Coding - Non-Synonymous: red	Untranslated: black	Intron: black
Splice Site: black	Reference (coding): black				

[View table schema](#)

**Data last updated:** 2006-10-02

**Description:**







**NCBI Entrez Protein**

Search | Protein for np\_001455

About Entrez  
Help & FAQ  
Entrez Tools  
Check sequence revision history  
LinkOut  
My NCBI  
Related resources  
BLAST  
Reference sequence project  
Search for Genes  
Clusters of orthologous groups  
Protein reviews on the web  
Search for full length cDNAs

Display: Summary Show 20 Sort by Relevance Send to

All: 1 Summary ASN.1

1: FASTA ADAM GenPept GI List Graphics TinySeq XML INSDSeq XML LinkOut Related Sequences Conserved Domain Links 3D Domain Links Gene Links Genome Links Gene Project Links HemoloGene Links Nucleotide Links NIH eDNA clone links

BLINK, Conserved Domains, Links

**NCBI Entrez Protein**

Search | Protein for np\_001455

Display: FASTA Show 20 Send to

Range: from begin to end Refresh

**I: NP\_001455. Reports ADAM metallopeptidase domain 2 proprotein [Homo sapiens]**

```
>gi|55743080|ref|NP_001455.3| ADAM metallopeptidase domain 2 proprotein [Homo sapiens]
MVRVFLLSGLGLRMDNSFDLPLVQITVPBKIRSIIKGKIESQASYKIVIEGKPYTVNLMKNFLPHNF
YQVHKHADQSVLYNEKD1ESSRLSLFKLQSVPPQDPAKYIEMHVIVBQLYNHMSDTTVAQKVFLIG
LTNAIPVSFNITIILSSELWIDENKIAZTGEANELLFTLWNKTSVLWLRPHDVFALLVREKSNVYGA
TPNCFSPEDPAPF1SK9KSCQCLNQPRLDPPFFQQAQCVCNAKLEAGEB3CDCCTGTQDCAL1GTCDD1ATCR
PSNGCSFEDPAPF1SK9KSCQCLNQPRLDPPFFQQAQCVCNAKLEAGEB3CDCCTGTQDCAL1GTCDD1ATCR
FKAGNSCAEGPCENCFLMSKERNCRPSFEECCLQEYCNGSSASCPEHNHYVGTCGCLNQWIC1DGVMC
SGDKQCTTGFKEVFFGSKCYSHLNLSKTDVSNGCG1SD8GTYQCEADNTCGG1CKYVGKFLL1OIFRA
TIIYANISGHU1CAVEFASDHADQSMMKIKOCGTFSGSNKVRCSNQCVCSSSYLGDCCTDKNDRGVCNNK
KUCHCSCASLYLPDCSVQSDLNPGCGSIDGNFPFPVAPARLPERRYIBNIYHSKPMRWFPLF1PFPIIFC
VLI1AIMVKVNFRQRKWRTEDYSSD8QPESESEPK0
```

**UCSC BLAT search**

Home Genomes Tables PCR Session FAQ Help

Chicken BLAT Search

**BLAT Search Genome**

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

Assembly: Feb. 2004

Query type: BLAT's guess

Sort output: query.score

Output type: hyperlink

Sequence: >gi|55743080|ref|NP\_001455.3| ADAM metallopeptidase domain 2 proprotein [Homo sapiens]

MVRVFLLSGLGLRMDNSFDLPLVQITVPBKIRSIIKGKIESQASYKIVIEGKPYTVNLMKNFLPHNF
YQVHKHADQSVLYNEKD1ESSRLSLFKLQSVPPQDPAKYIEMHVIVBQLYNHMSDTTVAQKVFLIG
LTNAIPVSFNITIILSSELWIDENKIAZTGEANELLFTLWNKTSVLWLRPHDVFALLVREKSNVYGA
TPNCFSPEDPAPF1SK9KSCQCLNQPRLDPPFFQQAQCVCNAKLEAGEB3CDCCTGTQDCAL1GTCDD1ATCR
PSNGCSFEDPAPF1SK9KSCQCLNQPRLDPPFFQQAQCVCNAKLEAGEB3CDCCTGTQDCAL1GTCDD1ATCR
FKAGNSCAEGPCENCFLMSKERNCRPSFEECCLQEYCNGSSASCPEHNHYVGTCGCLNQWIC1DGVMC
SGDKQCTTGFKEVFFGSKCYSHLNLSKTDVSNGCG1SD8GTYQCEADNTCGG1CKYVGKFLL1OIFRA
TIIYANISGHU1CAVEFASDHADQSMMKIKOCGTFSGSNKVRCSNQCVCSSSYLGDCCTDKNDRGVCNNK
KUCHCSCASLYLPDCSVQSDLNPGCGSIDGNFPFPVAPARLPERRYIBNIYHSKPMRWFPLF1PFPIIFC
VLI1AIMVKVNFRQRKWRTEDYSSD8QPESESEPK0

submit | I'm feeling lucky | clear |

Paste in a query sequence to find its location in the genome. Multiple sequences may be searched if separated by lines starting with > followed by the sequence name.

**File Upload:** Rather than pasting a sequence, you can choose to upload a text file containing the sequence.  
Upload sequence:  Browse... | submit file |

Only DNA sequences of 25,000 or fewer bases and protein or translated sequence of 10000 or fewer letters will be processed. Up to 25 sequences can be submitted at the same time. The total limit for multiple sequence submissions is 50,000 bases or 25,000 letters.

For locating PCR primers, use In-Silico PCR for best results instead of BLAT.

**About BLAT**

BLAT on DNA is designed to quickly find sequences of 95% and greater similarity of length 25 bases or more. It may miss more divergent or shorter sequence alignments. It will find perfect sequence matches of 33 bases, and sometimes find them down to 20 bases. BLAT on proteins finds sequences of 80% and greater similarity of length 20 amino acids or more. In practice DNA BLAT works well on primates, and protein BLAT on land vertebrates.

Chicken BLAT Results  
 Back Forward Reload Stop Home http://genome.ucsc.edu/cgi-bin/hgBlat

**UCSC BLAT search**

**Home Genomes Tables PCR Session FAQ Help**

**Chicken BLAT Results**

**BLAT Search Results**

ACTIONS	QUERY	SCORE	START	END	QSIZE	IDENTITY	CHRO	STRAND	START	END	SPAN
browser details	NP_001455.3	44	539	600	735	71.6%	Un	++	635370	635555	186
browser details	NP_001455.3	12	301	304	735	100.0%	1	++	67659709	67659720	12
browser details	NP_001455.3	12	437	440	735	100.0%	1	++	67660117	67660128	12
browser details	NP_001455.3	12	385	390	735	83.4%	1	++	67659961	67659978	18

**UCSC Genome Browser on Chicken Feb. 2004 Assembly**  
 move <<< | << | < | > | >> | zoom in 1.5x 3x 10x | base | zoom out 1.5x 3x 10x |  
 position/search chrUn:635,370–635,555 jump clear size 186 bp. configure |

635450 635460 635470 635480 635490 635500 635510 635520 635530 635540 635550 635560 635570 635580 635590 635600 635610 635620 635630 635640 635650 635660 635670 635680 635690 635700 635710 635720 635730 635740 635750 635760 635770 635780 635790 635800 635810 635820 635830 635840 635850 635860 635870 635880 635890 635900 635910 635920 635930 635940 635950 635960 635970 635980 635990 635600 635610 635620 635630 635640 635650 635660 635670 635680 635690 635700 635710 635720 635730 635740 635750 635760 635770 635780 635790 635800 635810 635820 635830 635840 635850 635860 635870 635880 635890 635900 635910 635920 635930 635940 635950 635960 635970 635980 635990 635991 635992 635993 635994 635995 635996 635997 635998 635999 6359999 63599999 635999999

move start Click on a feature for details. Click on base position to zoom in around cursor. Click gray/blue bars on left for track options and descriptions. move end < [2.0] >

default tracks | hide all | add custom tracks | configure | refresh |

Chicken BLAT Results  
 Back Forward Reload Stop Home http://genome.ucsc.edu/cgi-bin/hgBlat

**UCSC BLAT search**

**Home Genomes Tables PCR Session FAQ Help**

**Chicken BLAT Results**

**BLAT Search Results**

ACTIONS	QUERY	SCORE	START	END	QSIZE	IDENTITY	CHRO	STRAND	START	END	SPAN
browser details	NP_001455.3	44	539	600	735	71.6%	Un	++	635370	635555	186
browser details	NP_001455.3	12	301	304	735	100.0%	1	++	67659709	67659720	12
browser details	NP_001455.3	12	437	440	735	100.0%	1	++	67660117	67660128	12
browser details	NP_001455.3	12	385	390	735	83.4%	1	++	67659961	67659978	18

**Alignment of NP\_001455.3**

Click on links in the frame to the left to navigate through the alignment. Matching bases mark the boundaries of gaps in either sequence.

**NP\_001455.3**

```

mrcvlfllsqg lgqlrmdnsf dslpwqityp ekirsiikieg ieqsasaykiv ieqkpytvnl 60
mgknflphnf rvvaysgsgp iapkldqqfqn fchyggiieg ypkavwvwt ctglrgylgf 120
envsyiglepl eassvgfehvi ygvkhkkadni slynekdieis rdifsfklgv epqddfafkyi 180
emhmvskgl ynhmgeddtv waqkvflilg ltnafivsfn itillesel widenkiatt 240
enwvclvklqkysl vqdgkakmavt qyqyvawlyl rffsdqckqy wyttdqkqy vqgkqy 300
tisleslavli lalnlyqsdllqsl tlyddinkcq csgawciminae ealikayki lancneefida 360
hfliakpksgc lhnqppldp fkqgavcvcpna kleageecdc gteodcalig etccodiactr 420
fkagancage pocencnlms kermcrpsfe ecldpcyedn ssascpcheny vtqghpogl 480
qzGKCLLck v gkfkillgpmr sgdkgctdf gkevefgpse cyshlnsktd vsgncgiads gyttqcead L 540
qzGKCLLck v gkfkillgpmr sgdkgctdf gkevefgpse cyshlnsktd vsgncgiads gyttqcead L 600
yqgkqy vqgkqy 660
fppvaparl perryienly hskpmrwppfli iflipffliifc vilainmvkvn fqirkkwrted 720
yssdqepsea sepkq

```

**Chicken.chrUn**

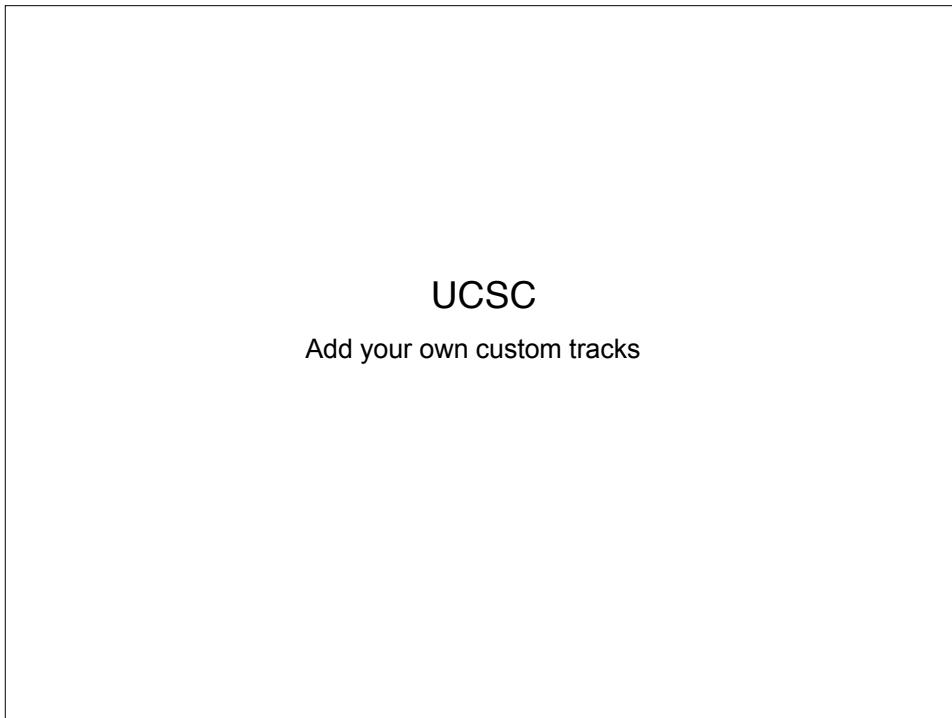
ATCGTGGATT GTGGAACACT CATCTGCCaca TA cccaaacg gggttccccctt caccggaaat 635429
 aasggt-GCA TCACTCTTGC Tcaagtgcga gagactCTCTC G ggtgtttt tggatgtatg 635489
 catgtccactt cggggcaca ta cccaaacg gggttccccctt gtt AGAGTC GcAcGaaATG CGGTccccggta 635549
 AMGGTA

**Side by Side Alignment\***

```

001615 N I Q C G K I I C K Y 001647
>>>>> | | Q | | C | | G | | K | | I | | I | | C | | K | | Y | >>>>
635370 aatctggccgtggaaaaactcatcgcacatac 635402

```



**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	genome	assembly	position or search term	image width	submit
Vertebrate	Human	Mar. 2006	chrX:151,073,054-151,383,976	620	<input type="button" value="submit"/>
<a href="#">Click here to reset the browser user interface settings to their defaults.</a> <a href="#">add custom tracks</a>   <a href="#">configure tracks and display</a>   <a href="#">clear position</a>					

