

Mining Genomic Sequence Data

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Current Topics in Genome Analysis
January 25, 2005

Accessing the 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/>

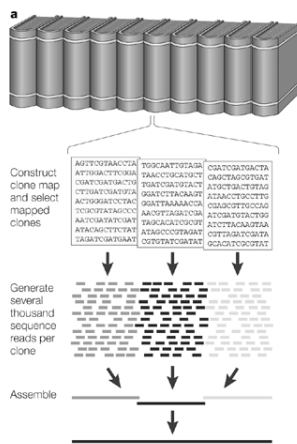
Ensembl
<http://www.ensembl.org>

Types of data integrated in genome browsers

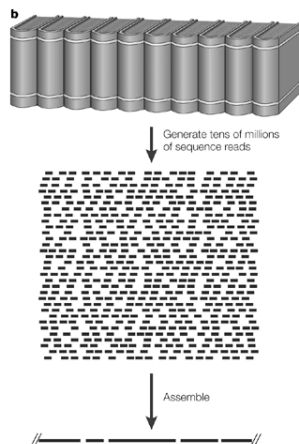
- Same starting material for all genome browsers: genomic sequence
- Annotations calculated independently by each genome browser
 - Genes
 - RefSeq mRNAs (non-redundant)
 - GenBank mRNAs (redundant)
 - ESTs
 - Gene predictions
 - SNPs
 - Homologous sequences from other organisms
 - STSs

Overview of genome sequencing strategies

Clone-by-clone shotgun sequencing



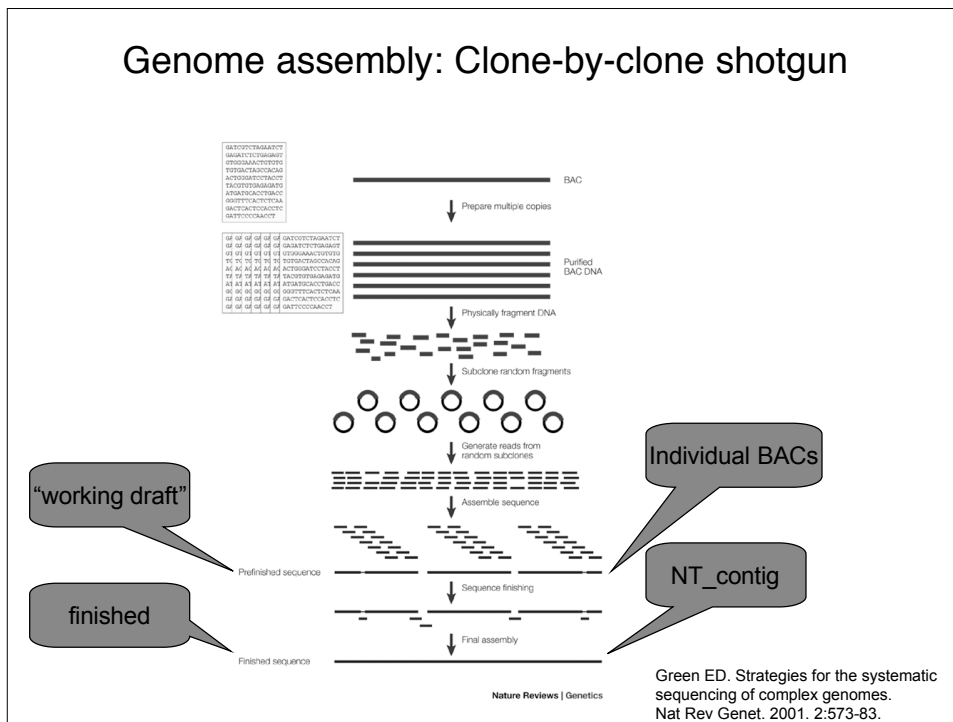
Whole-genome shotgun sequencing



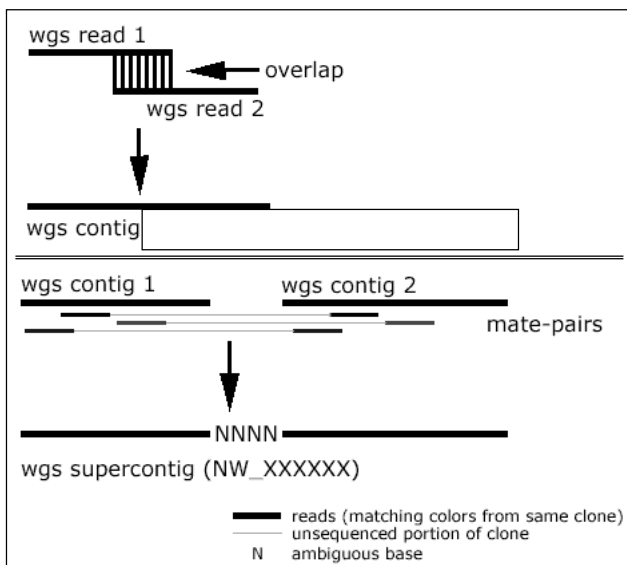
Nature Reviews | Genetics

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

Genome assembly: Clone-by-clone shotgun



Genome assembly: Whole genome shotgun (WGS)



<http://www.ncbi.nlm.nih.gov/genome/seq/NCBIContigInfo.html>

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
- UCSC is usually the first to display new assemblies, followed by NCBI and then Ensembl
 - “Pre-release” assemblies and annotations available at
 - UCSC: <http://genome-test.cse.ucsc.edu/>
 - pre!Ensembl: <http://pre.ensembl.org/>
 - UCSC provides access to older genome assemblies and annotations; NCBI and Ensembl do not
- 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	May 2004/hg17/Build 35	Build 35.1	Build 35
Mouse	Yes	May 2004/mm5/Build 33	Build 33.1	Build 33
Rat	Yes	June 2003/rn3/RGSC 3.1	Build 2.1	RGSC 3.1 (RGSC 3.2 on pre!)
Chicken	Yes(?)	February 2004/galGal2	Build 1.1	WASHUC1
Chimp	Yes, but NCBI is using a different chromosome numbering system	November 2003/ panTro1/NCBI Build 1.1	Build 1.1	CHIMP1
Fugu	Yes	August 2002/ fr1/v3.0	-	Fugu v2.0

UCSC

View a region in the genome by querying with a gene symbol

The screenshot shows the UCSC Human Genome Browser Gateway interface. At the top, there is a navigation menu with links: Home, Genomes, Gene Sorter, Blat, PCR, Tables, FAQ, and Help. Below the menu is the title "Human Genome Browser Gateway" and a copyright notice: "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." The main search area contains four input fields: "genome" (set to "Human"), "assembly" (set to "May 2004"), "position" (set to "adam2"), and "image width" (set to "620"). A "Submit" button is located to the right of the "image width" field. Below the search fields, there is a link: "Click here to reset the browser user interface settings to their defaults." and a button: "Add Your Own Custom Tracks".

About the Human May 2004 (hg17) assembly (sequences)

The May 2004 human reference sequence is based on NCBI Build 35 and was produced by the International Human Genome Sequencing Consortium.

Sample position queries

A genome position can be specified by the accession number of a sequenced genomic clone, an mRNA or EST or STS marker, or a cytological band, a chromosomal coordinate range, or keywords from the GenBank description of an mRNA. The following list shows examples of valid position queries for the human genome. See the [User's Guide](#) for more information.

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
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. Includes 100,000 bases on each side as well.
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 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

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Human adam2 - UCSC Genome Browser v91

http://genome.ucsc.edu/cgi-bin/hgTracks?org=Human&db=hg17&position=adam2&pix=620&hgsid=37847

Known Genes

ADAM2 at chr8:39720413-39814885 - (BC034957) a disintegrin and metalloproteinase domain 2 (fertilin beta)
 ADAM2 at chr8:39720413-39814886 - (BC064547) ADAM2 protein.
 ADAM20 at chr14:70058831-70061255 - (AF029899) a disintegrin and metalloproteinase domain 20
 ADAM21 at chr14:69993969-69996375 - (AF029900) a disintegrin and metalloproteinase domain 21
 ADAM22 at chr7:87208352-87471098 - (AF073291) a disintegrin and metalloproteinase domain 22
 ADAM23 at chr2:207134044-207308385 - (AJ005580) a disintegrin and metalloproteinase domain 23
 ADAM27 at chr8:24207560-24268547 - (AJ133004) a disintegrin and metalloproteinase domain 18
 ADAM28 at chr8:24207560-24268547 - (AJ242015) a disintegrin and metalloproteinase domain 28
 ADAM29 at chr4:176273863-176273986 - (AF134708) a disintegrin and metalloproteinase domain 29

RefSeq Genes

ADAM2 at chr8:39720413-39814936 - (NM_001464) a disintegrin and metalloproteinase domain 2
 ADAM20 at chr14:70058831-70071485 - (NM_003814) a disintegrin and metalloproteinase domain 20
 ADAM21 at chr14:69993969-69996375 - (NM_003813) a disintegrin and metalloproteinase domain 21
 ADAM22 at chr7:87208352-87471098 - (NM_018351) a disintegrin and metalloproteinase domain 22
 ADAM22 at chr7:87208352-87456079 - (NM_021721) a disintegrin and metalloproteinase domain 22
 ADAM22 at chr7:87208352-87471098 - (NM_021722) a disintegrin and metalloproteinase domain 22
 ADAM22 at chr7:87208352-87471098 - (NM_021723) a disintegrin and metalloproteinase domain 22
 ADAM22 at chr7:87208352-87456079 - (NM_004194) a disintegrin and metalloproteinase domain 22
 ADAM23 at chr2:207133873-207308383 - (NM_003812) a disintegrin and metalloproteinase domain 23
 ADAM28 at chr8:24207560-24268550 - (NM_014265) a disintegrin and metalloproteinase domain 28
 ADAM28 at chr8:24207568-24249555 - (NM_021777) a disintegrin and metalloproteinase domain 28
 ADAM29 at chr4:176271406-176273869 - (NM_021779) a disintegrin and metalloproteinase domain 29
 ADAM29 at chr4:176271406-176273692 - (NM_021780) a disintegrin and metalloproteinase domain 29
 ADAM29 at chr4:176226863-176273986 - (NM_014269) a disintegrin and metalloproteinase domain 29

Human Aligned mRNA Search Results

AJ005580 - Homo sapiens mRNA for adam23 protein.
 BC034957 - Homo sapiens a disintegrin and metalloproteinase domain 2 (fertilin beta), mRNA (cDNA clone MGC:26432 IMAGE:4826530), complete cds.
 BC064547 - Homo sapiens a disintegrin and metalloproteinase domain 2 (fertilin beta), mRNA (cDNA clone MGC:74935 IMAGE:5744846), complete cds.
 AF158637 - Homo sapiens metalloproteinase-disintegrin ADAM22-3 (ADAM22) mRNA, alternatively spliced, partial cds.
 BC025378 - Homo sapiens a disintegrin and metalloproteinase domain 20, mRNA (cDNA clone MGC:25993 IMAGE:4827383), complete cds.
 AF155381 - Homo sapiens metalloproteinase-like, disintegrin-like, cysteine-rich protein 2 delta (ADAM22) mRNA, alternative splice product, complete cds.
 AF155382 - Homo sapiens metalloproteinase-like, disintegrin-like, cysteine-rich protein 2 epsilon (ADAM22) mRNA, complete cds.
 AF134708 - Homo sapiens disintegrin and metalloproteinase domain 29 (ADAM29) mRNA, complete cds.
 AF171929 - Homo sapiens metalloproteinase-disintegrin (ADAM29) mRNA, complete cds.
 AF171930 - Homo sapiens metalloproteinase-disintegrin beta (ADAM29) mRNA, alternatively spliced, complete cds.
 AF171931 - Homo sapiens metalloproteinase-disintegrin gamma (ADAM29) mRNA, alternatively spliced, complete cds.
 AK129906 - Homo sapiens cDNA FLJ26396 fis, clone HRT08301, highly similar to Homo sapiens a disintegrin and metalloproteinase domain 23

Non-Human Aligned mRNA Search Results

Human chr8:39,720,413-39,814,936 - UCSC Genome Browser v91

http://genome.ucsc.edu/cgi-bin/hgTracks?position=chr8:39720413-39814936&hgsid=37847641&refGene=

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

UCSC Genome Browser on Human May 2004 Assembly

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

position chr8:39,720,413-39,814,936 size 94,524 bp. image width: 620 jump

move start Click on a feature for details. Click on base position to zoom in move end
 < 2.0 > around cursor. Click on left mini-buttons for track-specific options. < 2.0 >

reset all hide all Chromosome Guidelines Labels: left center refresh

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

Mapping and Sequencing Tracks

Base Position Chromosome Band STS Markers RGD/OTL FISH Clones

Go to 'http://genome.ucsc.edu/cgi-bin/hgTracks?hgsid=37847641&c=chr8&g=stsMap'

Human Gene ADAM2 Description and Page Index

Details: Known Genes Track

Home - Genomes - Genome Browser - Gene Sorter - Blat - PCR - Tables - FAQ - Help

Human Gene ADAM2 Description and Page Index

Description: a disintegrin and metalloproteinase domain 2 (fertilin beta)
Representative mRNA: BC034957 **Protein:** Q99965 (AD02_HUMAN)
RefSeq Summary: This gene encodes a disintegrin and metalloprotease (ADAM) domain 2, which is a member of the ADAM protein family. Members of this family are membrane-anchored proteins structurally related to snake venom disintegrins, and have been implicated in a variety of biologic 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.