**Add Custom Tracks**

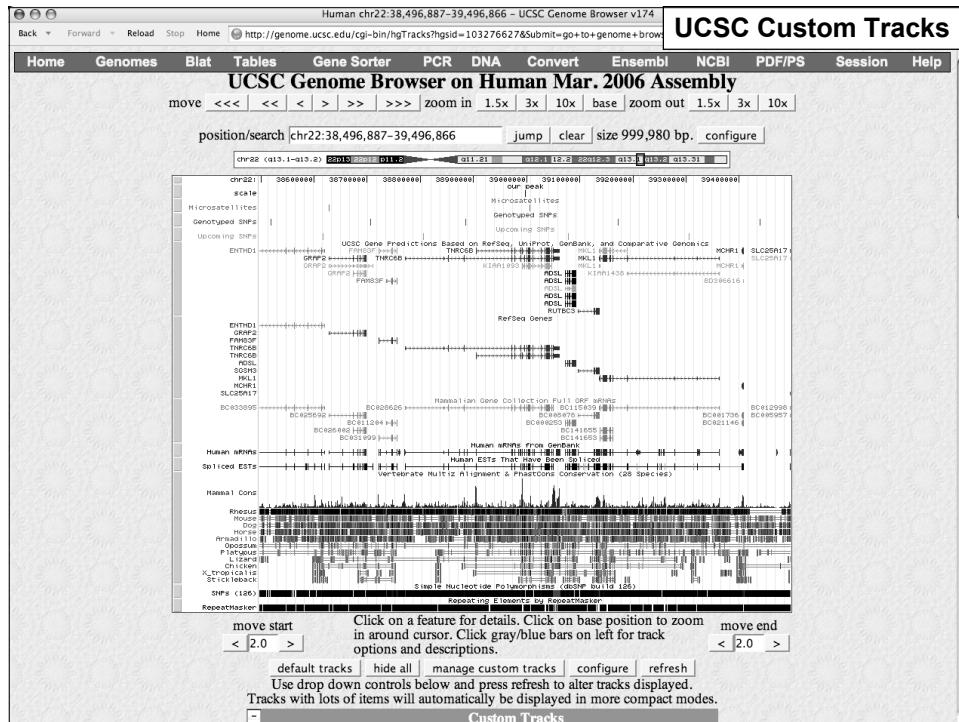
clade	genome	assembly	Mar. 2006	[hg18]
Display your own data as custom annotation tracks in the browser. Data must be formatted in BED, C display, set <a href="#">track</a> and <a href="#">browser</a> line attributes as described in the <a href="#">User's Guide</a> . Publicly available custom tr				
Paste URLs or data: Or upload: <input type="text"/> <input type="button" value="Browse..."/> <input type="button" value="Submit"/> <pre>track name="Microsatellites" description="Microsatellites" color=0,128,0 chr22 38627059 38627060 D22S276 chr22 39005417 39005418 D22S307 track name="Genotyped SNPs" description="Genotyped SNPs" color=0,0,255 chr22 38518342 38518343 ss146131 chr22 38705963 38705964 ss2941443 chr22 38884157 38884158 ss141110 chr22 39171391 39171391 ss22916</pre>				
Optional track documentation: <input type="text"/> <input type="button" value="Browse..."/> <input type="button" value="Clear"/>				

**Manage Custom Tracks**

Name	Description	Type	Doc Items	Pos	delete	add custom tracks
Upcoming SNPs	Upcoming SNPs	bed	4	chr22:	<input type="checkbox"/>	<a href="#">go to genome browser</a>
Genotyped SNPs	Genotyped SNPs	bed	5	chr22:	<input type="checkbox"/>	<a href="#">go to table browser</a>
Microsatellites	Microsatellites	bed	2	chr22:	<input type="checkbox"/>	
scale	our peak	bed	1	chr22:	<input type="checkbox"/>	
<input type="checkbox"/> check all / clear all <input type="button" value=""/>						<input type="button" value="-"/>

Click [here](#) for an HTML document template that

Nature Genetics: A user's guide to the human genome,  
 Question 7



## UCSC Table Browser

- Download track in text format
- Retrieve DNA sequence covered by a track
- Calculate intersections between tracks and view in the Genome Browser. For example:
  - Show all RefSeq genes that contain only one exon
  - Show transcription factor binding sites that overlap (intersect) with a SNP

**Table Browser**

Use this program to retrieve the data associated with a track in text format, to calculate intersections between track. For help in using this application see Using the Table Browser for a description of the controls in this form, sample queries, and the OpenHelix Table Browser tutorial for a narrated presentation of the software features and want to use Galaxy or our public MySQL server. Refer to the Credits page for the list of contributors and usage re

clade: Vertebrate    genome: Human    assembly: Mar. 2006

group: Genes and Gene Prediction Tracks    track: RefSeq Genes

table: refGene    describe table schema

region: \* genome position chr22:38496887-39496866    lookup    define regions

identifiers (names/acceessions): paste list    upload list

filter:

intersection:

correlation:

output format: all fields from selected table     Send output to Gala

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

file type returned: plain text    gzip compressed

[get output](#)    [summary/statistics](#)

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

**UCSC Table Browser: RefSeq genes that contain only one exon**

**Filter on Fields from hg18.refGene**

bin	is ignored	<input type="button" value="match"/>	AND
name	does	<input type="button" value="match"/>	AND
chrom	does	<input type="button" value="match"/>	AND
strand	does	<input type="button" value="match"/>	AND
txStart	is ignored	<input type="button" value="match"/>	AND
txEnd	is ignored	<input type="button" value="match"/>	AND
cdsStart	is ignored	<input type="button" value="match"/>	AND
cdsEnd	is ignored	<input type="button" value="match"/>	AND
exonCount	is =	<input type="button" value="1"/>	AND
exonStarts	does	<input type="button" value="match"/>	
exonEnds	does	<input type="button" value="match"/>	
id	is ignored	<input type="button" value="match"/>	AND
name2	does	<input type="button" value="match"/>	AND
cdsStartStat	does	<input type="button" value="match"/>	AND
cdsEndStat	does	<input type="button" value="match"/>	AND

#filter: refGene.exonCount = 1

bin	name	chrom	strand	txStart	txEnd	cdsStart	cdsEnd	exonCount	exonStarts	exonEnds	id	name2	cdsStartStat	cdsEndStat	exonFrames
5891	NM_0010054	chr1	+	58953	59971	58953	59871	1	58953,	59871,	OR4F5	cmpl	cmpl	cmpl	0,
5897	NM_0010052	chr1	+	357521	358458	357521	358458	1	357521,	358458,	OR4F3	cmpl	incmpl	cmpl	0,
5897	NM_0010052	chr1	+	357521	358458	357521	358458	1	357521,	358458,	OR4F16	cmpl	incmpl	cmpl	0,
5898	NM_0010052	chr1	+	610958	611897	610958	611897	1	610958,	611897,	OR4F16	cmpl	cmpl	cmpl	0,
5899	NM_0010052	chr1	-	610960	611897	610960	611897	1	610960,	611897,	OR4F29	incmpl	cmpl	cmpl	0,
5899	NM_0010052	chr1	-	610960	611897	610960	611897	1	610960,	611897,	OR4F3	incmpl	cmpl	cmpl	0,
5899	NM_0010052	chr1	-	610960	611897	610960	611897	1	610960,	611897,	OR4F16	incmpl	cmpl	cmpl	0,
5903	NM_000431	chr1	+	1297905	2929321	1297905	2929321	1	1297905,	2929321,	ACTR12	cmpl	cmpl	cmpl	0,
607	NM_080431	chr1	+	2927905	2929325	2928110	2929244	1	2927905,	2929325,	RSC1A1	cmpl	incmpl	cmpl	0,
88	NM_006511	chr1	+	15858950	15860803	15858950	15860803	1	15858950,	15860803,	LOC440567	cmpl	cmpl	cmpl	0,
707	NM_0010895	chr1	-	16006243	16006781	16006455	16006731	1	16006243,	16006781,					

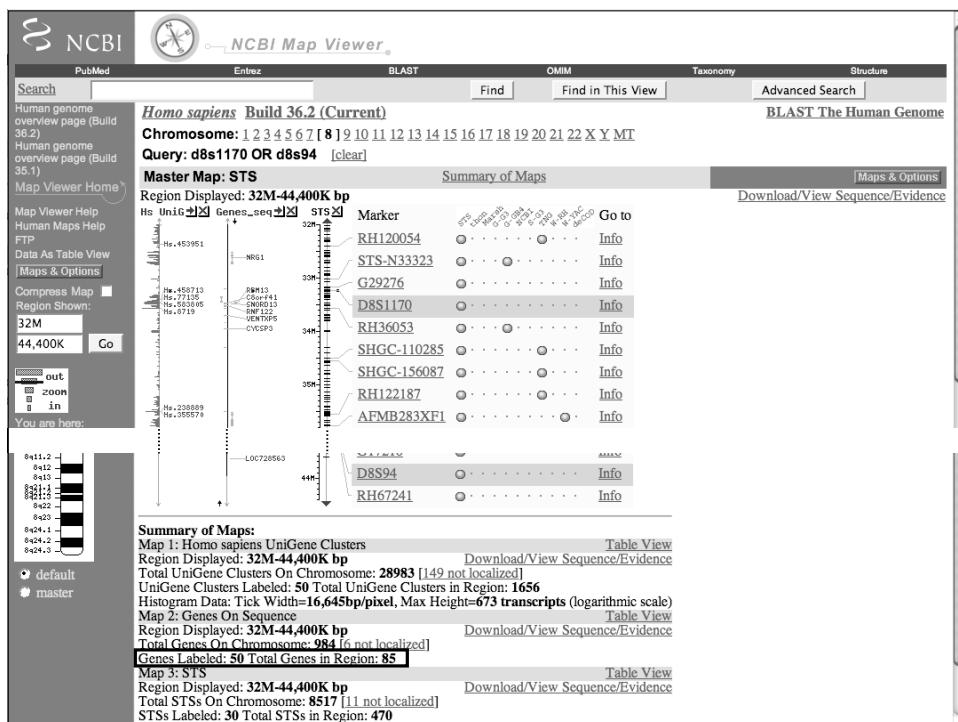
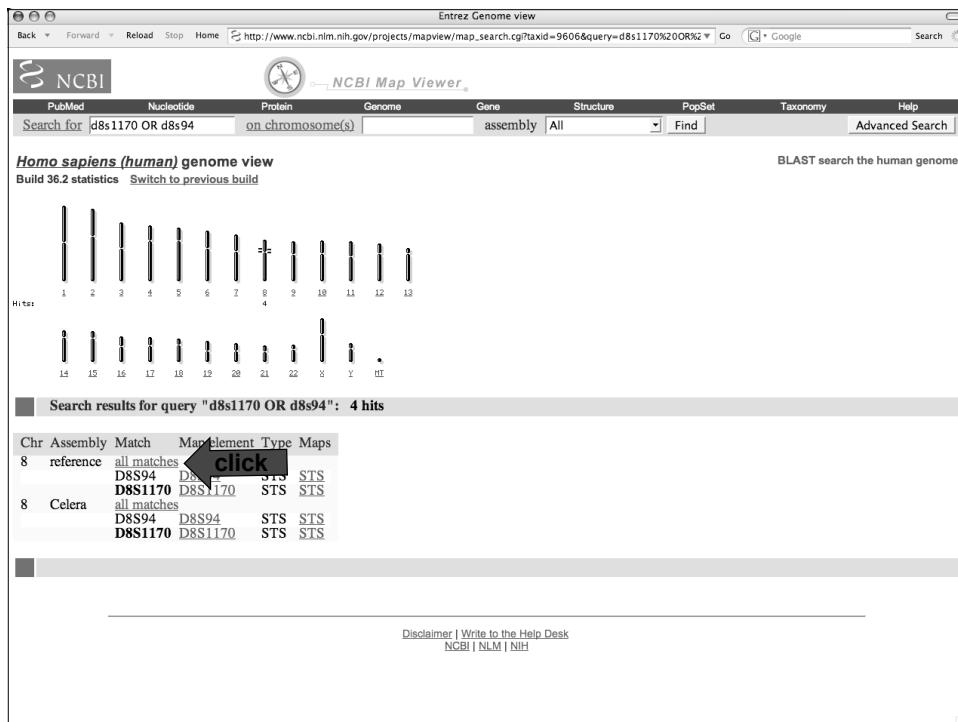
**NCBI**

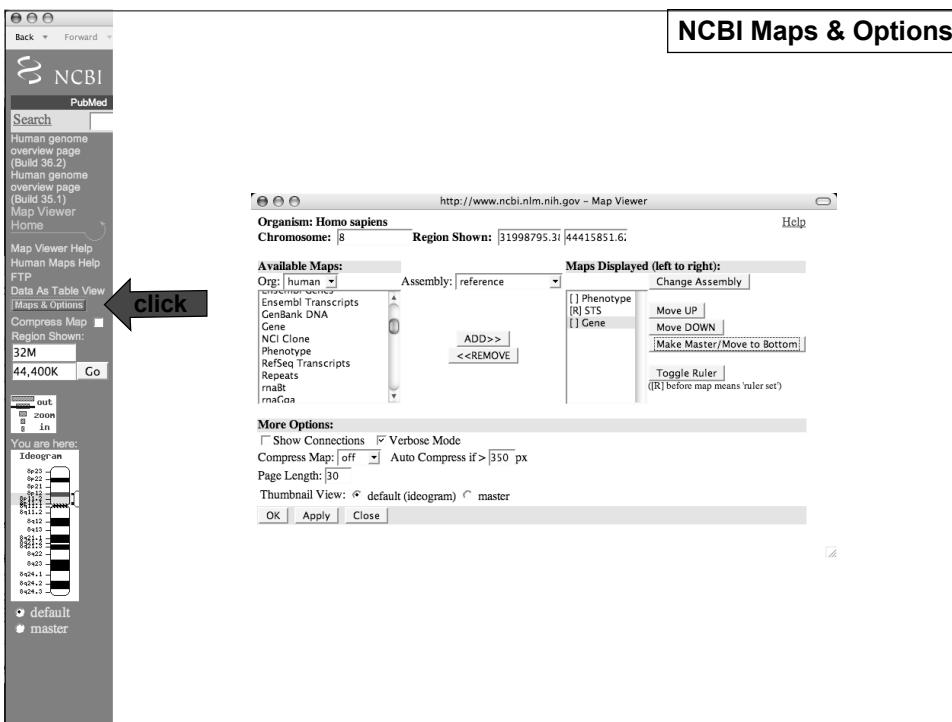
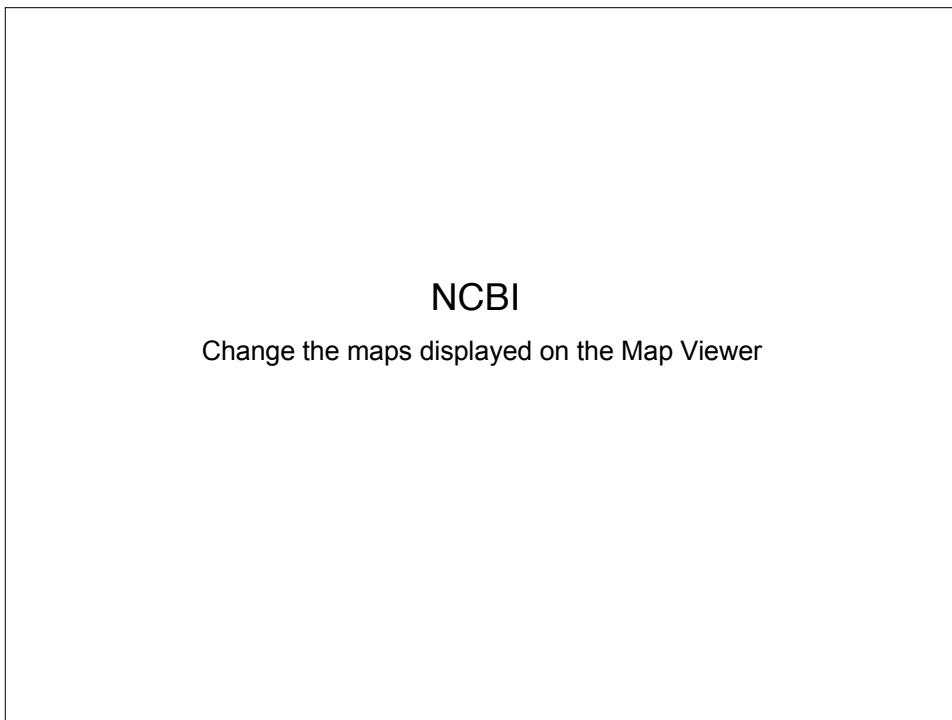
View a genomic region between two STS markers

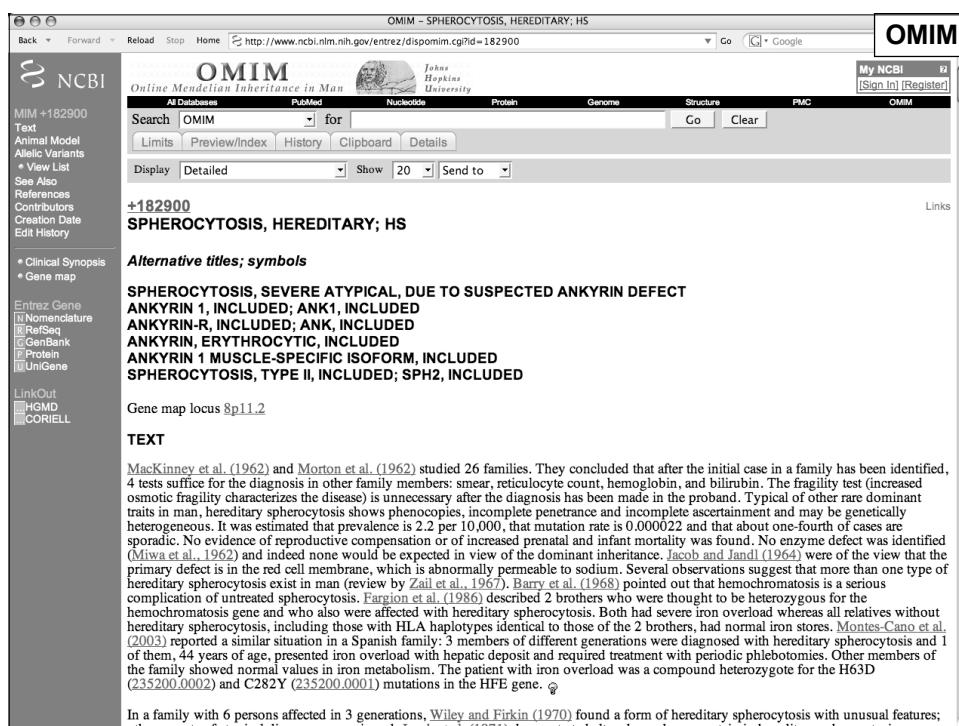
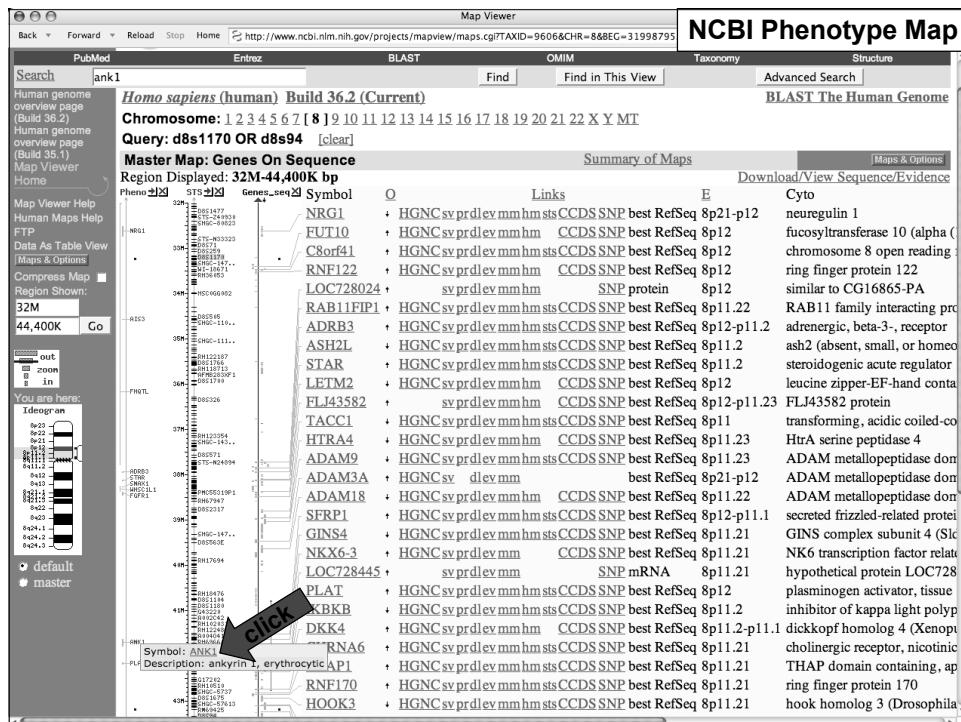
The screenshot shows the NCBI homepage with a sidebar on the left containing links to various databases and resources like PubMed, GenBank, and Molecular databases. The main content area features sections for 'What does NCBI do?' (Assembly Archive, Clusters of orthologous groups, etc.), 'Hot Spots' (GenBank 25th anniversary, Entrez Home, Entrez Tools, etc.), and 'PubMed Central' (an archive of biomedical and life sciences journals). A 'NCBI News' section is also present.

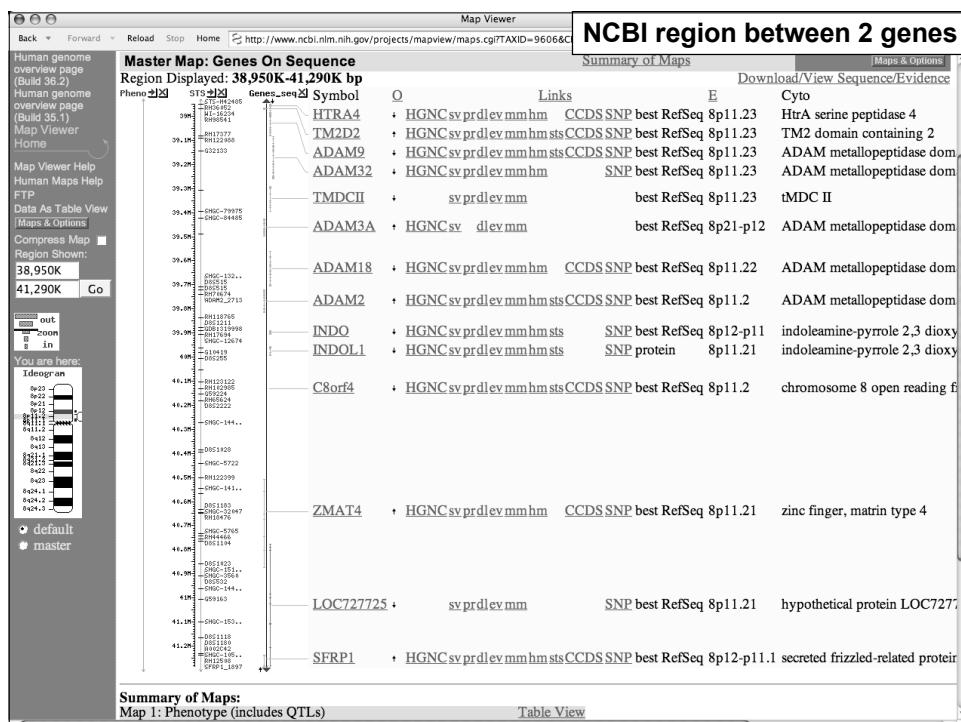
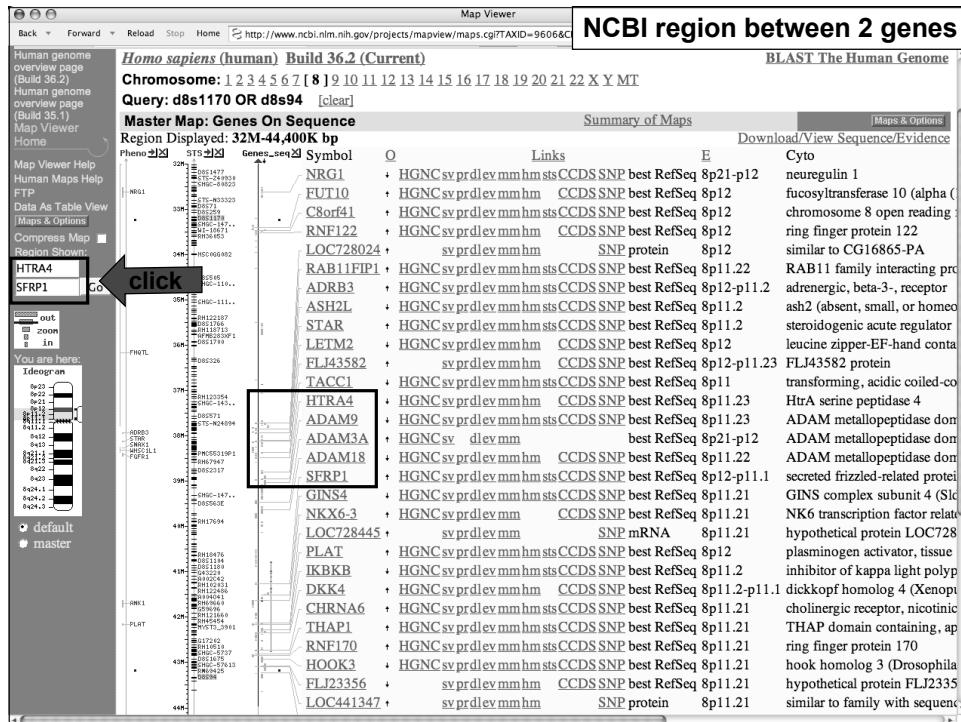
The screenshot shows the NCBI Map Viewer interface. It displays a hierarchical tree of genome mapping data for various organisms. The tree includes categories like Vertebrates, Mammals, Primates, Rodents, Monotremes, Marsupials, Other Mammals, Other Vertebrates, Invertebrates, Protozoa, Plants, and Fungi. Each node provides information about the build number and available tools.

Scientific name	Common name	Build	Tools
<i>Homo sapiens</i>	human	Build 36.2 Build 35.1	(15)
<i>Macaca mulatta</i>	rhesus macaque	Build 1.1	(13)
<i>Pan troglodytes</i>	chimpanzee	Build 2.1	(3)
<i>Mus musculus</i>	laboratory mouse	Build 37.1 Build 36.1	(2)
<i>Rattus norvegicus</i>	rat	RGSC v3.4	(2)
<i>Aspergillus fumigatus</i>		Build 2.1	(1)
<i>Aspergillus niger</i>		Build 1.1	(1)
<i>Candida glabrata</i>		Build 1.1	(6)
<i>Cryptococcus neoformans</i>		Build 2.1	(2)
<i>Debaromyces hansenii</i>		Build 1.1	(7)
<i>Encephalitozoon cuniculi</i>		Build 1.1	(7)
<i>Eremothecium gossypii</i>		Build 3.1	(42)
<i>Gibberella zeae</i>		Build 1.2	(16)
<i>Kluyveromyces lactis</i>		Build 1.1	(1)
<i>Magnaporthe grisea</i>	rice blast fungus	Build 3.1	(1)
<i>Neurospora crassa</i>		Build 1.1	(1)
<i>Pichia stipitis</i>		Build 1.1	(1)
<i>Saccharomyces cerevisiae</i>	baker's yeast	Build 2.1	(1)
<i>Schizosaccharomyces pombe</i>	fission yeast	Build 1.1	(1)
<i>Ustilago maydis</i>		Build 1.1	(1)
<i>Yarrowia lipolytica</i>		Build 1.1	(1)



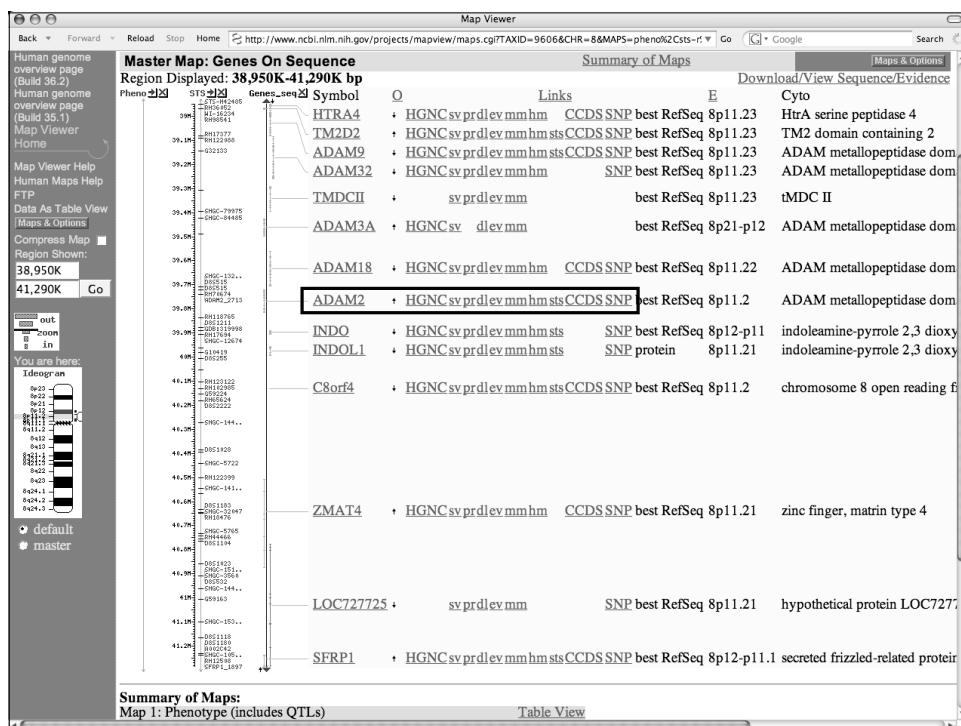
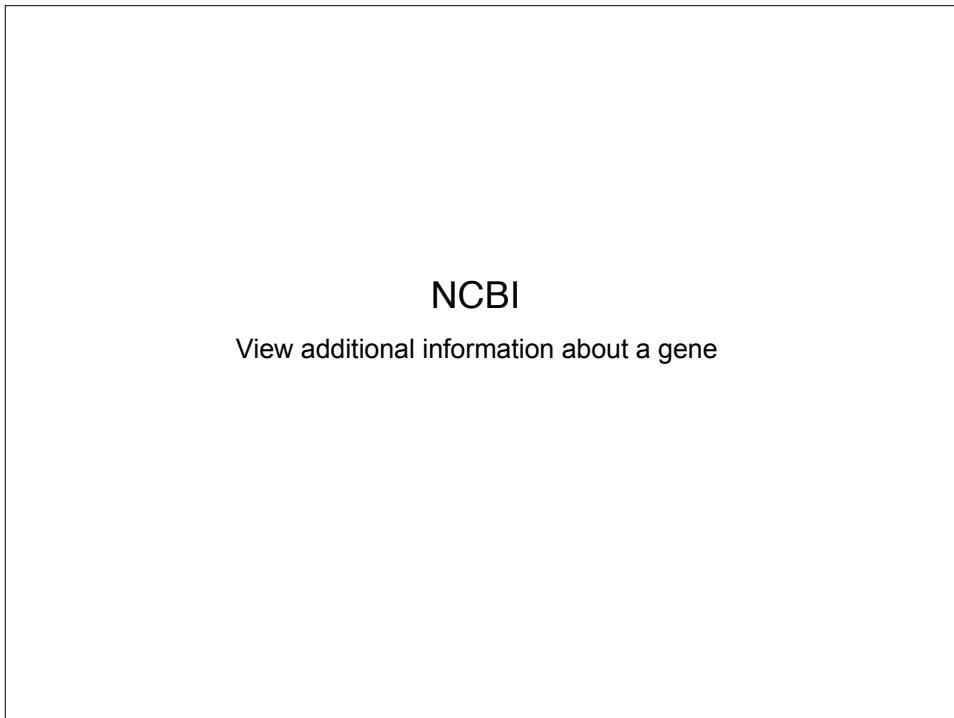