Page Index	Quick Links	SwissProt Comments	Sequence	Microarray	RNA Structure
Protein Structure	Other Species	GO Annotations	mRNA Descriptions	Methods	

Quick Links to Tools and Databases

Genome Browser	Proteome Browser	Gene Sorter	SwissProt	LocusLink	Entrez Gene
PubMed	OMIM	GeneLynx	GeneCards	CGAP	Stanford SOURCE
Jackson Labs					

Comments and Description Text from SwissProt

ID: AD02_HUMAN
DESCRIPTION: ADAM 2 precursor (A disintegrin and metalloproteinase domain 2) (Fertilin beta subunit) (PH-30) (PH30).
FUNCTION: Sperm surface membrane protein that may be involved in sperm-egg plasma membrane adhesion and fusion during fertilization. Could have a direct role in sperm-zona binding or migration of sperm from the uterus into the oviduct. Interactions with egg membrane could be mediated via binding between its disintegrin-like domain to one or more integrins receptors on the egg. This is a non catalytic metalloprotease-like protein.
SUBCELLULAR LOCATION: Type I membrane protein.
TISSUE SPECIFICITY: Expressed specifically in spermatogenic cells in the seminiferous cells. Not detected in fetal tissues.
DOMAIN: A tripeptide motif (FEE) within disintegrin-like domain could be involved in the binding to egg integrin receptor and thus could mediate sperm/egg binding.
PTM: The prodomain and the metalloprotease domain are cleaved during the epididymal maturation of the spermatozoa.
MISCELLANEOUS: In mammals, exists as a heterodimer composed of an alpha and beta subunits. In human, fertilin alpha is a pseudogene.
SIMILARITY: Belongs to peptidase family M12B.
SIMILARITY: Contains 1 disintegrin domain.
SIMILARITY: Contains 1 EGF-like domain

Human protein AD02_HUMAN - UCSC Proteome Browser

Home **UCSC Proteome Browser** PDF/PS Help

Human protein: **Q99965** (aka AD02_HUMAN) ADAM 2 precursor (A disintegrin and metalloproteinase domain 2) (Fertilin beta subunit) (PH-30) (PH30).

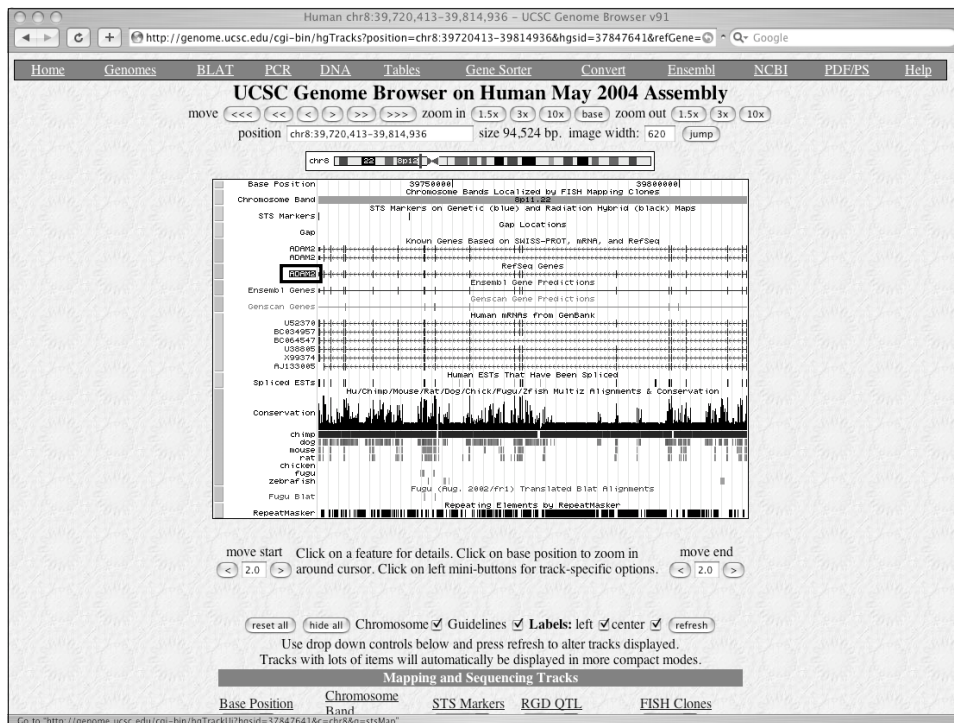
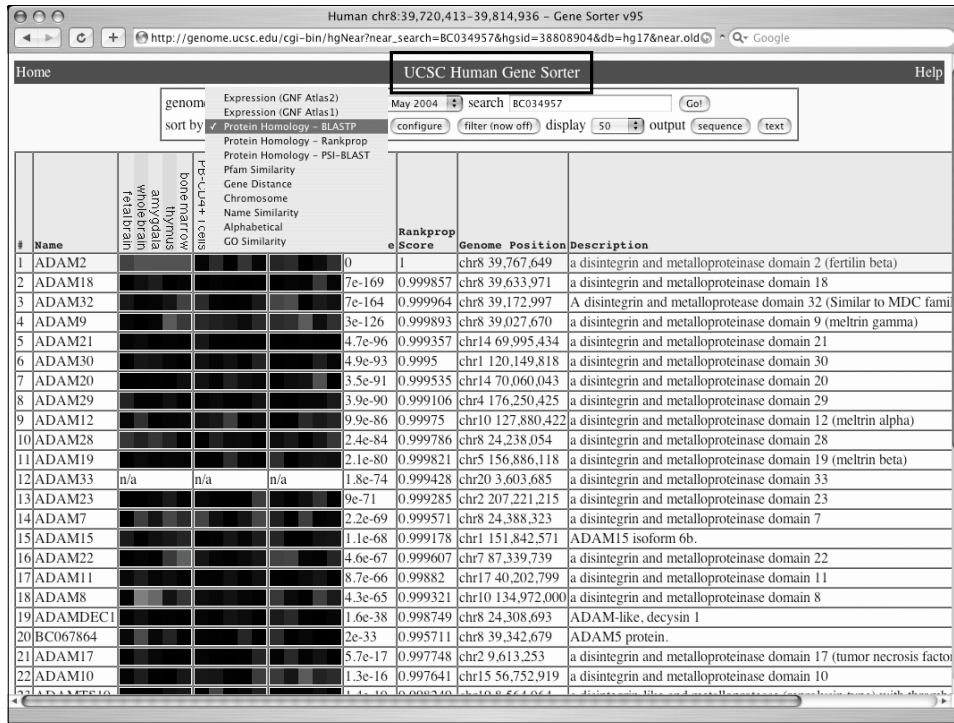
Move <<< << < > >> >>> Current scale: FULL Rescale to 1/6 1/2 FULL DNA

Explanation of Protein Tracks

pI 5.8	Molecular Weight 62456 Da	Number of Exons 19	Amino Acid Frequencies
InterPro Domains 6	Hydrophobicity -0.2	Number of Cysteines 43	Amino Acid Anomalies

Display a menu

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RefSeq Gene: **ADAM2** Details: RefSeq Genes Track

RefSeq: [NM_001464.3](#) Status: **Reviewed**
 CDS: 3' complete
 OMIM: [601533](#)
 LocusLink: [2515](#)
 Entrez Gene: [2515](#)
 PubMed on Gene: [ADAM2](#)
 PubMed on Product: [a disintegrin and metalloproteinase domain 2](#)
 GeneLynx: [ADAM2](#)
 GeneCards: [ADAM2](#)
 AceView: [ADAM2](#)
 Stanford SOURCE: [NM_001464](#)

Summary of ADAM2

This gene encodes a disintegrin and metalloprotease (ADAM) domain 2, which is a member of the ADAM protein family. Members of this family are membrane-anchored proteins structurally related to snake venom disintegrins, and have been implicated in a variety of biologic 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.

mRNA/Genomic Alignments

SIZE	IDENTITY	CHROMOSOME	STRAND	START	END	QUERY	START	END	TOTAL
2640	100.0%	8	-	39720414	39814936	NM_001464	1	2640	2657

Position: [chr8:39720414-39814936](#)
Band: [Sp11.22](#)
Genomic Size: 94523
Strand: -

Links to sequence:

- [Predicted Protein](#)
- [mRNA Sequence](#) may be different from the genomic sequence.
- [Genomic Sequence](#) from assembly

Genomic Sequence Near Gene

Home - Genomes - Genome Browser - Gene Sorter - Blat - PCR - Tables - FAQ - Help

Get Genomic Sequence Near Gene

Note: if you would prefer to get DNA for more than one feature of this track at a time, try the [Table Browser](#) using the output format sequence.

Sequence Retrieval Region Options:

Promoter/Upstream by bases
 5' UTR Exons
 CDS Exons
 3' UTR Exons
 Introns
 Downstream by bases
 One FASTA record per gene.
 One FASTA record per region (exon, intron, etc.) with extra bases upstream (5') and extra downstream (3')
 Split UTR and CDS parts of an exon into separate FASTA records