## NCBI

View additional information about a gene



Entrez Gene: ADAM2 ADAM metallopeptidase domain 2 (fertilin beta) [ Homo sapiens ]

Back Forward Reload Stop Home http://www.ncbi.nlm.nih.gov/sites/entrez?db=Gene&cmd=retrieve&dopt=full\_report&list\_uids=2515 Go

**Entrez Gene**

All Databases PubMed Nucleotide Protein Genome Structure PMC Taxonomy Books OMIM

Search Gene for Go Clear

Display Full Report Show 20 Send to

All: 1 Current Only: 1 Genes Genomes: 1 SNP GeneView: 1

1: ADAM2 ADAM metallopeptidase domain 2 (fertilin beta) [ Homo sapiens ] GeneID: 2515 updated 17-Jan-2008

**Summary**

Official Symbol ADAM2 provided by HGNC

Official Full Name ADAM metallopeptidase domain 2 (fertilin beta) provided by HGNC

Primary source HGNC:198

See related Ensembl:ENSG00000104755; HPRD:03322; MIM:601533

Gene type protein coding

RefSeq status Reviewed

Organism Homo sapiens

Lineage Eukarya; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Haplorhini; Catarrhini; Hominidae; Homo

Also known as FTNB; PH30; CRYN1; CRYN2; PH-30b

**Summary** This gene encodes a member of the ADAM (a disintegrin and metalloprotease domain) family. Members of this family are membrane-anchored proteins structurally related to snake venom disintegrins, and have been implicated in a variety of biological processes involving cell-cell and cell-matrix interactions, including fertilization, muscle development, and neurogenesis. This member is a subunit of an integral sperm membrane glycoprotein called fertilin, which plays an important role in sperm-egg interactions.

**Genomic regions, transcripts, and products**

(minus strand) Go to reference sequence details Try our new Sequence Viewer

NC\_000008\_9

Links Explain

- Order cDNA clone
- Conserved Domains
- Genome
- GEO Profiles
- HomoloGene
- Map Viewer
- CoreNucleotide
- EST
- Nucleotide
- OMIM
- Full text in PMC
- Probe
- Protein
- PubMed
- PubMed (GeneRIF)
- SNP
- SNP: Genotype
- SNP: GeneView
- Taxonomy
- UniSTS
- AceView
- CCDS
- Ensembl
- Evidence Viewer

Entrez Gene: ADAM2 ADAM metallopeptidase domain 2 (fertilin beta) [ Homo sapiens ]

Back Forward Reload Stop Home http://www.ncbi.nlm.nih.gov/sites/entrez?db=Gene&cmd=retrieve&dopt=full\_report&list\_uids=2515 Go

**NCBI Reference Sequences (RefSeq)**

RefSeqs maintained independently of Annotated Genomes

These reference sequences exist independently of genome builds. Explain

**mRNA and Protein(s)**

- NM\_001464.3→NP\_001455.3 ADAM metallopeptidase domain 2 protein**

Source sequence(s) BG71916, B1460477, U38805  
Consensus CDS CCDS34884.1  
UniprotKB/Swiss-Prot O99965  
Conserved Domains (5) summary

smart00050	DISIN; Homologues of snake disintegrins ; Snake disintegrins inhibit the binding of ligands to integrin receptors. They contain a 'RGD' sequence, identical to the recognition site of many adhesion proteins.
Location:39814936..39815000	A/C; ADAM Cysteine-Rich Domain;
Blast Score:281	ZnMc; Adamalysin_II like; Zinc-dependent metalloprotease; adamalysin_II_like
smart00608	subfamily ; Adamalysin_II is a snake venom zinc endopeptidase. This subfamily contains other snake venom metalloproteases, as well as membrane-anchored metalloproteases belonging to the ADAM family.
Location:472..609	ACR; Disintegrin; Disintegrin;
Blast Score:375	Location:178..372
cd04269	Pop_M12B_propep; Reprolysin family propeptide. This region is the propeptide for members of peptidase family M12B. The propeptide contains a sequence motif similar to the "cysteine switch" of the matrixins.
Location:39814936..39815000	Disintegrin; Disintegrin;
Blast Score:523	pfam00209
Location:39814936..39815000	Pop_M12B_propep; Reprolysin family propeptide. This region is the propeptide for members of peptidase family M12B. The propeptide contains a sequence motif similar to the "cysteine switch" of the matrixins.
pfam01562	
Location:2..141	
Blast Score:443	

**RefSeqs of Annotated Genomes: Build 36.2**

The following sections contain reference sequences that belong to a specific genome build. Explain

**Reference assembly**

**Genomic**

- NC\_000008.9 Reference assembly**

Range 39814936..39720411, complement  
Download GenBank FASTA Sequence Viewer (beta)
- NT\_007995.14**

Range 10016168..9921643, complement  
Download GenBank FASTA Sequence Viewer (beta)

**Alternate assembly (based on Celera assembly)**

**Genomic**

**HomoloGene Result**

**HomoloGene (hm)**

NCBI

HomoloGene - Discover Homologs

All Databases PubMed Nucleotides Protein Genome Structure Map Viewer Gene UniGene OMIM

Search HomoloGene for [Go] [Clear]

Limits Preview/Index History Clipboard Details

Display HomoloGene Show 20 Send to

All: 1 Fungi: 0 Mammals: 1

1: HomoloGene:1127. Gene conserved in Eutheria

Download, Links

**Genes**  
*Genes identified as putative homologs of one another during the construction of HomoloGene.*

ADAM2, *Homo sapiens*  
 ADAM metallopeptidase domain 2 (fertilin beta)

ADAM2, *Pan troglodytes*  
 ADAM metallopeptidase domain 2 (fertilin beta)

ADAM2, *Canis lupus familiaris*  
 ADAM metallopeptidase domain 2 (fertilin beta)

Adam2, *Mus musculus*  
 a disintegrin and metallopeptidase domain 2

Adam2, *Rattus norvegicus*  
 a disintegrin and metalloprotease domain 2

**Proteins**  
*Proteins used in sequence comparisons and their conserved domain architectures.*

NP\_001455.3  
 735 aa

XP\_519722.2  
 735 aa

XP\_532795.2  
 881 aa

NP\_033748.2  
 735 aa

NP\_064462.1  
 739 aa

**Alignment Scores**  
*Various evolutionary parameters derived from pairwise alignments have been saved.*

Show Table of Pairwise Scores

Alignments can be regenerated using BLAST for any selected pair of proteins.

Regenerate Alignments

NP\_001455.3 (Homo sapiens, ADAM2) XP\_519722.2 (Pan troglodytes, ADAM2)

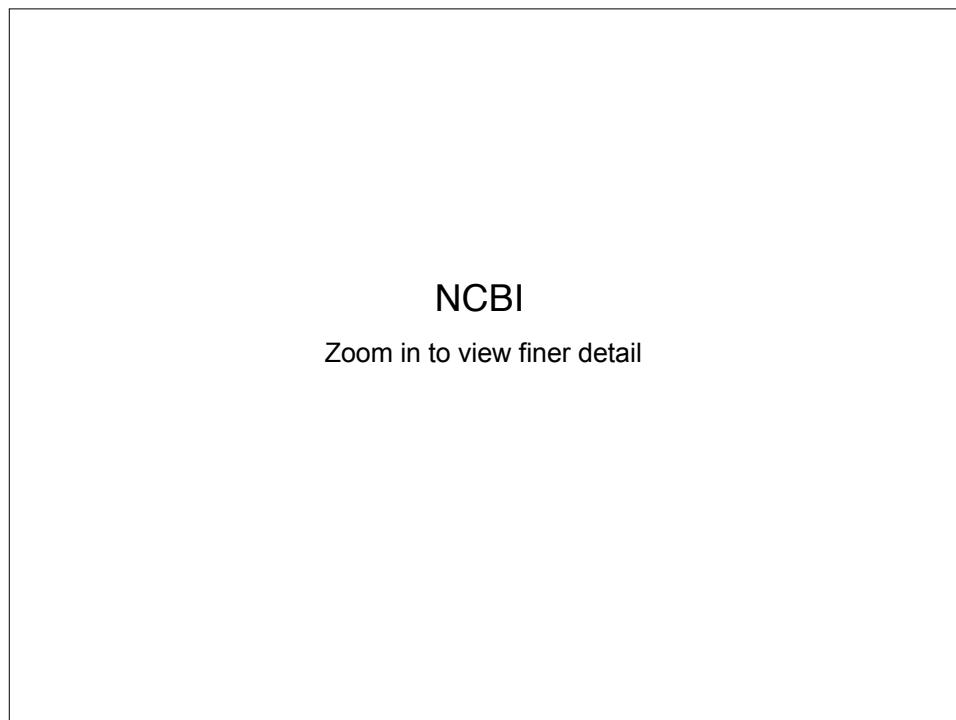
BLAST

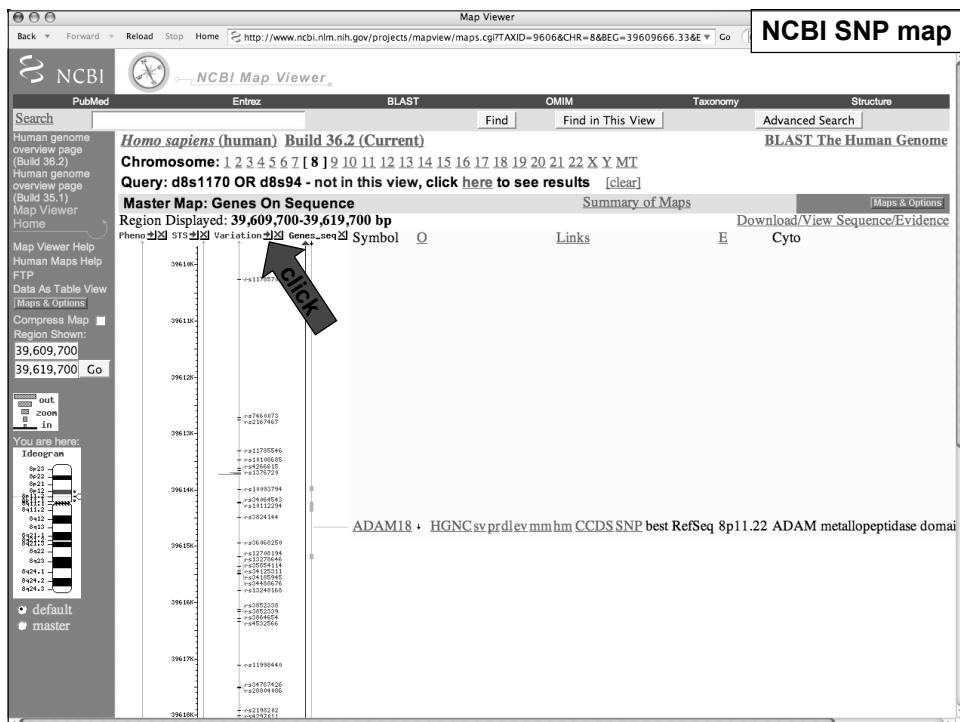
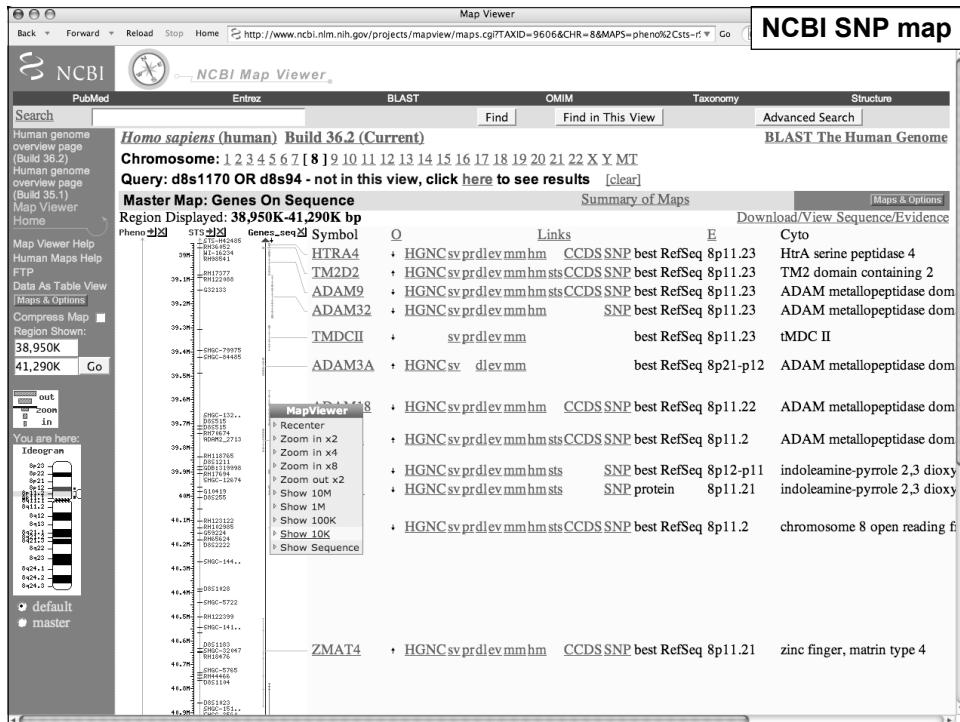
**Conserved Domains**  
*Conserved Domains from CDD found in protein sequences by rpsblast searching.*

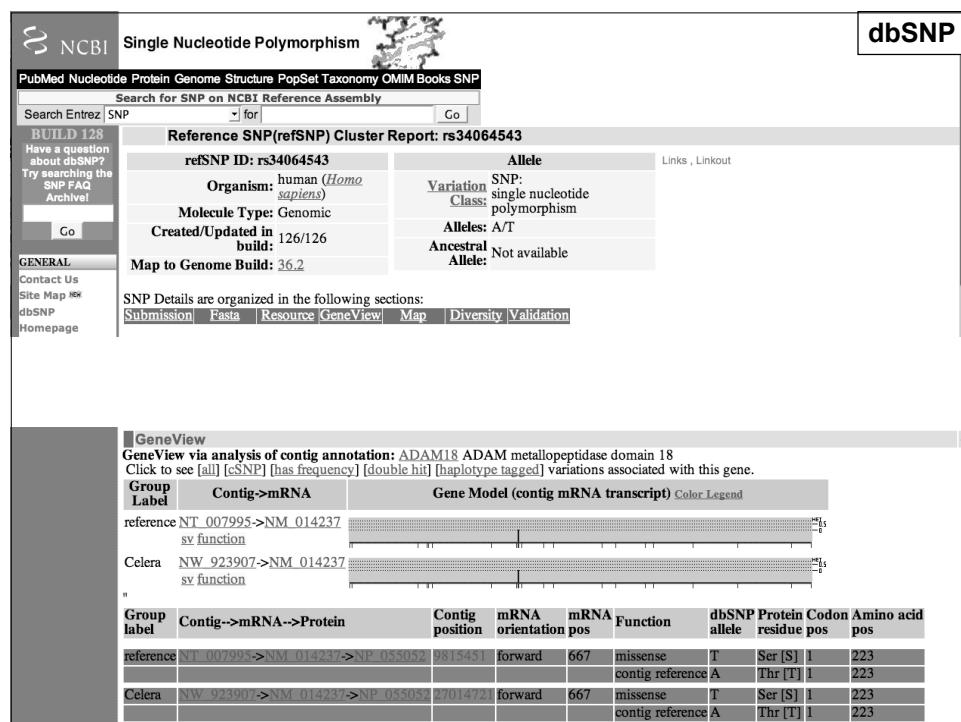
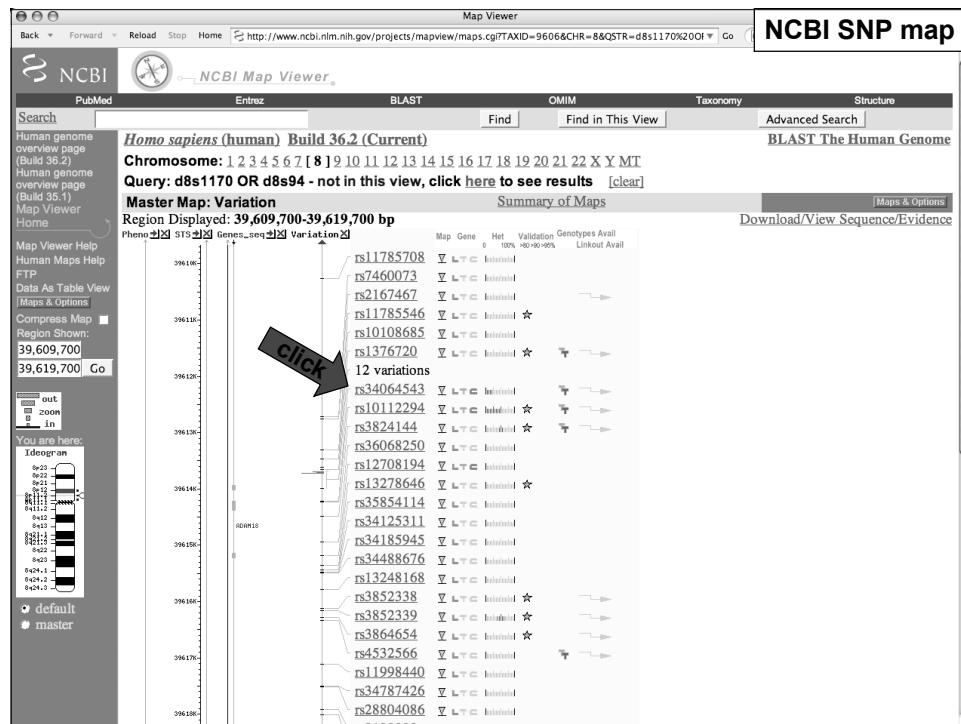
ZnMc\_adamalyisin\_II\_like (cd04269)  
 Zinc-dependent metalloprotease; adamalyisin\_II\_like subfamily. Adamalyisin II is a snake venom zinc endopeptidase. This subfamily contains other snake venom metalloproteinases, as well as membrane-anchored metalloproteinases belonging to the ADAM family.

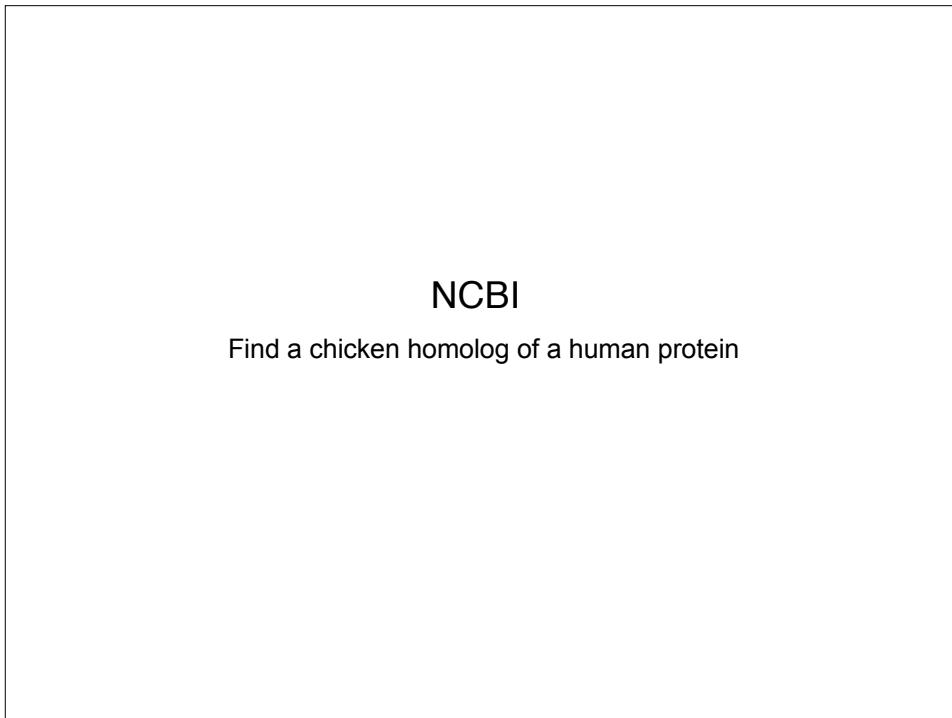
Pep\_M12B\_propep (pfam01562)  
 Reprolysin family propeptide. This region is the propeptide for members of peptidase family M12B. The propeptide contains a sequence motif similar to the "cysteine switch" of the matrixins.

ACR (smart00606)









BLAST Chicken Sequences. http://www.ncbi.nlm.nih.gov/genome/seq/BlastGen/BlastGen.cgi?taxid=9031

NCBI BLAST search

Search Map Viewer Go Clear

BLAST Overview FAQs News Manual References Retrieve results Genome Project

BLAST Chicken Sequences.

Enter an accession, gi, or a sequence in FASTA format:  
NP\_001455

Or, choose a file to upload Browse...

Set subsequence: (optional)  
From: To:

Database: genome reference only 17507 sequences

Program: tBLASTN: Compare a protein sequence against a nucleotide database

Optional parameters

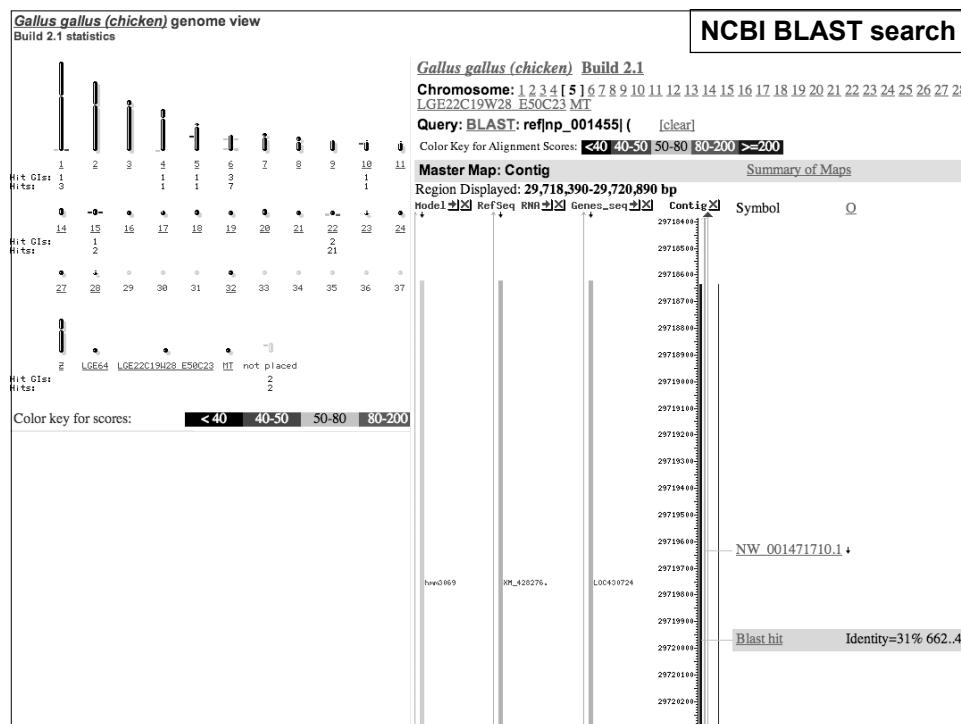
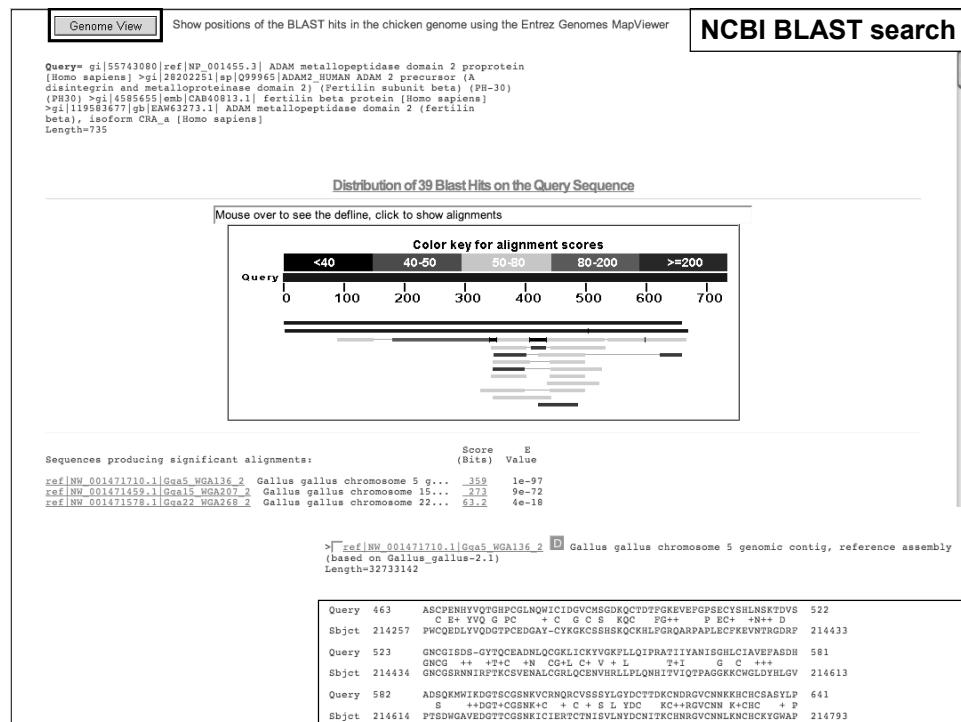
Expect Filter Descriptions Alignments  
0.01 low complexity 100 100

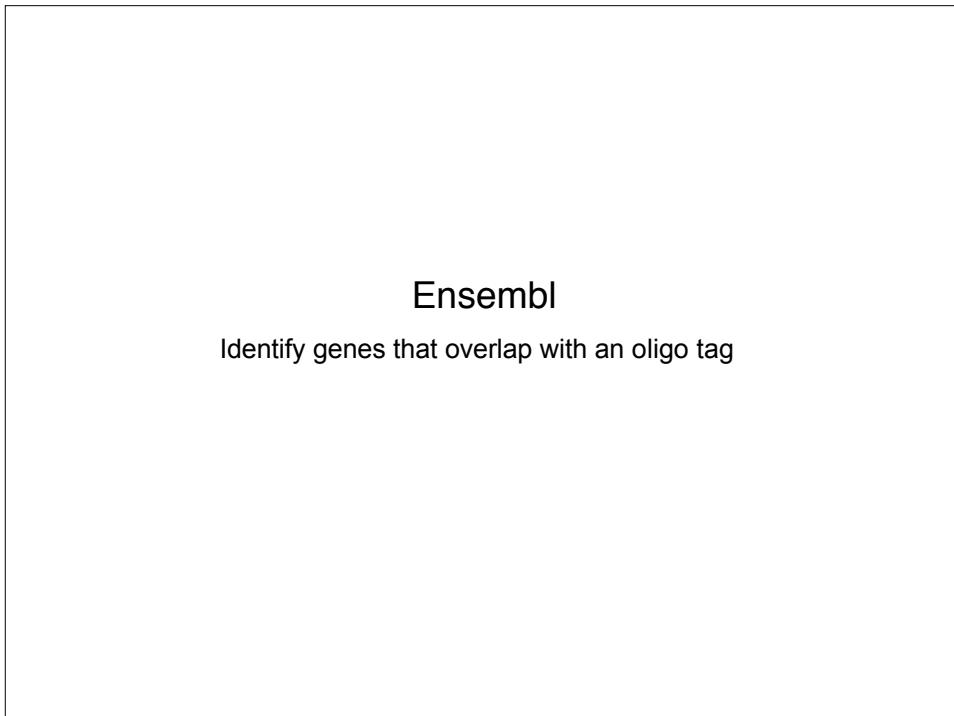
Advanced options:

Begin Search Clear Input

Get the URL with preset values? Get URL

A black arrow points from the text "select" to the dropdown menu labeled "Program: tBLASTN: Compare a protein sequence against a nucleotide database".