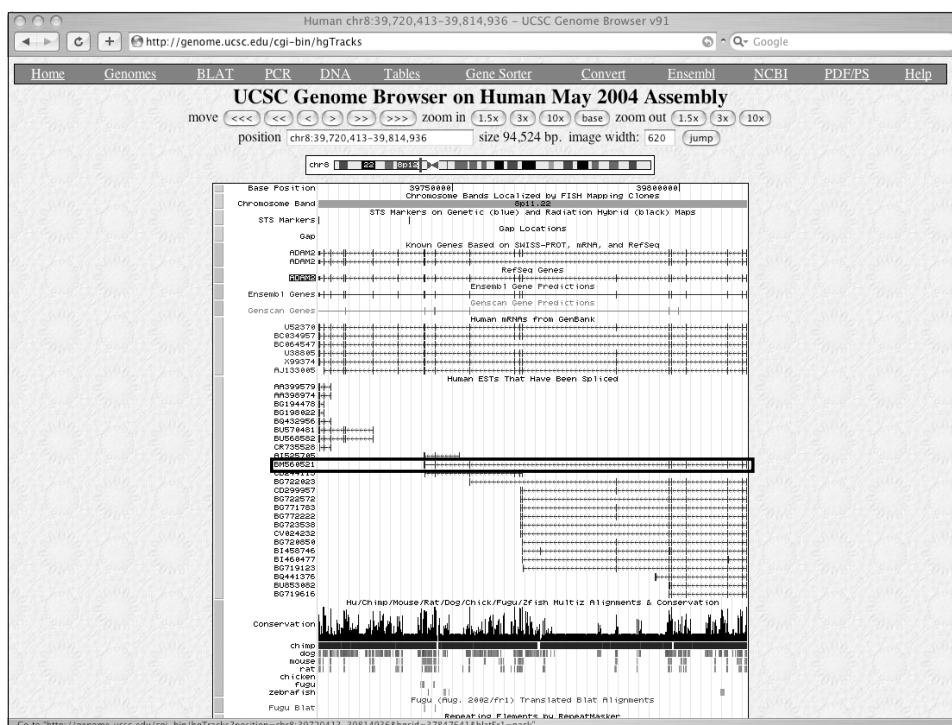
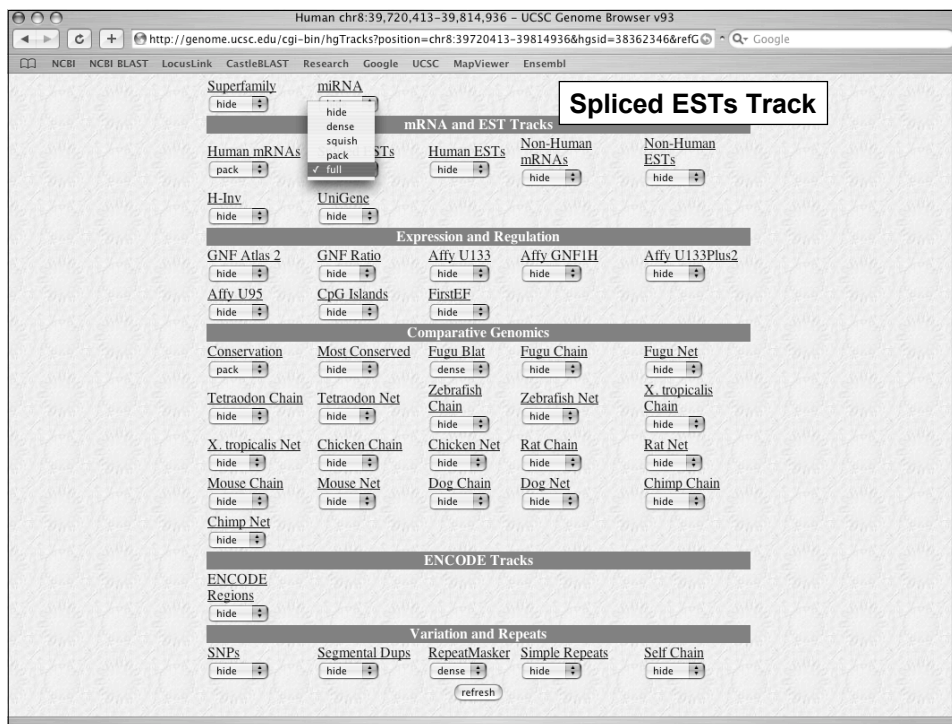
Note: if a feature is close to the beginning or end of a chromosome and upstream/downstream bases are added, they may be truncated in order to avoid extending past the edge of the chromosome.

Sequence Formatting Options:

Exons in upper case, everything else in lower case.
 CDS in upper case, UTR in lower case.
 All upper case.
 All lower case.
 Mask repeats: to lower case to N

Display a menu

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BM560521

http://genome.ucsc.edu/cgi-bin/hgChgsid=38386066&o=39743656&t=39814886&g=intronEst&i=BM560521

Home - Genomes - Genome Browser - Gene Sorter - Blat - PCR - Tables - FAQ - Help

Information on EST BM560521

Description: n/a
Gene: n/a
Product: n/a
Author: NIH-MGC <http://mgc.nci.nih.gov/>
Organism: [Homo sapiens](#)
Tissue: medulla
Development stage: n/a
Cell type: n/a
Sex: n/a
Library: NIH_MGC_119
Clone: IMAGE:5744846
Read direction: 5'
CDS: n/a
Date: 2002-01-20
Version: 1
Stanford SOURCE: [BM560521](#) [[Gene Info](#)] [[Clone Info](#)]
EST sequence: [BM560521](#)

EST/Genomic Alignments

SIZE	IDENTITY	CHROMOSOME	STRAND	START	END	QUERY	START	END	TOTAL
1004	98.43	8	-	39743657	39814886	BM560521	20	1051	1063

Description

The Spliced EST track displays Expressed Sequence Tags (ESTs) from [GenBank](#) that show signs of splicing when aligned against the genome. To be considered spliced, an EST must show evidence of at least one canonical intron, i.e. one that is at least 32 bases in length and has GT/AG ends. By requiring splicing, the level of contamination in the EST databases is drastically reduced at the expense of eliminating many genuine 3' ESTs. For a display of all ESTs (including unspliced), see the human EST track.

BM560521 vs Genomic

http://genome.ucsc.edu/cgi-bin/hgChgsid=38386066&g=htcCdnaAli&i=BM560521&c=chr8&l=397204

Alignment of BM560521 and chr8:39743657-39814886

Click on links in the frame to the left to navigate through the alignment. Matching bases in cDNA and genomic sequences are colored blue and capitalized. Light blue bases mark the boundaries of gaps in either sequence (often splice sites).

Alignment of BM560521

[BM560521](#)
[Human chr8](#)
[block1](#)
[block2](#)
[block3](#)
[block4](#)
[block5](#)
[block6](#)
[block7](#)
[block8](#)
[block9](#)
[block10](#)
[together](#)

cDNA BM560521

```

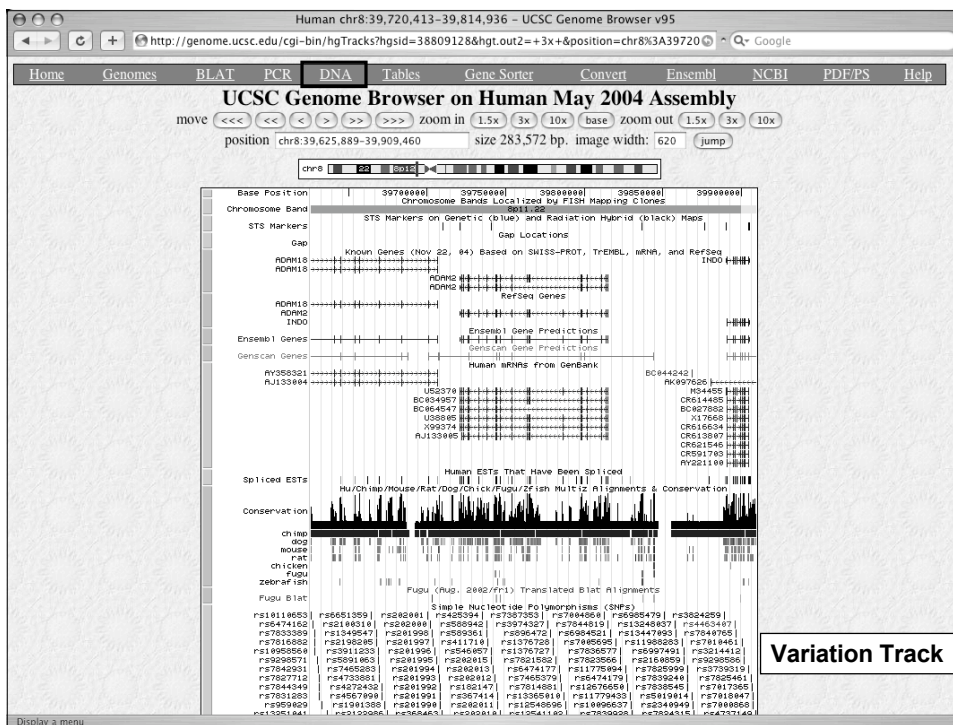
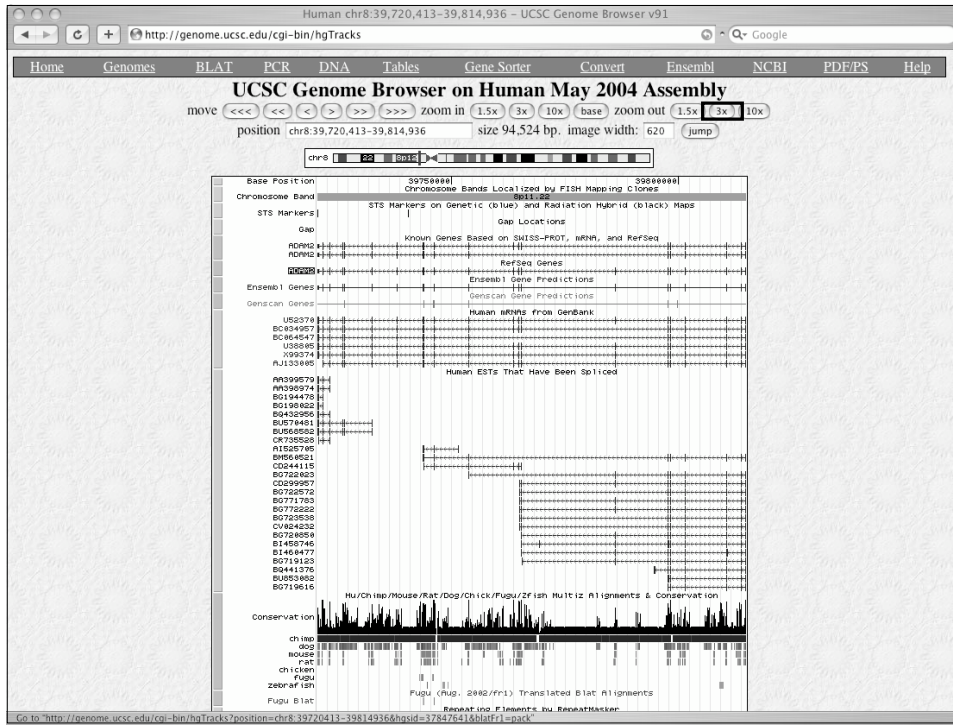
ggteccggaat tceccgggat GGCTGGGACC CAGGACTTCA AGCCATGTGG 50
CGCGTCTTGT TTCCTCTCAG CGGGCTCGGC GGGCTGCGGA TGGACATGAA 100
TTTTGATAGT TTACCTGTGC AAATTACAGT TCCGGAGAAA ATACGGTCAA 150
TAATAAAGGA AGGAATTGAA TCGCAATCAT CTACAAAAT TGTAAATTGAA 200
GGGAACCAT ATACTGTGAA TTTAATGCAA AATAACTTTT TACCCCATRA 250
TTTTAGAGTT TACAGTTATA GTGGCAGAG AATTATGAAA CCACCTGACC 300
AAGATTTTCA GAATTTCTGC CACTACCAAG GGATATTTGA AGGTATGCA 350
AAATCTGTGG TGATGTTTAC CACATGTACT GGAATCAAGG CGCTACTACA 400
GTTTGAATAA GTTAGTTATG GAATAGAACC COTGGAGTCT TCAATTGGCT 450
TTGACATGAT AATTACCAA GTAACAATA AGAAGCAGA TGTTCCTTA 500
TATATGAGA AGGATTTTCA ATCAGAGAT CTGTCCCTTA AATTACAAAG 550
CTAGAAAC CCAGAACCA TAAGTCTGGA ATCACTTGCA GTTATTTAG 600
CTCAATTAAT GAGCCTTAGT ATGGGGATCA CTTATGATGA CATTAAACAA 650
TGCCAGTGCT CAGGACTGAT CTGCATTATG AATCCAGAAG CAATTCATT 700
CAGTGGTGTG AAGATCTTTA GTAACCTGAG CTTGCAAGAC TTGCACATT 750
TTATTTCAA GCAGAAGTCC CAGTGTCTTC ACAATCAGCC TCCTTTAGAT 800
CTTTTTC AACAACAAGC AGTGTGTGT AATGCAAGC TGGAAACAG 850
AGAGAGTGT GACTGTGGGA CTGACAGAG a GTGCCCTT ATGGAGAAA 900
aCTGCTGTGA TATTCCTCC ATGTAATTT TAAAGCCGT TCAACTGTG 950
CctgaAGGA CATGCTGCGA AAactggcc aTTTATGTCA AAAGAAAGA 1000
ATggTAGG CctTCCTTT GAaAATGC GaaCTCCC TGAaaATTG 1050
Ceattggaat cct
                    
```