Ensembl Genome Browser

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e! Ensembl

Ensembl release 48 - Dec 2007

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Pika Ochotona princeps Now in Ensembl

Search Ensembl

Search: All species for e.g. mouse chromosome 2 or rat X:10000..20000 or human gene BRCA2

Ensembl tools

- Start a sequence search → click
- Search Ensembl for nucleotide or peptide sequences with BLAST and SSAHA.
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- Customise Your Ensembl → Register with Ensembl to bookmark your favourite pages, customise your home page and much more!
- Fetch data with the Ensembl API → Learn how to extract data from the public Ensembl database with this tutorial.

Ensembl 48 Pre species Popular genomes · Log in to customize

- Human NCBI 36 | Vega
- Mouse NCBI m37 | Vega
- Zebrafish Zv7 | Vega

All genomes -- Select a species -- Other pre-build species are available in Ensembl Prel →

Ensembl headlines: Release 48 (December 2007)

- e! MySQL upgrades on external databases (all species)
- Human variation updates (*Homo sapiens*)
- Updated Mouse variation data (*Mus musculus*)
- Updated Rat variation data (*Rattus norvegicus*)
- New species - American Pika (*Ochotona princeps*)

**BlastView**

**Ensembl BLAST search**

**e! Ensembl Multi BlastView**

Ensembl release 48 - Dec 2007

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Pika *Ochotona princeps*  
 Now in Ensembl!

Enter the Query Sequence

Either Paste sequences (max 30 sequences) in FASTA or plain text:  
>MPSS\_1  
AAAAATGTCGGCTGAAGAG

Or Upload a file containing one or more FASTA sequences

Or Enter a sequence ID or accession (EMBL, UniProt, RefSeq)

Or Enter an existing ticket ID:

dna queries  
 peptide queries

Select the databases to search against

Select species:  
 Use 'ctrl' key to select multiple species  
 Gallus\_gallus  
 Gasterosteus\_aculeatus  
 Homo\_sapiens

dna database  
 peptide database  
 Known Consensus CDS Peptides (CCDS peptides)

Select the Search Tool

BLASTN  
 BLAT  
 SSAHA2

(configure)

Search sensitivity:  
 Optimise search parameters to find the following alignments

Near-exact matches (oligo)  
 Exact matches  
 Near-exact matches  
 Near-exact matches (oligo)  
 Allow some local mismatch  
 Distant homologies  
 No optimisation

About BlastView  
 BlastView provides an integrated interface to both BLAST and Ensembl databases, offering access to both BLAST and Ensembl databases.

**BlastView**

**Ensembl BLAST search**

**Mouse Lemur *Microcebus murinus***  
 Now in Ensembl!

**100% identity over 100% of the query length**

Query Start	End	% ID
1	20	100.00

Alignment Locations vs. Query (click arrow to hide)

+ rep coverage  
>MPSS\_1

Alignment Summary (click arrow to hide)

Select rows to include in tables, and type of sort.  
 (Use the 'ctrl' key to select multiple.)

Links	Query Start	Chromosome	Supercontig	Clone	Contig	Chromosome	Stats	Sort By
Links	Start	Name	Start	Name	Start	Name	Score	Chromosome
Links	Start	Start	Start	Start	Start	Start	E-val	Off
Links	Start	End	Ori	Name	Start	End	Ori	Off
Links	Start	End	Ori	Name	Start	End	Ori	Score
Links	Start	End	Ori	Name	Start	End	Ori	%ID
Links	Start	End	Ori	Name	Start	End	Ori	Length
[A]	[S]	[G]	[C]	1	20	+	Chr15	54998168 54998187 +
[A]	[S]	[G]	[C]	1	20	+	Chr15	54998168 54998187 +
[A]	[S]	[G]	[C]	72205113	72205129	-	Chr8	72205113 72205129 -
[A]	[S]	[G]	[C]	72205113	72205129	-	Chr8	72205113 72205129 -

**click**

**Ensembl Human ContigView**

Ensembl release 48 - Dec 2007

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- View of Chromosome 15
- Graphical view
- Genomic overview
- Resequencing alignment
- View alignment with ...
- View alongside ...
- View Syntenic regions ...
- View region at UCSC
- View region at NCBI

Chromosome 15  
 54,996,168 - 55,000,187

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- Export SNP info in region
- Export Vega info in region

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Ensembl release 48: Homo sapiens Features on Chromosome 15 54996168-550000

http://www.ensembl.org/Homo\_sapiens/contigview?chr=15&vc\_start=54996168&vc\_end=55000

e.g. AL138722.15.1.44776, AL355340.17.1.112442

HOME · BLAST · BIOMART · SITEMAP · HELP · Search>>

Chr. 15

Chr. 15 band

DNA(contigs)

Markers

Ensembl Genes

mRNA Genes

Gene legend

TE39 SUHWA UN025 TCF12 CGN

ENST0000031725 > TCF12 > NP\_056920.1 >

AC000694.1.22.1966 > RP11-32F24

Reg. Features

Length

Common Known ProteinCoding EST gene Unpublished Promoter associated

**Ensembl ContigView**

Ensembl release 48: Homo sapiens Features on Chromosome 15 54996168-550000

http://www.ensembl.org/Homo\_sapiens/contigview?panel\_zoom=on:&v=15K4A54996168-550001

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Chr. 15

Chr. 15 band

DNA(contigs)

Human tispath clones

Reg. Features

Length

Common Known ProteinCoding EST gene Unpublished Promoter associated

AC000694.1.22.1966 > RP11-32F24

AC000694.1.22.1966 > RP11-32F24

Reg. Features

Length

Common Known ProteinCoding EST gene Unpublished Promoter associated

AC000694.1.22.1966 > RP11-32F24

Amino acids

Sequence

DNA(contigs)

Human tispath clones

Amino acids

Sequence

DNA(contigs)

Human tispath clones

Reg. Features

Length

Common Known ProteinCoding EST gene Unpublished Promoter associated

Ensembl Homo sapiens version 48.36 (NCBI 36) Chromosome 15 54,996,168 - 55,000,187

Basepair view

Chr. 15

Chr. 15 band

DNA(contigs)

Human tispath clones

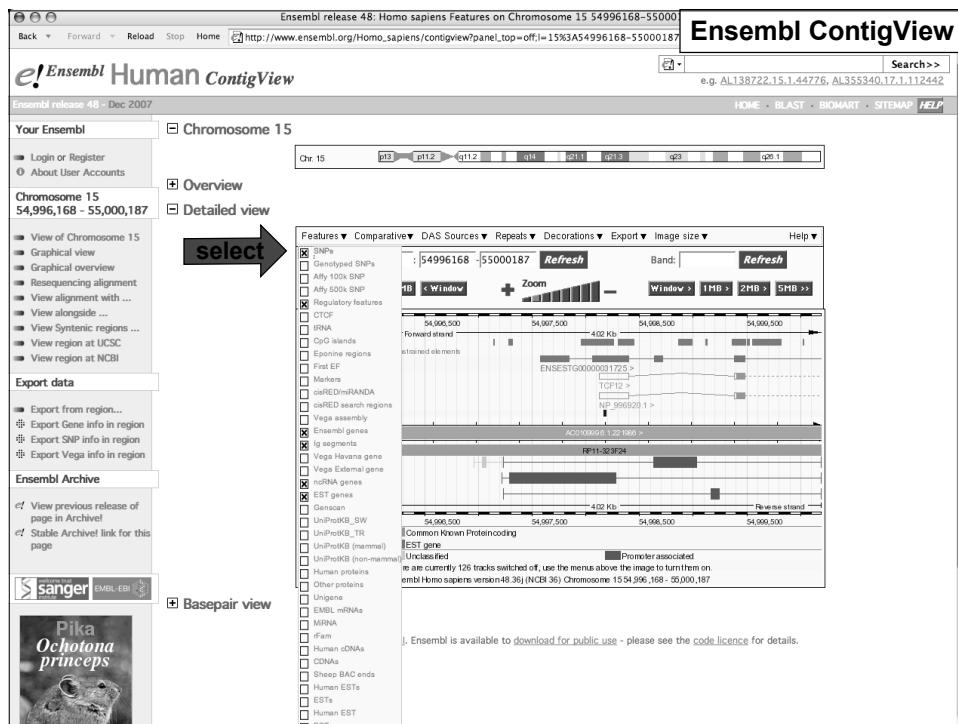
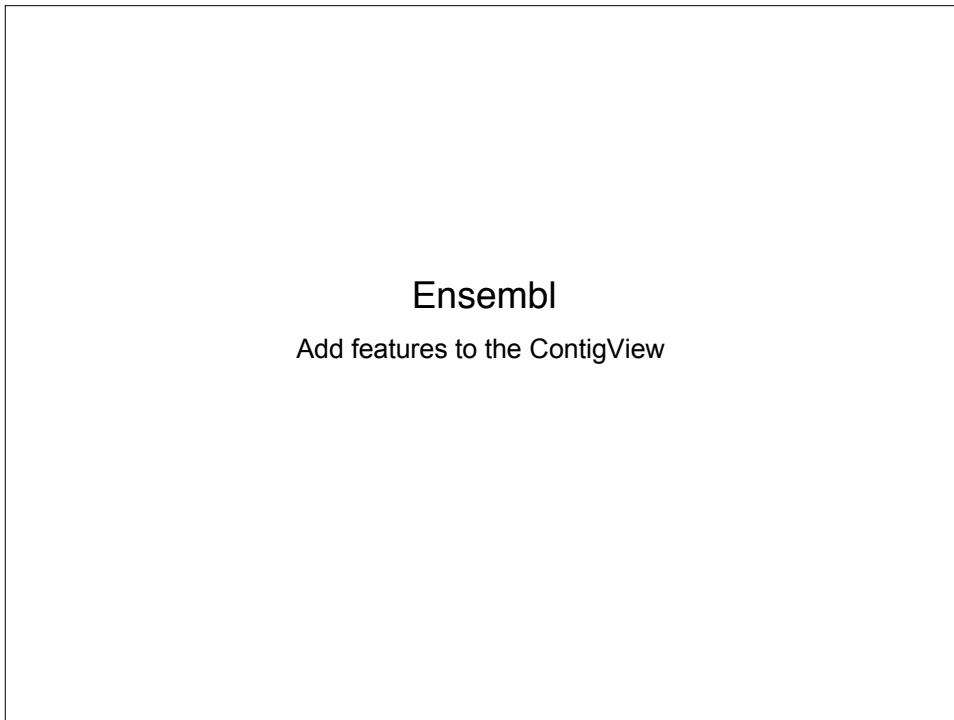
Reg. Features

Length

Common Known ProteinCoding EST gene Unpublished Promoter associated

Ensembl Homo sapiens version 48.36 (NCBI 36) Chromosome 15 54,996,168 - 55,000,187

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**Ensembl ContigView**

Ensembl release 48: Homo sapiens Features on Chromosome 15 54996168-55000187

http://www.ensembl.org/Homo\_sapiens/contigview?panel\_top=off,!=15%3A54996168-55000187;h=BLAST\_NEW%3ABLA\_K

e! Ensembl Human ContigView

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Chromosome 15 54,996,168 - 55,000,187

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- View alignment with ...
- View alongside ...
- View synteny regions ...
- View region at UCSC
- View region at NCBI

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- Export Gene info in region
- Export SNP info in region
- Export Vega info in region

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Mouse Lemur *Microcebus murinus*  
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Basepair view

Ensembl ContigView

Chromosome 15

Chr 15 p11.2 q11.2 q15 q21.3 q23 q26.1

Features Comparative DAS Sources Repeats Decorations Export Image size Help

Jump to region 15 54996168-55000187 Refresh Band: Refresh

<< 5MB < 2MB < 1MB < Window Zoom Window > 1MB > 2MB > 5MB >

Chr 15 length Forward strand Constraints elements 402 Kb 54,996,500 54,997,500 54,998,500 54,999,500

Conservation EST trans. Ensembl transcripts Blast hits DNA(contigs) Human telepath clones SNPs Reg. Features Length

Reg. Features Length

Gene legend Common Known Protein coding EST gene SNP legend Upstream Intron Reg. feats legend Unclassified Promoter associated

Length

SNP: rs12915002 SNP properties bp: 54999139 class: SNP ambiguity code: M alleles: C/A source: dbSNP type: Intron

There are currently 125 tracks switched off, use the menu above the image to turn them on.

Ensembl Homo sapiens version 48 (NCBI 36) Chromosome 15 54,996,168 - 55,000,187

click

**Ensembl SNPView**

SNPview

Back Forward Reload Stop Home http://www.ensembl.org/Homo\_sapiens/snpview?snp=rs12915002;source=dbSNP;c=15:54999139

54,999,139

Flanking sequence

SNP rs12915002 is located in the following transcripts

Genomic location (strand)	Gene	Transcript relative SNP position	Translation relative SNP position	Type	GeneSNPView
15: 54999139-54999139 (+)	ENSG00000140262	ENST00000333725: n/a	ENSP00000331057: n/a	INTRONIC	SNPs in gene context
	ENSG00000140262	ENST00000267811: n/a	ENSP0000267811: n/a	INTRONIC	SNPs in gene context

Population genotypes and allele frequencies

This SNP has no allele or genotype frequencies per population.

Individual genotypes for SNP rs12915002

SNP Context - 15 54999139

Features Source SNP class SNP type Decorations Export Image size Help

No ncRNAs in the region No Ensembl transcripts in this region

No Ensembl transcripts in this region

No Ensembl transcripts in this region

No Ensembl transcripts in this region

No EST transcripts in this region

8 of the 105 variations in this region have been filtered out by the Source, Class and Type menus.

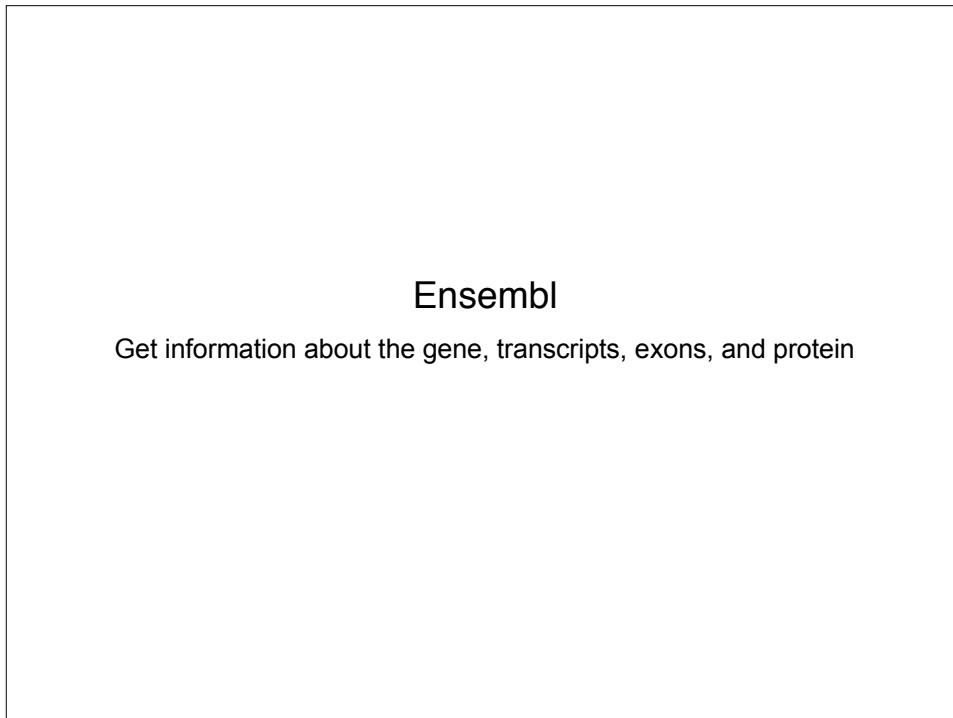
SNPs

Germlined SNPs

SNP legend

Intergenic Upstream Intron

54.99 Mb 55.00 Mb 55.01 Mb



**Ensembl ContigView**

Ensembl release 48: Homo sapiens Features on Chromosome 15 54996168-55000

Human ContigView

Chromosome 15 54,996,168 - 55,000,187

Overview Detailed view

Features Comparative DAS Sources Repeats Decorations Export Image size Help

Jump to region Chr 15 54,996,168 - 55,000,187 Refresh Band: Refresh

<< 5MB < 2MB < 1MB < Window Zoom Window > 1MB > 2MB > 5MB >>

Chr 15 Length Forward strand 54,996,500 54,997,500 54,998,500 54,999,500

Constrained elements ENST000000031725 > ENSP00000140262 Gene:ENSG00000140262 Transcript:ENST00000333725

EST trans. TCP10 > Ensembl Gene RefSeq\_protein NP\_996920.1 Gene:ENSG00000140262 Transcript:ENSP00000331057

Ensembl transcripts

Blast hits DNA(contigs) AC0000005 Human depth clones RPT11\_32.8

SNPs

Reg. Features Length 4.02 Kb Reverse strand

Reg. Features Length 4.02 Kb Reverse strand

Gene legend Common Known Protein coding Gene

SNP legend SNP gene Intergenic

Reg. Feature legend Unclassed Promoter associated

There are currently 125 tracks switched off, use the menus above the image to turn them on.

Ensembl Homo sapiens version 48.36 (NCBI 36) Chromosome 15 54,996,168 - 55,000,187

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ENSG00000140262

- Gene information
- Gene regulation info.
- Genomic sequence
- Genomic sequence alignment
- Resequencing alignment
- Gene splice site image
- Gene tree info.
- Gene variation info.
- ID info
- ID history
- Compare SNPs in transcript
- Transcript information
- Exon information
- Protein information
- Export gene data

Chromosome 15  
54,998,125 - 55,368,004

- View of Chromosome 15
- Graphical view
- Graphical overview
- Export from region...
- Export Gene info in region
- Export SNP info in region
- Export Vega info in region

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Ensembl release 48: Homo sapiens Gene report for ENSG00000140262

Gene TCF12 (HGNC Symbol) Synonyms: HEB, HsT17266, HTF4 This gene is a member of the Human CCDS set: CCDS10159, CCDS10160

Ensembl Gene ID ENSG00000140262

Genomic Location This gene can be found on Chromosome 15 at location 54,998,125-55,368,004. The start of this gene is located in Contig AC010999.6.1.221986.

Description Transcription factor 12 (Transcription factor HTF-4) (E-box-binding protein) (DNA-binding protein HTF4). Source: UniProt/SwissProt Q99Q81

Prediction Method Gene containing both Ensembl predicted transcripts and Havana manual annotation

Transcripts

TCF12	ENST00000267811	ENSP00000267811	[Transcript info]	[Exon info]	[Peptide info]
NP_996920_1	ENST00000333725	ENSP00000331057	[Transcript info]	[Exon info]	[Peptide info]
NP_996923_1	ENST00000343827	ENSP00000342459	[Transcript info]	[Exon info]	[Peptide info]

Features ▾

Alignments

This gene can be viewed in genomic alignment with other species

- view genomic alignment with 7 eutherian mammals Pecan
- view genomic alignment with 10 amniota vertebrates Pecan
- view genomic alignment with Rattus norvegicus
- view genomic alignment with Macaca mulatta
- view genomic alignment with Loxodonta africana
- view genomic alignment with Oryctolophus tenuirostris
- view genomic alignment with Dasyprocta novemcincta
- view genomic alignment with Canis familiaris
- view genomic alignment with Pan troglodytes
- view genomic alignment with Gallus gallus
- view genomic alignment with Ornithodorhynchus anatinus
- view genomic alignment with Bos taurus

click

**Ensembl AlignSliceView**

Ensembl release 48 - Dec 2007

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Chromosome 15  
54,998,125 - 55,368,004

- View of Chromosome 15
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- Resequencing alignment
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- View alongside ...
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- View region at UCSC
- View region at NCBI

Export data

- Export from region...
- Export Gene info in region
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Mouse Lemur *Microcebus murinus*

Ensembl Human AlignSliceView

Chromosome 15

Overview

Detailed view

Features ▾ Comparative ▾ Repeats ▾ Decorations ▾ Export ▾ Image size ▾ Help ▾

Jump to region 15 : 54998125 - 55368004 Refresh

Zoom Window 3 1MB 2MB 5MB

Homo\_sapiens

BEST tran.

Ensembl tran.

TCF12 Common Known Protein coding

DNA(cot�)

rRNA

Gene\_familiaris

Ensembl tran.

ENSCAF00000025746 Ensembl Novel Protein Coding

DNA(cot�)

Ensembl tran.

ENSCAF10000000579 Ensembl Novel Protein Coding

ENSCAF10000000579 Ensembl Novel Protein Coding

Basepair view

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**Ensembl GeneView**

Ensembl release 48: Homo sapiens Gene report for ENSG00000140262

Orthologue Prediction

The following gene(s) have been identified as putative orthologues:  
 (N.B. If you don't find a homologue here, it may be a 'between-species paralogue'. Please view the [gene tree info](#) or export between-species paralogues with Biomart to see more.)

Species	Type	Gene identifier
<i>Aedes aegypti</i>	1-to-many	<a href="#">AAFL010226</a> (Novel Ensembl prediction)
<i>Anopheles gambiae</i>	1-to-many	<a href="#">AGAP008114</a> (Novel Ensembl prediction)
<i>Bos taurus</i>	1-to-1	<a href="#">ENSBTA0000000002586</a> (ADJN84_BOVIN)
<i>Caenorhabditis elegans</i>	1-to-many	<a href="#">MO53S_5</a> (Nhr-2)
<i>Canis familiaris</i>	1-to-1	<a href="#">ENSCAFG00000016200</a> (TCF12)
<i>Cavia porcellus</i>	1-to-1	<a href="#">ENSCCP00000016200</a> (TCF12)
<i>Ciona intestinalis</i>	1-to-many	<a href="#">ENSGING00000009523</a> (Q4H3N7_CION)
<i>Ciona savignyi</i>	1-to-many	<a href="#">ENSGS000000011705</a> (Novel Ensembl prediction)
<i>Danio rerio</i>	1-to-1	<a href="#">ENSDARG00000004714</a> (zgc:85956)
<i>Dasyurus novemcinctus</i>	1-to-1	<a href="#">ENSDNOG00000003842</a> (TCF12)
<i>Drosophila melanogaster</i>	1-to-many	<a href="#">CG5102 (da)</a>
<i>Echinops telfairi</i>	1-to-1	<a href="#">ENSEETL000000016200</a> (TCF12)
<i>Erinaceus europaeus</i>	1-to-1	<a href="#">ENSEELU00000002182</a> (TCF12)
<i>Felis catus</i>	1-to-1	<a href="#">ENSEFCAG0000001867</a> (TCF12)

Transcription factor 12 (Transcription factor HTF-4) (E-box-binding protein) (DNA-binding protein HTF4). [Source:Uniprot/SwISSPROT, Acc:099081]  
 Transcription factor 12 (Transcription factor HTF-4) (E-box-binding protein) (DNA-binding protein HTF4). [Source:Uniprot/SwISSPROT, Acc:099081]  
 Transcription factor 12 (Transcription factor HTF-4) (E-box-binding protein) (DNA-binding protein HTF4). [Source:Uniprot/SwISSPROT, Acc:099081]  
 Transcription factor 12 (Transcription factor HTF-4) (E-box-binding protein) (DNA-binding protein HTF4). [Source:Uniprot/SwISSPROT, Acc:099081]  
 Transcription factor 12 (Transcription factor HTF-4) (E-box-binding protein) (DNA-binding protein HTF4). [Source:Uniprot/SwISSPROT, Acc:099081]  
 Transcription factor 12 (Transcription factor HTF-4) (E-box-binding protein) (DNA-binding protein HTF4). [Source:Uniprot/SwISSPROT, Acc:099081]  
 Transcription factor 12 (Transcription factor HTF-4) (E-box-binding protein) (DNA-binding protein HTF4). [Source:Uniprot/SwISSPROT, Acc:099081]  
 Transcription factor 12 (Transcription factor HTF-4) (E-box-binding protein) (DNA-binding protein HTF4). [Source:Uniprot/SwISSPROT, Acc:099081]  
 Transcription factor 12 (Transcription factor HTF-4) (E-box-binding protein) (DNA-binding protein HTF4). [Source:Uniprot/SwISSPROT, Acc:099081]

**Ensembl GeneView**

Ensembl release 48: Homo sapiens Gene report for ENSG00000140262

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ENSG00000140262

- Gene information
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- Genomic sequence alignment
- Resequencing alignment
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- Gene tree info.
- Gene variation info.
- LD info
- ID history
- Compare SNPs in transcript
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- Protein information
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Chromosome 15  
54,998,125 - 55,368,004

- View of Chromosome 15
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Ensembl Gene Report for ENSG00000140262

Gene TCF12 (HGNC Symbol) **Synonyms:** HEB, HsT17266, HTF4  
 This gene is a member of the Human CCDS set: [CCDS10159](#), [CCDS10160](#)

Ensembl Gene ID ENSG00000140262

Genomic Location This gene can be found on Chromosome 15 at location 54,998,125-55,368,004.  
 The start of this gene is located in Contig AC01099\_6.1\_221986.

Description Transcription factor 12 (Transcription factor HTF-4) (E-box-binding protein) (DNA-binding protein HTF4). [Source:Uniprot/SwISSPROT](#) 099081

Prediction Method Gene containing both Ensembl predicted transcripts and Havana manual annotation

Transcripts

Gene	ENSG00000140262	ENSG00000140262	ENSG00000140262	ENSG00000140262	ENSG00000140262
TCF12	ENST00000267811	ENSP00000267811	[Transcript info]	[Exon info]	[Peptide info]
NP_996920_1	ENST0000033725	ENSP00000331057	[Transcript info]	[Exon info]	[Peptide info]
NP_996923_1	ENST0000034527	ENSP00000342459	[Transcript info]	[Exon info]	[Peptide info]

Features ▾

Chr 15  
Length 35.00 Mb - 55.30 Mb  
Forward strand 36.88 Kb  
Reverse strand 35.20 Mb - 55.30 Mb

Ensembl transcripts  
TCF12 > Common Known Protein coding  
NP\_996920\_1  
Common Known Protein coding

DNA contigs  
AC01099\_6.1\_221986 > AC01099\_6.1\_19850  
Length 35.00 Mb - 55.30 Mb  
Reverse strand

Alignments

This gene can be viewed in genomic alignment with other species  
[view\\_genomic\\_alignment\\_with\\_Z\\_eutherian\\_mammals\\_Pecan](#)  
[view\\_genomic\\_alignment\\_with\\_Bivalvia\\_molluscs\\_Pecan](#)  
[view\\_genomic\\_alignment\\_with\\_Rattus\\_norvegicus](#)  
[view\\_genomic\\_alignment\\_with\\_Macaca\\_mulatta](#)  
[view\\_genomic\\_alignment\\_with\\_Loxodonta\\_africana](#)  
[view\\_genomic\\_alignment\\_with\\_Echinospermum\\_telfairi](#)  
[view\\_genomic\\_alignment\\_with\\_Oryctolagus\\_cuniculus](#)  
[view\\_genomic\\_alignment\\_with\\_Dasyurus\\_novemcinctus](#)  
[view\\_genomic\\_alignment\\_with\\_Cavia\\_familiaris](#)  
[view\\_genomic\\_alignment\\_with\\_Pan\\_troglodytes](#)  
[view\\_genomic\\_alignment\\_with\\_Gallus\\_gallus](#)  
[view\\_genomic\\_alignment\\_with\\_Omniorthorhynchus\\_anatinus](#)  
[view\\_genomic\\_alignment\\_with\\_Bos\\_taurus](#)

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**ENST00000267811**

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**Chromosome 15**  
54,998,125 - 55,368,004

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**Mouse Lemur**

**Ensembl Transcript Report**

**Transcript** TCF12 (HGNC Symbol) **Synonyms:** HEB, HsT17266, HTF4  
This transcript is a member of the Human CCDS set: CCDS10159  
Havana transcript having same CDS: OTTHUMT00000255069

**Ensembl Transcript ID** ENST00000267811

**Transcript information** Exons: 20 Transcript length: 4,712 bps Translation length: 682 residues  
This transcript is a product of gene: ENSG00000140262

**Genomic Location** This transcript can be found on Chromosome 15 at location 54,998,125-55,368,004.  
The start of this transcript is located in Contig AC01099.6.1.221986.