Genomic chr8 (reverse strand):

```

cccacctggg ctctcccage cgcctacctc tcccagggtg cgtggccggg 39814937
gegtcatctc gegcttccaa ctgcccctgta accaccaact gccattatcc 39814887
CGGCTGGGAC CCAGAGCTTC AAGCCATGTG CGCGCTCTTG TTTCTGTCTCA 39814837
CGGGCTCTCG CGGGCTCGGC ATGGACAGT: gtaagcagaa aaacctctcc 39814787
ctctggccttt tgggaccttc agcctacttc tccctttgce tccagttac 39814737
attgacatcc ctgggagat getttctgta ggggtttcca actcagaat 39814687
tccaaagctca tcccagacat tgcgccttc cccaccceaaa atccacgtag 39814637
aataaggctc tcccacacg tctggaaac ctaggaaaga taagatagga 39814587
ggctctagge cttaaagggt tgettcaaaa tgtagtttc agactctgct 39814537
ttgatttttt cagcagagt aaagttgaga cccaacttct gactctcta 39814487
                    
```

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Extended DNA Case/Color

Extended DNA Case/Color Options

Use this page to highlight features in genomic DNA text. DNA covered by a particular below for details about color, and for examples. Tracks in "hide" display mode are not:

Position Reverse complement

Letters per line Default case: Upper Lower

Track Name	Toggle Case	Underline	Bold	Italic	Red	Green	Blue
ChromosomeBand	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
STS Markers	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Known Genes	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
RefSeq Genes	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Ensembl Genes	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Genscan Genes	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Human mRNAs	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Spliced ESTs	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Conservation	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Fugu Blat	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
SNPs	<input type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
RepeatMasker	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

SNPs in RefSeq exon

```

...
tgttactaatttttgcgactcctgaagggaatcagcagtgtagactgaaataat
catccttactttaaagtctatatacttgaagtgtatctttttccagcagatggt
gaaaatttttccaaatctagpatcattgecttggatgattcagactccacactat
aacataacacataccctgcaatttGAAATGTACAGTGGCTTTAATCAATACCCACC
TCATTACATTATGATCTCTTATACCCATGCCCTACTGACAGAAATGATGCTACTAA
CCAACAGAAATAGCTTAAGATAACTTTAAAAAACAGAAATTTGTAATGAAAACATT
ATCATGCAATTTTATAAGTAAATTTATCTGGAATTTATACATATATATCATATCT
GGTCAAAATTAATCTCAATATGACAGTTTGAATCCAGGAACCCAGGTAAATGACTTC
TTTTCTTCATTTTGTGTCCATCCACAGATATAATTAATCTAGAGactgaaataa
atgaaaaaaaagtcaccgttatcaacaagtgaatgatagctagtccaacaatgca
tgcgaagtaacctattgcaagtagactataatgcaattatttttagccaaccttt
taaatatacaaatgctaaagaagtaaaatcaactaaagaasaatataaaaaatacaact
gtaataaatttttaagtggaaatttcaaatcaattataacttgatcatgtctattata
aatctgaaaaataaatccgaaaaatagctctcaatttattttatataatagcta
tatgaactgcaactactaataatataattttaaataatagctctctctctctctcc
ccccaa.cccccatccttaaaatatttgaactcactcattgctccttgaaggcaatgg
atgaagctggaagcactatctcaacaacaacacaggaataaaacccaacaccgc
atgtctcactcataagtggagctgaatcatgaaaacacatggacagaggggggaac
atcacaccagaccctgctgggggttggaggccaaggagggaagacatggagaccaa
tacctaatgatgtggggttaaaacctagatgacgggttgataggtgcagcaaacacc
atggcactgtataacctatgtaaaaacctgcaactctgcaactctgctcccagtaacta
aagtaaaataaatttataaaatgaagaaactcaactgataatcagctcctctctgtt
GTCCAGACTACCCCTTAGGTTCACTCTCACTTTCAGGTGCTGCTGATttaaataaaa
ataatattatactttctcaagatgatttcaaacctctgtatattctgctagcttga
aacagttcgaattttacctaagaactgggttcaacctattttgagataacctatca
ctgagaacatctctgcccgtgtaatacaagactaataatgaacatcttgcactcagtgat
atgaaaaaaaagcagtgatataatgtttaccgaaaaagaaccttagatcatgtttat
ctatgatatattggttaagtacttttaacttttgcactcattttcaacctattaa
ttctctcattttactagctatgcttttccctgaagatattgatatcagttatccc
acaatgattccaataagaaacaccttagtatattaaataagtaagttatt
actggttagtgctgatatatttcagcaatttcaactcaacttttaaatatattcc
atttcattacattgctcttttaagtattttaaagaactcaactcatttcaagtaaaa
tgaaggacatggttatatactcccaggagaactcatttggtagcatttctctgaaa
taataaaacagtttaagtcttactgtatcttagcacaattatcagatattttat
ttttaaagaagttattgtttatatttctgtagactagctgggctatcctagat
    
```

UCSC

Find a chicken homolog of a human protein

NHGRI Current Topics in Genome Analysis 2005
Mining Genomic Sequence Data

NCBI Entrez Protein search results for **np_001455**. The search results show the protein **Disintegrin and metalloproteinase domain 2 proprotein [Homo sapiens]** with accession number **NP_001455.3**. The amino acid sequence is displayed as follows:

```
>gi|55743080|ref|NP_001455.3| a disintegrin and metalloproteinase domain 2 proprotein
MNRVLLFLLSGLGRMDSNFDLPVQITVPEKIRSIIEKIGESQASYKIVIEGPKYTVNLMQKNFLPHNF
RVVSYSGTGIMKPLDQDFQNFCHYQYIEGYPKSVVMVSTCTGLRGLVQFENVSYGIEPLESSVGFHEVI
YQVHRKADVSLYNEKDIERSDLSFKLQSVPEQDFAKYIEMHIVIEKQLYNHMGSDTTVAQKVFQIIG
LTNAIFVSNITLISLELWIDENKIATTEANELLTFLRWKTSYLVLPHDVAFLVYREKSNVYGA
TFQGMCDANYAGGVVLPRTISLESLELAVLQQLLSMGIYDDINKCQCSGAVCMNPEAIHFSGVKI
FNSCSFEDFAHFISKRQSQCLHNPRLDPFFKQAVCGNAKLEAGEECDCGTEQDCALIGETCCDIATCR
FKAGSNCAEGPCENCLFMSKERMCRPSFECDLPEYCNSSASCENHYVOTGHPGCLNQMVICIDGVCN
SGDRQCTDTPGKEVFPFSGVSBYSBLNSITDVSQNGSIDSGYTCEDANLQCGKLICTVGRFLLQIPRA
TTIYANISGHLCIAVEFASDHADSQMKWIKDGTSCSNKRCRNRQCVSSYLGVYDCTDKCNDRGVCNKK
KHCHCSASYLPPDCSQSDLPGGSDSNFPVAIPARLPERRYENIYHSKPMRWPFLLPFIPIIFC
VLIAMVKNVFRQKKWRTEYSSDEQPESESEPKG
```

Chicken BLAT Search

BLAT Search Genome

Genome: **Chicken** Assembly: **Feb. 2004** Query type: **BLAT's guess** Sort output: **query.score** Output type: **hyperlink** [Submit] [Reset]

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

```
>gi|55743080|ref|NP_001455.3| a disintegrin and metalloproteinase domain 2 proprotein [Homo sapiens]
MNRVLLFLLSGLGRMDSNFDLPVQITVPEKIRSIIEKIGESQASYKIVIEGPKYTVNLMQKNFLPHNF
RVVSYSGTGIMKPLDQDFQNFCHYQYIEGYPKSVVMVSTCTGLRGLVQFENVSYGIEPLESSVGFHEVI
YQVHRKADVSLYNEKDIERSDLSFKLQSVPEQDFAKYIEMHIVIEKQLYNHMGSDTTVAQKVFQIIG
LTNAIFVSNITLISLELWIDENKIATTEANELLTFLRWKTSYLVLPHDVAFLVYREKSNVYGA
TFQGMCDANYAGGVVLPRTISLESLELAVLQQLLSMGIYDDINKCQCSGAVCMNPEAIHFSGVKI
FNSCSFEDFAHFISKRQSQCLHNPRLDPFFKQAVCGNAKLEAGEECDCGTEQDCALIGETCCDIATCR
FKAGSNCAEGPCENCLFMSKERMCRPSFECDLPEYCNSSASCENHYVOTGHPGCLNQMVICIDGVCN
SGDRQCTDTPGKEVFPFSGVSBYSBLNSITDVSQNGSIDSGYTCEDANLQCGKLICTVGRFLLQIPRA
TTIYANISGHLCIAVEFASDHADSQMKWIKDGTSCSNKRCRNRQCVSSYLGVYDCTDKCNDRGVCNKK
KHCHCSASYLPPDCSQSDLPGGSDSNFPVAIPARLPERRYENIYHSKPMRWPFLLPFIPIIFC
VLIAMVKNVFRQKKWRTEYSSDEQPESESEPKG
```

Rather than pasting a sequence, you can choose to upload a text file containing the sequence.
Upload sequence: **Choose File** no file selected **Submit File**

Only DNA sequences of 25,000 or fewer bases and protein or translated sequence of 10,000 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.

About BLAT

BLAT on DNA is designed to quickly find sequences of 95% and greater similarity of length 40 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 21 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.

NHGRI Current Topics in Genome Analysis 2005
Mining Genomic Sequence Data

Chicken BLAT Results

http://genome.ucsc.edu/cgi-bin/hgBlat

Home - Genomes - Gene Sorter - Blat - Tables - 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

Display a menu

Chicken chrUn:635,370-635,555 - UCSC Genome Browser v9.5

http://genome.ucsc.edu/cgi-bin/hgTracks?position=chrUn:635370-635555&db=galGal2&ss=.../trash/hg

Home Genomes BLAT DNA Tables Convert PDE/PS Help

UCSC Genome Browser on Chicken Feb. 2004 Assembly

position chrUn:635,370-635,555 size 186 bp. image width: 620

Base Position: 635440 635450 635550

Gap Locations

NP_001455.3 Your Sequence from BLAT Search

RefSeq Genes

Non-Chicken RefSeq

Ensembl Genes

Ensembl Gene Predictions

Spliced ESTs

Fugu Blat

Hg16 Het

BGI SNPs

Seif Het

RepeatMasker

move start Click on a feature for details. Click on base position to zoom in around cursor. Click on left mini-buttons for track-specific options. move end

reset all hide all Guidelines Labels: left center refresh

Chromosome Color Key: 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 X Y M Un

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

Mapping and Sequencing Tracks

Base Position full hide

Supercontigs hide

Assembly hide

Gap dense

BAC End Pairs hide

GC Percent hide

Quality Scores hide

Contamination hide

Short Match hide

BLAT Sequence pack

Genes and Gene Prediction Tracks

Display a menu

Chicken BLAT Results

http://genome.ucsc.edu/cgi-bin/hgBlat

Home - Genomes - Gene Sorter - Blat - Tables - 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

Display a menu

User Sequence vs Genomic

http://genome.ucsc.edu/cgi-bin/hgc?o=635369&g=htcUserAll&i=../trash/hg5s_genome_27788_110608

Alignment of NP_001455.3

NP_001455.3
Chicken.chrUn
block1
block2
block3
together

Alignment of NP_001455.3 and chrUn:635370-635555

Click on links in the frame to the left to navigate through the alignment. Matching bases are colored blue and capitalized. Light blue bases mark the boundaries of gaps in either sequence.

NP_001455.3

```

mwrvfllsg lgglrmdsnf dslpvqitvp ekirsiikeg iesqasykiv iegkpytvnl 60
mqknflphnf rvsyasygtgi mkpldqdfqn fchygygieg ypksvvmvat ctglrgvlqf 120
envsygiopl esvsygehvi yqvkhhkadv slynekdiess rdlsfklqav epqgdakyl 180
emhvivkeql ynhmsgdtv vaqkvfqlg ltnaifsvfn ltiilssalel widsnkiaat 240
geanelhtf lrwktaylv rphdvafllv yreksnyvga tfqgkmedan yaggvvihpr 300
tisleslavi laqlslismg ityddinkc csqavcimp eaihfsqyki fsncsfedfa 360
hfiskqkagc lhnqprldpf fkqgavcna kleageecdc gteqdcalg etccdiater 420
fkagsnaeag pccencfms kermcrpsfe ecdlpeycng ssasopenhy vqtghpcgln 480
qwicidgvcm sgdkqctdtf gkevefpgse cyshinsktid vsngcngsids gytgceadL 540
qCCKLICKV gkfligipra IIPAnisgh LLaivefaad hadsqkwiL DGTGQanRV 600
cncgrevass ylydcttdk cndrgvennk khchcsasyl ppdcsvqsd wpggsidsgn 660
fppvaiparl perryieniy hskpmrvpff lfipffiiic vliainvkvn fgrkkwrtd 720
yssdeqese sepkg
    
```

Chicken.chrUn :

```

AATCTGggcT GTGGAAACT CATCTGcaca TA^ccaaaac gagttccctt caccaaatta 635429
aagggt^CCA TCATCTATGC Teaagtcaaa gaaCATCTGT G:gtgtcttt tgatgtaatg 635489
catgcacct ccgggacaga tcctctectg gtt^AGGATG GCACGaaaTG CGGTcccgga 635549
AAGGT^
    
```

Side by Side Alignment*

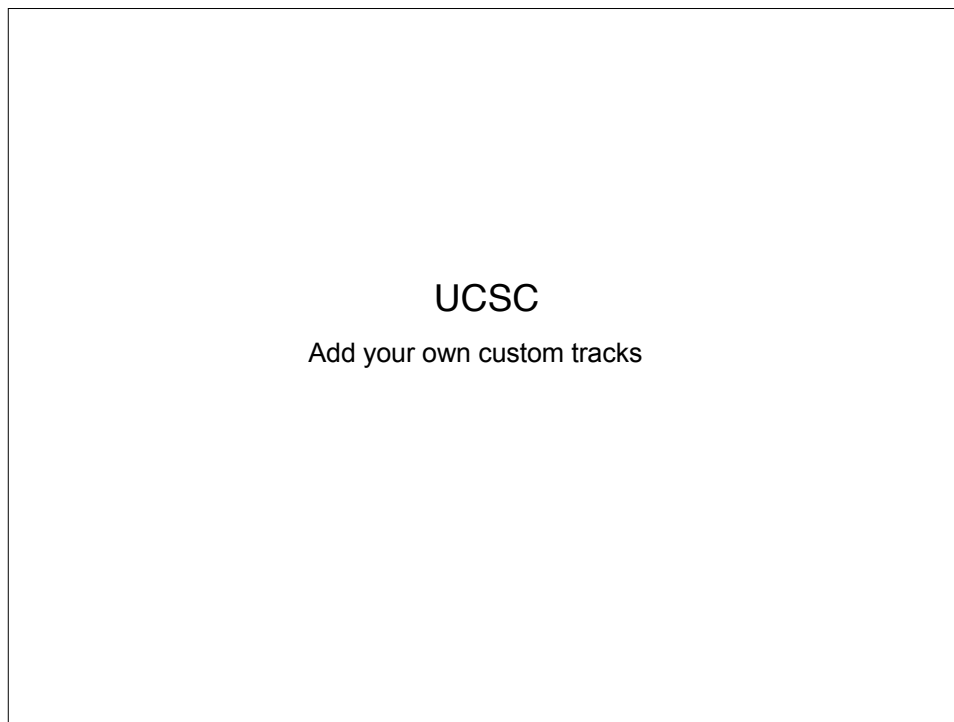
```

001615 N L O C G K L I C K Y 001647
>>>>> | | G | | | | | | | T | >>>>>
635370 aatctgggctgtggaanaactcatctgcacatac 635402

001681 T I I Y A N I S G H L C 001716
>>>>> | | | | | O V O E | | | >>>>>
635436 accatcatctatgctcaagtcaagaacctctgtgc 635471

001768 K D G T S C G S N K V 001800
>>>>> | | | | K | P G | | >>>>>
    
```

Display a menu



Add Your Own Custom Track

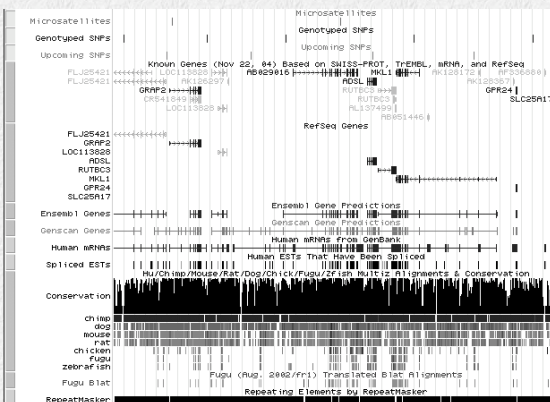
Display your own annotation tracks in the browser using the [procedure described here](#). Annotations may be uploaded from files or pasted into the text box below. You can also paste a URL or a list of URLs into the large text box that refer to files in one of the supported formats.

Click [here](#) to view a collection of custom annotation tracks submitted by Genome Browser users.

Annotation File: no file selected

```
browser position chr22:38496887-39496866
browser hide cytoBand
browser hide stsMap
browser hide gap
browser hide ClonePos
browser full refGene
browser dense mirna
track name="scale" description="our peak"
chr22 38996887 38996888 peak
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
```

Nature Genetics User's Guide,
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 all SNPs that are contained within a RefSeq coding region

NCBI

Identify all the genes between two STS markers

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Mining Genomic Sequence Data

Map Viewer

http://www.ncbi.nlm.nih.gov/mapview/

NCBI NCBI Map Viewer

Genome Taxonomy Entrez BLAST Help

Search for

Now Available! - NCBI's annotation of the **dog** (*Canis familiaris*) genome assembly (build 1.1). The dog is a useful model organism due to extensive genetic diversity and morphological variation within the species and to aggressive breeding practices that have resulted in inbred populations of dogs. [Map Viewer](#) and many [other resources](#) at NCBI now provide a more comprehensive resource for dog.

Click the to BLAST, the to search the group

Mammals 9 organisms

Other Vertebrates 2 organisms

Fungi 11 organisms

Invertebrates

Insects 3 organisms

Nematode 1 organism

Protozoa 1 organism

Plants 8 organisms

See more about [Bacteria](#), [Organelles](#), [Viruses](#)

The Map Viewer supports search and display of genomic information by chromosomal position. Regions of interest can be retrieved by text queries (e.g. gene or marker name) or by sequence alignment (BLAST). View results at the whole genome level, and select what to display in more detail. Multiple options exist to configure your display, download data, navigate to related data, and analyze supporting information using the tools provided. [More...](#)

Display a menu

Entrez Genome view

http://www.ncbi.nlm.nih.gov/mapview/map_search.cgi?taxid=9606&query=D8S1170%20OR%20D8S94

NCBI

PubMed Nucleotide Protein Genome Structure PopSet Taxonomy OMIM Help

Search for assembly

Show related entries

Homo sapiens genome view BLAST search the human genome

build 35.1 statistics

Hits: 1 2 3 4 5 6 7 8 9 10 11 12 13

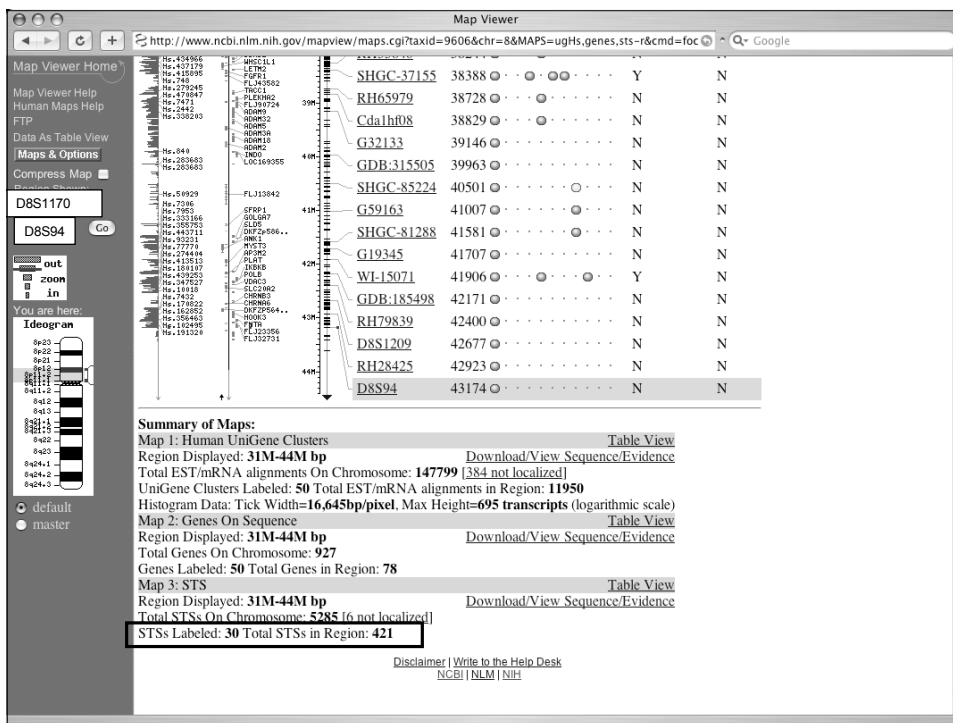
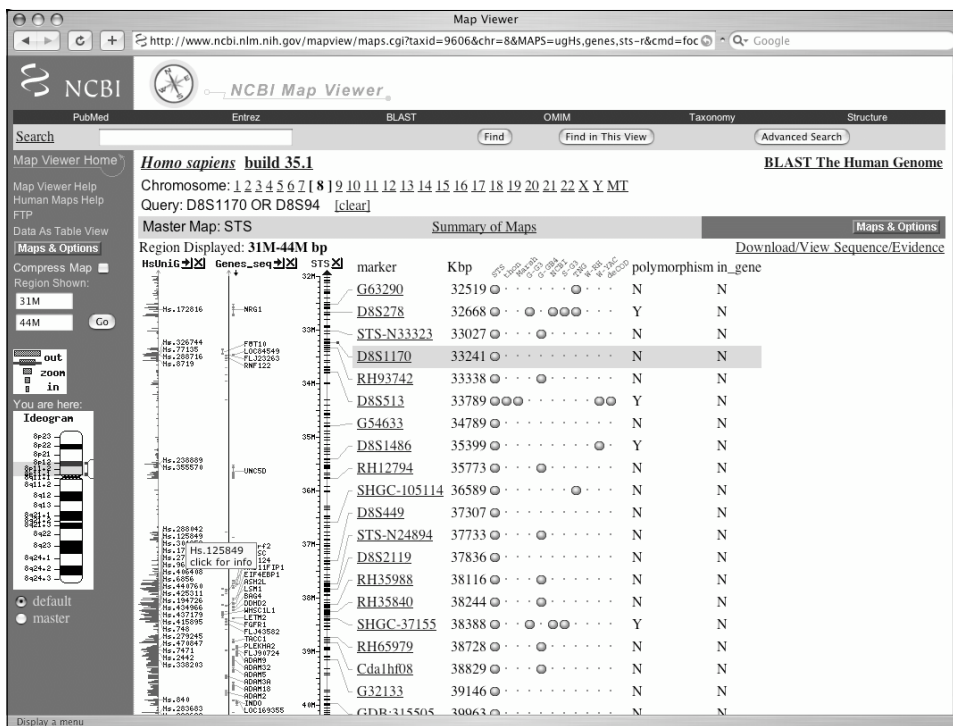
Hits: 14 15 16 17 18 19 20 21 22 X Y HT