**Description** Transcription factor 12 (Transcription factor HTF-4) (E-box-binding protein) (DNA-binding protein HTF4). Source: Uniprot/SwissProt 029081

**Prediction Method** Transcript where the Ensembl predicted transcript and Havana manually annotated transcript have at least the same CDS

**Alternate transcripts** This Ensembl\_havana\_transcript entry corresponds to the following database identifiers:  
Havana transcript having same CDS: OTTHUMT00000255069

**Similarity Matches** This Ensembl\_havana\_transcript entry corresponds to the following database identifiers:  
**HGNC Symbol:** TCF12 transcription factor 12 (HTF4, helix-loop-helix transcription factors 4)  
**UCSC Stable ID:** uc002ae1  
**CCDS:** CCDS10159.1  
**UniProtKB/Swiss-Prot:** HTF4\_HUMAN [Target %id: 100, Query %id: 100] [align]  
**RefSeq peptide:** NM\_002205.1 [Target %id: 100, Query %id: 100] [align]  
**RefSeq DNA:** NM\_002205 [align]  
**UniProtKB/TriEMBL:** Q86TC1\_HUMAN [Target %id: 99, Query %id: 99] [align]  
Hypothetical protein DKPz451F163.  
Q9NYQ1\_HUMAN [Target %id: 5, Query %id: 100] [align]  
Transcription factor 12 (Fragment).  
Q9NYQ2\_HUMAN [Target %id: 3, Query %id: 100] [align]  
Transcription factor 12 (Fragment).  
Q9NYQ4\_HUMAN [Target %id: 3, Query %id: 100] [align]  
Transcription factor 12 (Fragment).  
Q9NYQ5\_HUMAN [Target %id: 8, Query %id: 100] [align]  
Transcription factor 12 (Fragment).  
Q9NYQ6\_HUMAN [Target %id: 7, Query %id: 100] [align]  
Transcription factor 12 (Fragment).

**Ensembl Human TransView**

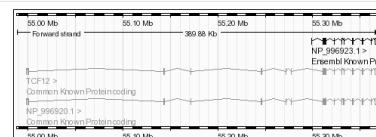
Ensembl release 48 - Homo sapiens Transcript Report for ENST00000267811

**GO** The following GO terms have been mapped to this entry via UniProt and/or RefSeq:  
GO:0003677 [DNA binding] **IEA**  
GO:0003702 [RNA polymerase II transcription factor activity] **TAS**  
GO:0005634 [nucleus] **IEA**  
GO:0006357 [regulation of transcription from RNA polymerase II promoter] **TAS**  
GO:0006955 [immune response] **TAS**  
GO:0007275 [multicellular organismal development] **IEA**  
GO:0007517 [muscle development] **TAS**  
GO:0030528 [transcription regulator activity] **IEA**  
GO:0045449 [regulation of transcription] **IEA**

**InterPro** IPR009057 Homeodomain-like - [View other genes with this domain]  
IPR001092 Basic helix-loop-helix dimerization region bHLH - [View other genes with this domain]

**Protein Family** ENSF0000001032 : TRANSCRIPTION FACTOR TRANSCRIPTION FACTOR  
This cluster contains 3 Ensembl gene member(s) in this species.

**Transcript structure** 

**Transcript neighbourhood** 

**Transcript sequence** The sequence is a long string of nucleotides (A, T, C, G) representing the transcript sequence, spanning from approximately 55.00 Mb to 55.30 Mb.

**Ensembl GeneView**

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**Ensembl Human GeneView**

Ensembl release 48: Homo sapiens Gene report for ENSG00000140262

Back Forward Reload Stop Home http://www.ensembl.org/Homo\_sapiens/geneview?gene=ENSG00000140262

**Gene** TCF12 (HGNC Symbol) **Synonyms:** HEB, HsT17266, HTF4  
This gene is a member of the Human CCDS set: CCDS10159, CCDS10160

**Ensembl Gene ID** ENSG00000140262

**Genomic Location** This gene can be found on Chromosome 15 at location 54,998,125-55,368,004.  
The start of this gene is located in Contig AC010999.6.1.221986.

**Description** Transcription factor 12 (Transcription factor HTF-4) (E-box-binding protein) (DNA-binding protein HTF4). Source: UniProt/SwissProt Q99081

**Prediction Method** Gene containing both Ensembl predicted transcripts and Havana manual annotation

**Transcripts**

TCF12	ENST00000267811	ENSP00000267811	[Transcript info]	[Exon info]	[Peptide info]
NP_996920_1	ENST00000333725	ENSP00000331057	[Transcript info]	[Exon info]	[Peptide info]
NP_996923_1	ENST00000343827	ENSP00000342459	[Transcript info]	[Exon info]	[Peptide info]

**Features ▾**

**Alignments**

This gene can be viewed in genomic alignment with other species

- view genomic alignment with 7 eutherian mammals Pecan
- view genomic alignment with 10 amniota vertebrates Pecan
- view genomic alignment with Rattus norvegicus
- view genomic alignment with Macaca mulatta
- view genomic alignment with Loxodonta africana
- view genomic alignment with Oryctolagus cuniculus
- view genomic alignment with Dasypus nemestrinus
- view genomic alignment with Canis familiaris
- view genomic alignment with Pan troglodytes
- view genomic alignment with Gallus gallus
- view genomic alignment with Ornithodoros chihuahuensis
- view genomic alignment with Bos taurus

**Chromosome 15 54,998,125 - 55,368,004**

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**click**

**Ensembl ExonView**

Ensembl release 48: Homo sapiens Exon Report for ENST00000267811

Back Forward Reload Stop Home http://www.ensembl.org/Homo\_sapiens/exonview?db=core;transcript=ENST00000267811

**Exons**

18	ENSE00001126365	15	1	55,352,520	55,352,752	2	1
Intron 18-19							
19	ENSE00001103528	15	1	55,352,753	55,361,934	1	-
Intron 19-20							
20	ENSE00001405489	15	1	55,362,089	55,365,646	-	-

**3' downstream sequence**

**Supporting Evidence**

The supporting evidence below consists of the sequence matches on which the exon predictions were based and are sorted by alignment score.

There are a large number of supporting evidence hits for this transcript. Only the top ten hits have been shown. Click to view all 23 supporting evidence hits.

**Score:** 100 → -99 → -97 → -90 → -85 → -80 → -75 → -70 → -60 → -50 → -40 → -30 → -20 → NO EVIDENCE

1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20
ML_020706.1	gi 3030791 refM_020305.1  Homo sapiens transcription factor 12 (HTF-4), helo-helix transcription factor 12 (HTF-4), transcript variant 1 mRNA	gi 3030791 refM_020305.3  Homo sapiens transcription factor 12 (HTF-4), helo-helix transcription factor 4 (HTF-4), transcript variant 3 mRNA	BC051790.2	BC051790.2  Homo sapiens cDNA clone IMAGE:5782025, containing frame-shifts	BC050596.1	BC050596.1  Homo sapiens transcription factor 12 (HTF-4), helo-helix transcription factor 4, mRNA (cDNA clone MGC:576579), complete cds	ENST02907.1												

**Ensembl GeneView**

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### Ensembl Human GeneView

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**Gene** TCF12 (HGNC Symbol) **Synonyms:** HEB, HsT17266, HTF4  
This gene is a member of the Human CCDS set: CCDS10159, CCDS10160

**Ensembl Gene ID** ENSG00000140262

**Genomic Location** This gene can be found on Chromosome 15 at location 54,998,125-55,368,004. The start of this gene is located in Contig AC010999.6.1.221986.

**Description** Transcription factor 12 (Transcription factor HTF-4) (E-box-binding protein) (DNA-binding protein HTF4). Source: UniProt/SwissProt Q99081

**Prediction Method** Gene containing both Ensembl predicted transcripts and Havana manual annotation

**Transcripts**

TCF12	ENST00000267811	ENSP00000267811	[Transcript info]	[Exon info]	[Peptide info]
NP_396920_1	ENST00000333725	ENSP00000331057	[Transcript info]	[Exon info]	[Peptide info]
NP_396923_1	ENST00000343827	ENSP00000342459	[Transcript info]	[Exon info]	[Peptide info]

**Features ▾**

**Alignments**

This gene can be viewed in genomic alignment with other species

- view genomic alignment with *7* eutherian mammals Pecan
- view genomic alignment with *10* amniota vertebrates Pecan
- view genomic alignment with *Rattus norvegicus*
- view genomic alignment with *Macaca mulatta*
- view genomic alignment with *Loxodonta africana*
- view genomic alignment with *Onychophora* tefrapi
- view genomic alignment with *Drosophila melanogaster*
- view genomic alignment with *Danio rerio*
- view genomic alignment with *Canis familiaris*
- view genomic alignment with *Pan troglodytes*
- view genomic alignment with *Gallus gallus*
- view genomic alignment with *Oryzias latipes*
- view genomic alignment with *Bos taurus*

**Ensembl ProtView**

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### Ensembl Human ProtView

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**Peptide** TCF12 (HGNC Symbol) To view all Ensembl genes linked to the name click here.  
This peptide is a member of the Human CCDS set: CCDS10159

**Ensembl Peptide ID** ENSP00000267811

**Translation information** This protein is a translation of transcript ENST00000267811, which is a product of gene ENSG00000140262.

**Genomic Location** This peptide can be found on Chromosome 15 at location 54,999,404-55,362,077. The start of this peptide is located in Contig AC010999.6.1.221986.

**Description** Transcription factor 12 (Transcription factor HTF-4) (E-box-binding protein) (DNA-binding protein HTF4). Source: UniProt/SwissProt Q99081

**Prediction Method** Gene containing both Ensembl predicted transcripts and Havana manual annotation

**InterPro** IPR009057 Basic Helix-loop-helix dimerisation region BHHL - [View other genes with this domain]  
IPR001092 Basic helix-loop-helix dimerization region BHHL - [View other genes with this domain]

**Protein Family** ENSF0000001037 : TRANSCRIPTION FACTOR TRANSCRIPTION FACTOR  
This cluster contains 3 Ensembl gene member(s) in this species.

**Protein Features**

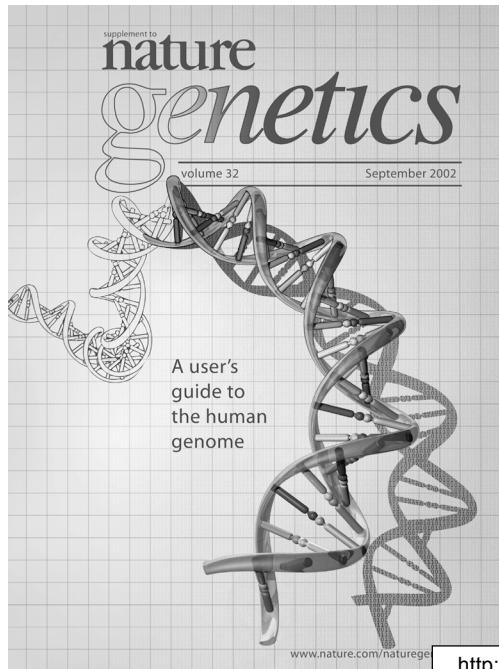
**Protein Sequence**

```
MHQGDPPWGIQIATKEL251LL373KQE2399VNSQ597PTDLSQ550592601K366P729
0750672572350T735027250GQ5314064L212791P9H42W53675LY
AIVMQRV9101250729723023323232323XYYDFTWYTTT173404N9320263
RDPYV2676530933062010151224435669954S3P3F7351472371109316774
S35527PWB86721158228905290260L094XQK91213975313169
S76769211L9702903932564583P7105051247L53123121H5742A8969
M77712225550593084467280616310583573219243458613096801090356
S71257479553108273561635805730761274342623155309573
S73135775053108273561635805730761274342623155309573
S71257479553108273561635805730761274342623155309573
S71257479553108273561635805730761274342623155309573
S71257479553108273561635805730761274342623155309573
GPTG1R2R12R1XNLLL12592115215940231213133397519334011
GPTG1R2R12R1XNLLL12592115215940231213133397519334011
GPTG1R2R12R1XNLLL12592115215940231213133397519334011
GPTG1R2R12R1XNLLL12592115215940231213133397519334011

Show the following features: 
Number residues: 
```

## Additional resources

- UCSC Human Genome Browser User Guide  
<http://genome.ucsc.edu/goldenPath/help/hgTracksHelp.html>
- NCBI Genomic Biology  
<http://www.ncbi.nih.gov/Genomes/>
- NCBI MapViewer Help  
<http://www.ncbi.nlm.nih.gov/mapview/static/MapViewerHelp.html>
- Ensembl Help & Tutorials  
<http://www.ensembl.org/info/using/website>



Current Protocols in Bioinformatics	
<b>The UCSC Genome Browser</b> <small>UNIT 1.4</small>	<b>Using the NCBI Map Viewer to Browse Genomic Sequence Data</b> <small>UNIT 1.5</small>
<p>The rapid progress of public sequencing and mapping efforts on vertebrate genomes has increased the demand for tools that offer quick and effective ways to analyze these levels and facilitate comparative data analysis. The University of California Santa Cruz (UCSC) Genome Bioinformatics Web site at <a href="http://genome.ucsc.edu">http://genome.ucsc.edu</a> contains a variety of genome analysis tools, most notably the UCSC Genome Browser (UNIT 1.4), a graphical tool for viewing genomic sequence and annotation tracks, and a collection of aligned annotation "tracks." Another tool, the MySQL MyS表 Browser—supplies convenient access to the MySQL database underlying the Genome Browser annotations. The custom annotation tracks feature that enables users to upload their own data and compare it against the genome sequence.</p> <p>This main protocol of this unit (see Basic Protocol) describes how to access and use the Genome Browser to browse genomic sequence data. It also includes a section on how to use the MySQL MyS表 Browser to access and use the MySQL database underlying the Genome Browser annotations.</p>	<p>The NCBI Map Viewer is an interface to a large, integrated set of genomic data, including sequence, cytogenetic, genetic linkage, and radiation hybrid maps, as well as the assembled and annotated genomic sequence itself. Along with the UCSC Genome Browser (UNIT 1.4) and Ensembl (UNIT 1.5), it is one of the primary Web sites from which genomic sequence data can be accessed.</p> <p>This unit includes an introduction to the Map Viewer (see Basic Protocol), which describes how to perform a simple text-based search of genome annotations to view the results, navigate along a chromosome, zoom in and out, and change the display to show information. It also describes some of NCBI's Map Viewers, which are provided as links from the Map Viewer. The Alternate Protocol 1 shows how to perform a BLAST search against the human genome. Alternate Protocol 2 shows how to retrieve a list of all genes between two STS markers. Alternate Protocol 3 shows how to find all annotated members of a genome.</p>
<b>Using the Ensembl Genome Server to Browse Genomic Sequence Data</b> <small>UNIT 1.15</small>	<p>Access through <a href="http://nihlibrary.nih.gov/ResearchTools/OnlineJournals.htm">http://nihlibrary.nih.gov/ResearchTools/OnlineJournals.htm</a></p>

Current Topics in Genome Analysis
<p>Next Lecture:</p> <p><b>Evolutionary Analysis</b></p> <p><i>Fiona Brinkman, Ph.D. Simon Fraser University</i></p>