Search results for query "D8S1170 OR D8S94": 4 hits

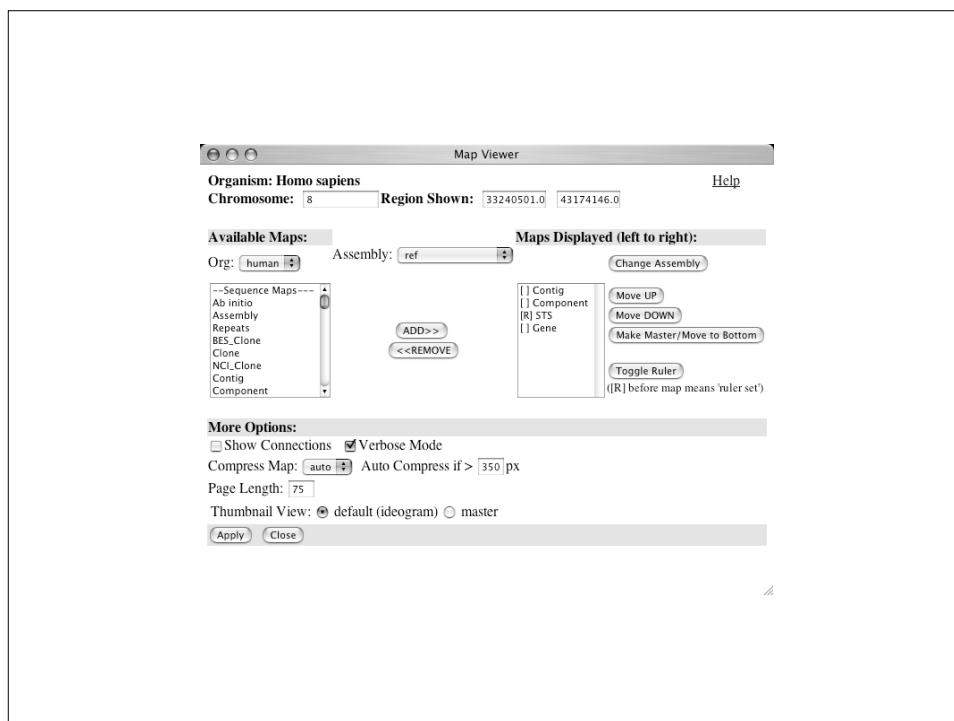
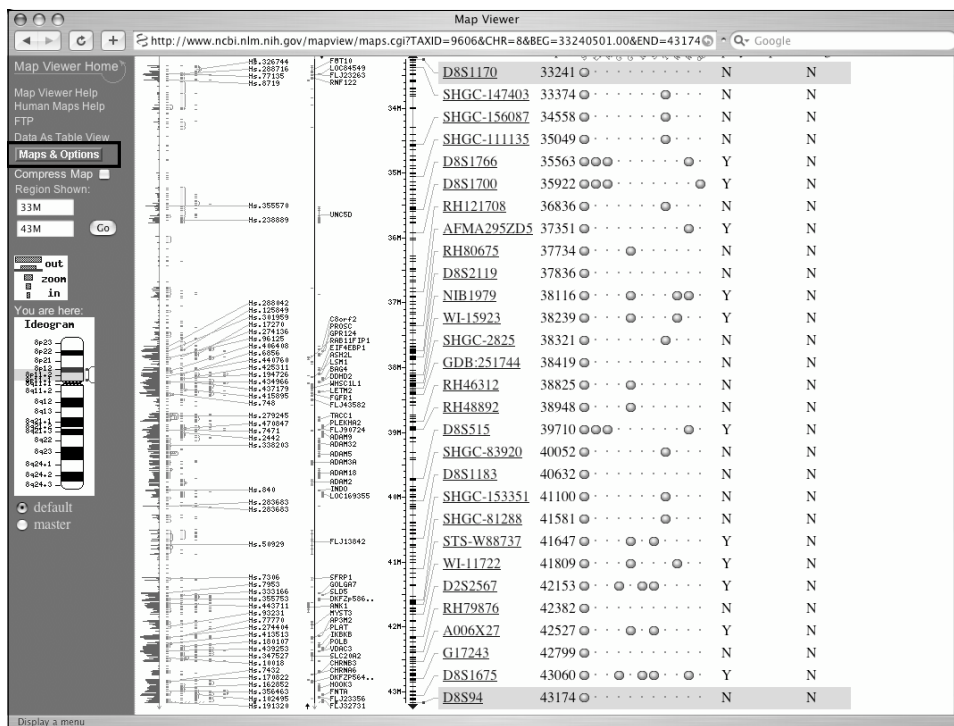
Chr	Assembly	Match	Map element	Type	Maps
8	reference	all matches			
		D8S94	D8S94	STS	STS
		D8S1170	D8S1170	STS	STS
8	Celera	all matches			
		D8S94	D8S94	STS	STS
		D8S1170	D8S1170	STS	STS

Display a menu

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Mining Genomic Sequence Data



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Mining Genomic Sequence Data



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Mining Genomic Sequence Data

Query: D8S1170 OR D8S94 [clear]

Master Map: Genes On Sequence

Region Displayed: 33M-43M bp

Contig	Comp	STS	Genes_seq	Symbol	O	LinkOut	E	Cyto	Description
RC967835..	33,293	2065-114..	+	FUT10	+	sv pr dl ev mm hm	C	8p12	fucosyltransferase
RC991544..	33,493	2065-114..	+	LOC84549	+	sv pr dl ev mm hm	C	8p12	RNA binding pro
RC912640..	33,466	2065-114..	+	FLJ23263	+	sv pr dl ev mm hm	C	8p12	hypothetical prote
RC945779..	33,799	2065-114..	+	RNF122	+	sv pr dl ev mm hm	C	8p12	ring finger protei
RF279073..	33,799	2065-114..	+	MGC1136	+	sv pr dl ev mm hm	C	8p12	hypothetical prote
RC911664..	34,011	2065-114..	+	LOC442384	+	sv dl ev mm	?	8p12	similar to VENT-
NT_007995..	38,286	2065-114..	+	BLP1	+	sv pr dl ev mm hm	C	8p11.23	BBP-like protein
RC007362..	38,286	2065-114..	+	ADAM9	+	OMIM sv pr dl ev mm hm	C	8p11.23	a disintegrin and t
RC007623..	38,493	2065-114..	+	ADAM32	+	sv pr dl ev mm hm	C	8p11.23	a disintegrin and t
RC009120..	38,493	2065-114..	+	ADAM5	+	sv dl ev mm	C	8p11.23	a disintegrin and t
RC010013..	38,799	2065-114..	+	ADAM3A	+	OMIM sv dl ev mm	C	8p21-p12	a disintegrin and t
RC007817..	38,893	2065-114..	+	ADAM18	+	sv pr dl ev mm hm	C	8p11.22	a disintegrin and t
RC108863..	39,000	2065-114..	+	ADAM2	+	OMIM sv pr dl ev mm hm	C	8p11.2	a disintegrin and t
RC105091..	39,193	2065-114..	+	INDO	+	OMIM sv pr dl ev mm hm	C	8p12-p11	indoleamine-pyrr
RC105105..	39,293	2065-114..	+	LOC169355	+	sv pr dl ev mm hm	C	8p11.21	hypothetical prote
RC106113..	39,293	2065-114..	+	FLJ23356	+	sv pr dl ev mm hm	C	8p11.21	hypothetical prote
RC099634..	40,093	2065-114..	+	LOC441347	+	sv pr dl ev mm	E	8p11.21	similar to family v
RC110275..	40,093	2065-114..	+	FLJ32731	+	sv pr dl ev mm hm	C	8p11.1	hypothetical prote

Summary of Maps:

Map 1: Contig [Table View](#)
 Region Displayed: 33M-43M bp [Download/View Sequence/Evidence](#)
 Total Contigs On Chromosome: 18 [8 not localized]
 Contigs Labeled: 1 Total Contigs in Region: 1

Map 2: Component [Table View](#)
 Region Displayed: 33M-43M bp [Download/View Sequence/Evidence](#)
 Total Components On Chromosome: 1200 [8 not localized]
 Components Labeled: 85 Total Components in Region: 85

Map 3: STS [Table View](#)
 Region Displayed: 33M-43M bp [Download/View Sequence/Evidence](#)
 Total STSs On Chromosome: 5285 [6 not localized]
 STSs Labeled: 125 Total STSs in Region: 372

Map 4: Genes On Sequence [Table View](#)
 Region Displayed: 33M-43M bp [Download/View Sequence/Evidence](#)
 Total Genes On Chromosome: 927
 Genes Labeled: 74 Total Genes in Region: 74

Download Sequence Region

http://www.ncbi.nlm.nih.gov/mapview/seq_reg.cgi?chr=8&from=39720414&to=39814932

Homo sapiens Genome (build 35.1)
 Region to retrieve (in chromosome coordinates):
 Chromosome: 8 Strand: plus
 from: 39720414 adjust by: -OK
 to: 39814932 adjust by: +OK [Change Region/Strand](#)

Sequence Format: FASTA

This chromosome region corresponds to the contig region(s):

Contig	start	stop	strand
NT_007995.14	9921646	10016164	+

+ Display Save to Disk View Evidence ModelMaker

Sequence download (dl)

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Mining Genomic Sequence Data

Evidence Viewer

http://www.ncbi.nlm.nih.gov/sutils/ev.cgi?contig=NT_007995.14&gene=ADAM2&lid=2515

Exon 12
NT_007995.14: 9947300-9947483 minus strand
BC034957: 996-1179
AJ133005: 1033-1216
U38805: 1097-1280
U52370: 1036-1219
NM_001464: 1097-1280
BC064547: 676-859
X99374: 1029-1212

preceding intron phase: 2
flank-gtggattgttag--flank

frame 1 (1):
9947483 H F S Ø V K I F S N C S F E D F A H F I S K Q K S Q C L H N
996 TCATTTCAGTGTGTGTGAGATCTTTTAACTCAGCTTCGAAAGCTTTCACATTTTATTCAGAGCAGAGTCCGAGTGTCTTCAGAA
1033 H F S Ø V K I F S N C S F E D F A H F I S K Q K S Q C L H N
1097 H F S Ø V K I F S N C S F E D F A H F I S K Q K S Q C L H N
1036 H F S Ø V K I F S N C S F E D F A H F I S K Q K S Q C L H N
1097 H F S Ø V K I F S N C S F E D F A H F I S K Q K S Q C L H N
676 H F S Ø V K I F S N C S F E D F A H F I S K Q K S Q C L H N
1029 H F S Ø V K I F S N C S F E D F A H F I S K Q K S Q C L H N

frame 1 (1):
1086 Q P R L D P F F K Q Q A V C G N A K L E A G E E C D C G T E
1123 TCACCTCCTTAGATCTTTTTCAGAGCAGAGTGTGTGATTCAGAGCTGGAAGCAGAGAGGAGTGTGACTGTGGAGCTGA
1107 Q P R L D P F F K Q Q A V C G N A K L E A G E E C D C G T E
1126 Q P R L D P F F K Q Q A V C G N A K L E A G E E C D C G T E
1187 Q P R L D P F F K Q Q A V C G N A K L E A G E E C D C G T E
766 Q P R L D P F F K Q Q A V C G N A K L E A G E E C D C G T E
1119 Q P R L D P F F K Q Q A V C S N A K L E A G E D E F D C G T E

intron phase: 0
flank-gtggattgttagctactg

frame 1 (1):
9947303 ACAG
1176 ..
1213 ..
Q

Display a menu

Evidence viewer (ev)

Model Maker

http://www.ncbi.nlm.nih.gov/mapview/modelmaker.cgi?contig=NT_007995.14&gene=ADAM2&lid=2515

Model Maker (Make Your Own Model by selecting an evidence help legend)
exon "set" and/or add/remove individual putative exons for inclusion in your model

Organism: Homo sapiens Chromosome: 8 Contig: NT_007995.14 Locus: ADAM2

Evidence:
10016164<<< mv sv ev seq >>>9921646

minus strand

change strand

expand ESTs

BC034957.1
BC064547.1
U38805.1
U52370.1
NM_001464.1

Putative exons (graphic view):

Your model: clear

BC064547.1

CATCTCGGCTTCCAAGTCCCTGTAACCACCAACTGCCATTATCCGGCTGGGA
CCCAGCATTCAAGCATGTGGCGCTCTGTTCTGCTCAGCGGCTCGCGGG
CTCGGATCGACATTAATTTGATAGTTACTCTGCAAAATACACTCCCGAGA
AAATACGGTCAATAAAGGAAGGAATTGAATCGCAGCATCTACAAAAATTGT

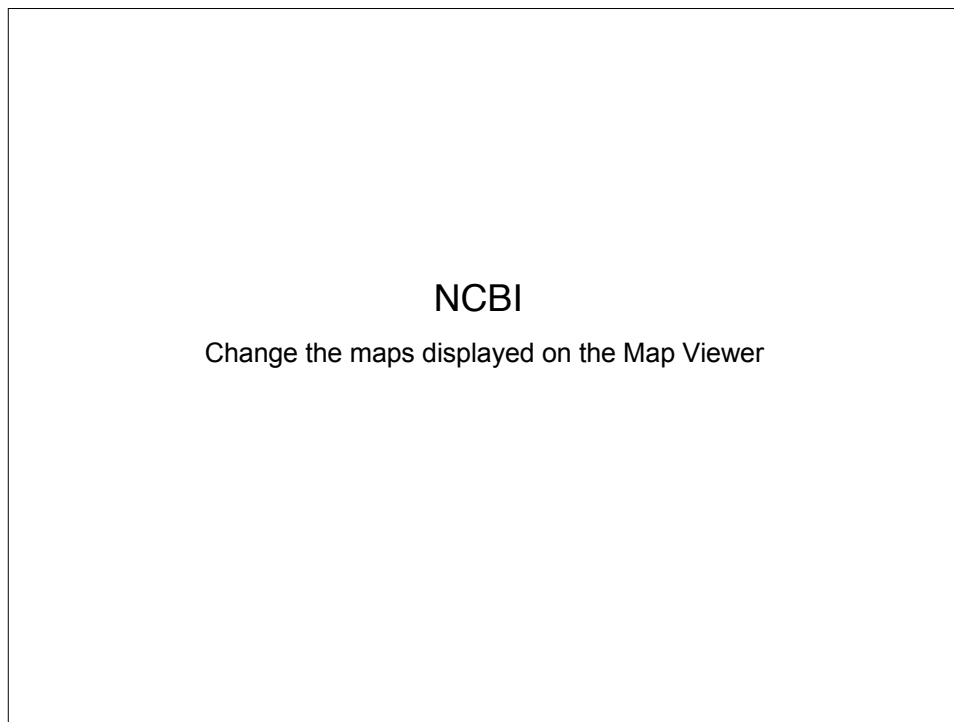
ORF Finder

Frame 1, ORF= 90 CDD Frame 2, ORF= 29 CDD Frame 3, ORF= 579 CDD

Putative exons (table view):

Exon	Sequence	Start	End	Strand
1	AG GCA	10016303	10016039	GTA GT >> 3
2	10016164 GT [CAT...CC ATG	10016093	10016039	GTA GT >> 3
3	1 or 2 <= AG ATT	10015120	10015044	CAG GT >> 4
4	3 <= AG GCA	10011907	10011852	AAA GT >> 5
5	4 <= AG AAA	10002805	10002727	CAG GT >> 6
6	5 <= AG AAT	9999570	9999494	CAG GT >> 7
7	6 <= AG GGG	9999078	9998910	GAG GT >> 8 or 9 or 12
8	7 <= AG CCA	9987374	9987318	TTG GT >> 9
9	7 or 8 <= AG TAT	9966648	9966577	GCT GT >> 10
10	9 <= AG ATT	9966159	9965993	TGT GT >> 11
11	10 <= AG TTA	9964963	9964882	CTG GT >> 12
12	7 or 11 <= AG CAC	9955069	9954933	AAT GT >> 13

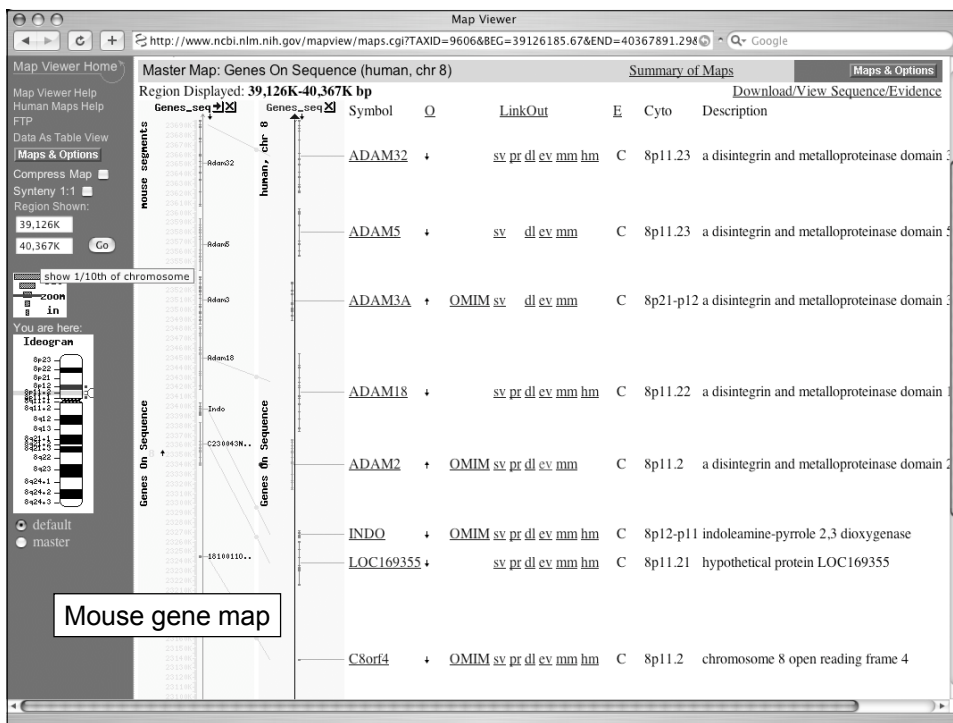
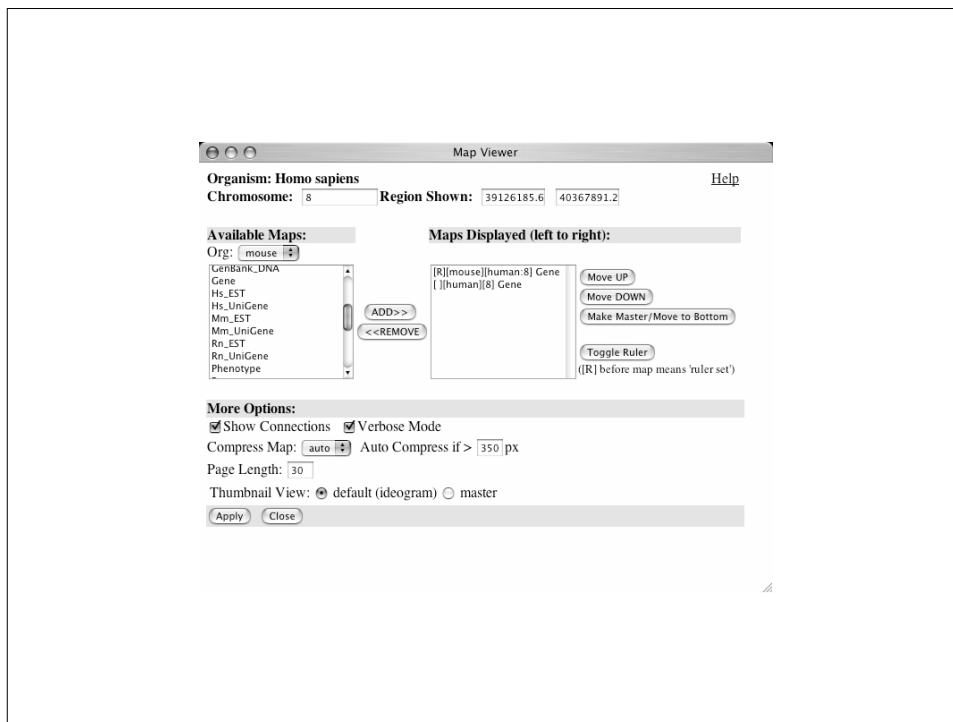
Model maker (mm)



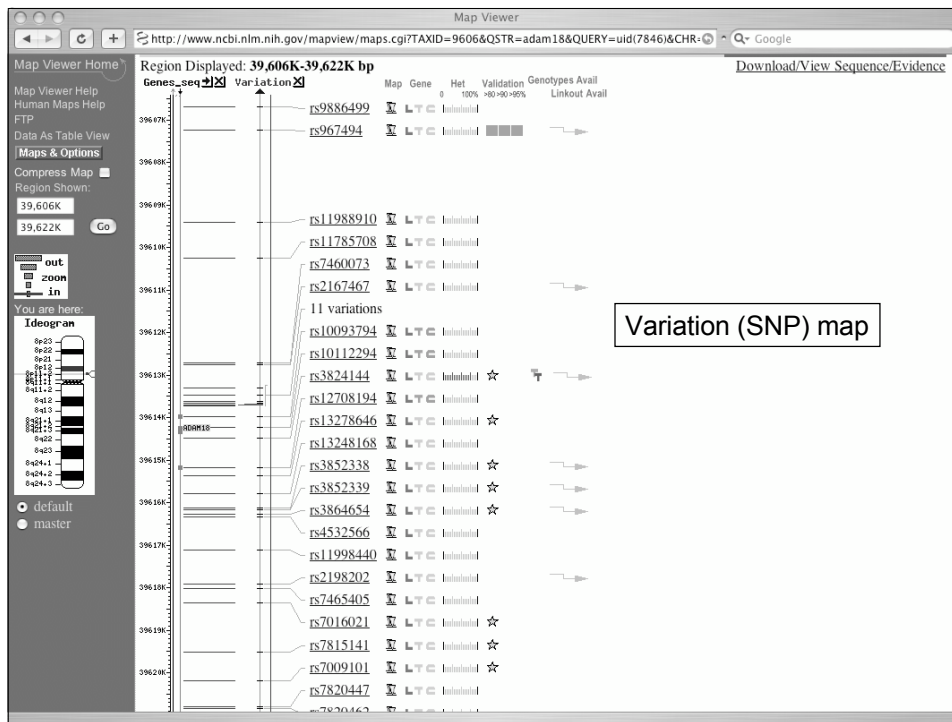
The screenshot shows the NCBI Map Viewer interface. The browser address bar displays the URL: <http://www.ncbi.nlm.nih.gov/mapview/maps.cgi?TAXID=9606&CHR=8&BEG=33240501.00&END=43174>. The interface includes a navigation pane on the left with options like 'Map Viewer Home', 'Map Viewer Help', and 'Human Maps Help'. The main area shows a genomic map with various features and labels. On the right, a list of gene annotations is displayed, including:

Gene Name	Accession	Species	Gene Type	Gene Description
RAB11FIP1	+	OMIM	sv pr dl ev mm hm	C 8p11.22 RAB11 family inte
ASH2L	+	OMIM	sv pr dl ev mm hm	C 8p11.2 ash2 (absent, small
BAG4	+	OMIM	sv pr dl ev mm hm	C 8p12 BCL2-associated at
WHSC1L1	+	OMIM	sv pr dl ev mm hm	C 8p11.2 Wolf-Hirschhorn sy
FGFR1	+	OMIM	sv pr dl ev mm hm	C 8p11.2-p11.1 fibroblast growth fa
TACC1	+	OMIM	sv pr dl ev mm hm	C 8p11 transforming, acidic
PLEKHA2	+	OMIM	sv pr dl ev mm hm	C 8p11.23 pleckstrin homolog
ADAM9	+	OMIM	sv pr dl ev mm hm	C 8p11.23 a disintegrin and m
ADAM32	+	sv pr dl ev mm hm	C 8p11.23 a disintegrin and m	
ADAM5	+	sv dl ev mm hm	C 8p11.23 a disintegrin and m	
ADAM3A	+	OMIM	sv dl ev mm hm	C 8p21-p12 a disintegrin and m
ADAM18	+	sv pr dl ev mm hm	C 8p11.22 a disintegrin and m	
ADAM2	+	OMIM	sv pr dl ev mm hm	C 8p11.2 a disintegrin and m
LOC169355	+	sv pr dl ev mm hm	C 8p11.21 hypothetical protei	
FLJ13842	+	sv pr dl ev mm hm	C 8p11.21 hypothetical protei	
SFRP1	+	OMIM	sv pr dl ev mm hm	C 8p12-p11.1 secreted frizzled-rel
DYKFP586M1819	+	OMIM	sv pr dl ev mm hm	C 8p11.21 putative lysosposl
	+	OMIM	sv pr dl ev mm hm	C 8p11.1 ankyrin 1, erythrocy
	+	OMIM	sv pr dl ev mm hm	C 8p11 MYST histone acet
	+	OMIM	sv pr dl ev mm hm	C 8p12 plasminogen activa
	+	OMIM	sv pr dl ev mm hm	C 8p11.2 inhibitor of kappa l
	+	OMIM	sv pr dl ev mm hm	C 8p11.2 polymerase (DNA
	+	OMIM	sv pr dl ev mm hm	C 8p12-q21 solute carrier family
	+	OMIM	sv pr dl ev mm hm	C 8p11.2 cholinergic receptor
DKFZP564A022	+	sv pr dl ev mm hm	C 8p11.21 hypothetical protei	
HOOK3	+	OMIM	sv pr dl ev mm hm	C 8p11.21 hook homolog 3 (E
FLJ32731	+	sv pr dl ev mm hm	C 8p11.1 hypothetical protei	

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NCBI

Find a chicken homolog of a human protein

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BLAST the Chicken Genome

http://www.ncbi.nlm.nih.gov/genome/seq/GgaBlast.html

NCBI Home > Genomic Biology > Chicken Genome Resources > BLAST

Search

BLAST the Chicken Genome

Blast your sequence against Chicken specific sequences

Database: Program:

use MegaBLAST

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

```
>gi55743080[ref|NP_001455.3] a disintegrin and metalloproteinase domain
2 proprotein [Homo sapiens]
MWRVLELLSGLCGLRMDSNFDSLVPQITVPEKIRSIKESQASYKIVIEGKPYTVNLMQK
NFLPHNF
RVYSYSGTGIMKPLDQDFQNFCHYQYIEGPKSVMMVSTCTGLRGLQFENVSYGIEPL
ESSVGFHVI
```

Optional parameters

Expect: Filter: Descriptions: Alignments:

Advanced options:

Show positions of the BLAST hits in the chicken genome using the Entrez Genomes MapViewer

Taxonomy reports

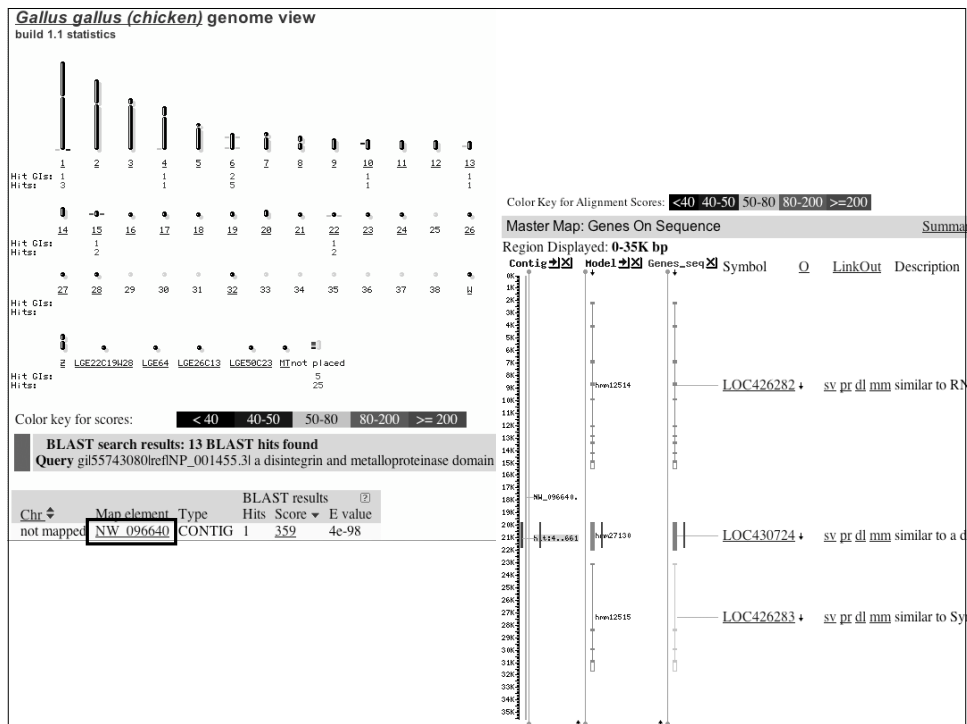
Distribution of 40 Blast Hits on the Query Sequence

Mouse-over to show details and scores. Click to show alignments

Color Key for Alignment Scores

<40 40-50 50-80 80-200 >200

```
>ref|NW_036440.1|GgaNW_041444.1| Gallus gallus chromosome Un genomic contig, whole genome shotgun
Sequence
Length = 35588
Score = 359 bits (922), Expect = 4e-98
Identities = 215/681 (31%), Positives = 345/681 (50%), Gaps = 22/681 (3%)
Frame = +1
Query: 4 VLFLSLGLGLLMDNSDFSLVQIT-----VSRIRSIIRKIESQASVIVIEKRPITV 58
          VL +L GL G + +G P+ +T VP + + S + + + + + + + + + + + + + + + +
Sbjct: 19795 VLVLVLLGLVCCPTTDDSGSFLVDMVTVFPGD--SPRADNPLTVSYLVQVEGRPVL 19971
Query: 59 NLR-QNPLFRFRVSYSGTGIKELDDQDFQVYQYISGVKSVVYVSTC--TGLRG 116
          L +R F + + + G + + D+ C YGG + +G P + + + + + + + + + + + + + + +
Sbjct: 19972 ELRPFKGLASRPFLLVTDGCAKREQVTVQNCPIYQGVGSGFGLVALGCTCGLAG 20151
Query: 117 VLQFENVSYGIEPLSSVGFERNIVYVYKRAKVS---LVNNDIESRDLSPKLSQVDP 172
          VL D + + Y IEP+ F + + + + + + + + + + + + + + + + + + + + + + + +
Sbjct: 20152 VLNRHGPTIEIPIIDDFAFQMLFRMEADSDWPGFCULPREEIQVQVTLVPGQAPWT 20331
Query: 173 QQDF-----ARYIENNVIVEKQLTNHGGDPTVVAQKVFQILGILNAINVSPFITII 224
          + + + + + + + + + + + + + + + + + + + + + + + + + + + + + + + +
Sbjct: 20332 EDKTYLKNWHTFRVYLVVVVDVFRVDRHESVLRQLEVVNIGDGLVQGLVLP 20511
Query: 225 LSELELWIDENIATTCENELLEITLWKTSTYLVLR--PDAVPLVYREKSNVYATQ 283
          L LER+ + N I ? A+ + L F R+ S S L R SD A L + + + + + + + + + + +
Sbjct: 20512 LVGLWITRNSPINITASATLAFDRWRESDIPFQHTALFAPQKRELATL 20691
Query: 284 GENCANYAGGVVHFRPTISLESVLIAQLLSELSNGITDIDINKQCQCAVNCIPEAI 343
          G +CD + + V + + L S V L + +G+ +D+ C+C CIN
Sbjct: 20692 GSICDRQNSAAVDVYVNR-RLSSPVTTFVHGLNGLMHBDE--RCKCRKRCIMYSE- 20862
Query: 344 HFGVQVIFNGSDFEDFAHFLSKQKQCLMNPRLDFFPF--QAVCGNLAELGRCDCOT 402
          S F+ + + + + + + + S CL+ P L + + + + + + + + + + + + + + + + + +
Sbjct: 20863 --SDTANFSDVITDITDGLRGGG--LVDQALGAGITTLREYICGRIEVSQKCKGS 21033
Query: 403 EQDCALIGETCCDIATCRPKAGSAGPCCNCLFMSKERKCRPFSEKDLPEYNGSS 462
          + DC + CC C AGS CA G CC+ C + + + + + + + + + + + + + + + + + +
Sbjct: 21034 REDCRN--DFCC--PFCCTLTAGVCAKCKGCGQILFAGTLCAARTGDCDLPEYNGTS 21204
Query: 463 AGCFENHVYQHPFCGLNQCIDVCGSGDQCTGFGREYFGRFCEYSLNLRKTVS 522
          C E+ + + G RC + + C G C S RGC FQ+ + P EC+ +R+ + D
Sbjct: 21205 PWCQEDLVYDQGTFCDEGAY-CYKCKSSKQKRLGQARFAPLECFKRVHTGDRF 21381
Query: 523 GNCISDS--GYTQCEADNLQCKLICKYKFLLLQIPRATIIYANISGLCIAVPAEDR 581
          GRCD + + + + + + + + + + + + + + + + + + + + + + + + + + + + + +
Sbjct: 21382 GNCGRNIRITFGEVVALCGKQCEVVELLQGHITVLIOTFAGKCKGLVYELDV 21561
Query: 582 ADSGRNWKDGTCCGSRVCRMQVSSGLGDCDTRDCHGRVCCNKRKCHRYAFLP 641
          S + + + + + + + + + + + + + + + + + + + + + + + + + + + + + +
Sbjct: 21562 PTDHWAGVEDDTCCGSRKICRTCTNISVLNVDCHITRCHGRVCCNKRKCHRYWAP 21741
Query: 642 PDCVQSDLVFGGSIDSGNFP 662
          P C + + + + + + + + + + + + + + + + + + + + + + + + + + + + +
Sbjct: 21742 FYCEBQDF--GGVSDGPPP 21795
```



Ensembl

Identify genes and SNPs in a chromosomal band

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Ensembl Genome Browser

Search all species for with

About Ensembl

Ensembl is a joint project between EMBL - EBI and the Sanger Institute to develop a software system which produces and maintains automatic annotation on metazoan genomes. Ensembl is primarily funded by the Wellcome Trust.

This site provides free access to all the data and software from the Ensembl project. Click on the species buttons to the right to browse the data.

Access to all the data produced by the project, and to the software used to analyse and present it, is provided free and without constraints. Some data and software may be subject to third-party constraints [\[details\]](#).

For all enquiries, please contact the Ensembl [HelpDesk](#) (helpdesk@ensembl.org).

Help and documentation

- Take the [Ensembl tour](#), go through a step-by-step [worked example](#), or read [these papers](#).
- For help on any web page click [Help](#).
- There is also an [index](#) of help pages, and a set of guided [How do I...?](#) trails.

Display your own data in Ensembl

Questions or suggestions? Try the [Documentation](#) (includes tutorial on direct data access & instructions for installing Ensembl on your own site)

Try the site map as a good starting point for exploring what Ensembl has to offer

Species - Ensembl v27

Human	NCBI 35	Dec 04
Mouse	NCBI m33	Dec 04
Zebrafish	WTSL Zv4	Sep 04
Rat	pre! RGSC 3.1	Jul 04
Chicken	WASHUC1	Jul 04
Mosquito	MOZ 2	Apr 04
Fugu	Fugu v2.0	May 04
Fruittly	BOGF 3.1	Dec 04
Chimp	CHIMP1	May 04
Honeybee	Amel1.1	Sep 04
Tetraodon	TETRAODON7	Sep 04
Dog	BROADD1	Dec 04
C. elegans	WS 130	Dec 04
Cow	pre! Btau_1.0	
X.tropicalis	pre! JGI3	
Opossum	pre! BROAD0.5	

Data

Sequence similarity searches

Batch data/sequence retrieval

Vertebrate Genome Annotation (VEGA)

Access to whole genome shotgun data (includes additional species)

Download Ensembl data via FTP

Have you tried?

[Ensembl Assembly Preview Browser](#)
Opossum

Go to "http://pre.ensembl.org/Monodelphis_domestica/"

Ensembl Genome Browser

Search for with

Display Chr From To

Retrieve a sequence

Search your sequence

Advanced data retrieval tool

Finishing the Genome

The International Human Genome Sequencing Consortium have published their scientific analysis of the finished human genome. [Nature 431, 931 - 945 \(21 October 2004\)](#)

[WT Sanger Institute Press Release](#)

Browse a Chromosome

Current Release 27.35a.1

This release is based on the NCBI 35 assembly of the human genome.

View the [status](#), [history](#) of the human assemblies.

Last Update: 14-12-2004

Ensembl gene predictions: 24194 (incl. 1978 pseudogenes)

GenScan gene predictions: 68101

Ensembl gene exons: 245215

Ensembl gene transcripts: 35838

ENCODE regions

The ENCODE (ENCYclopedia OF DNA Elements) project aims to find functional elements in the human genome. More information about the [ENCODE resources](#) at Ensembl.

Documentation & Help

About Ensembl

For context-sensitive help on any web page click

Questions or suggestions? Try

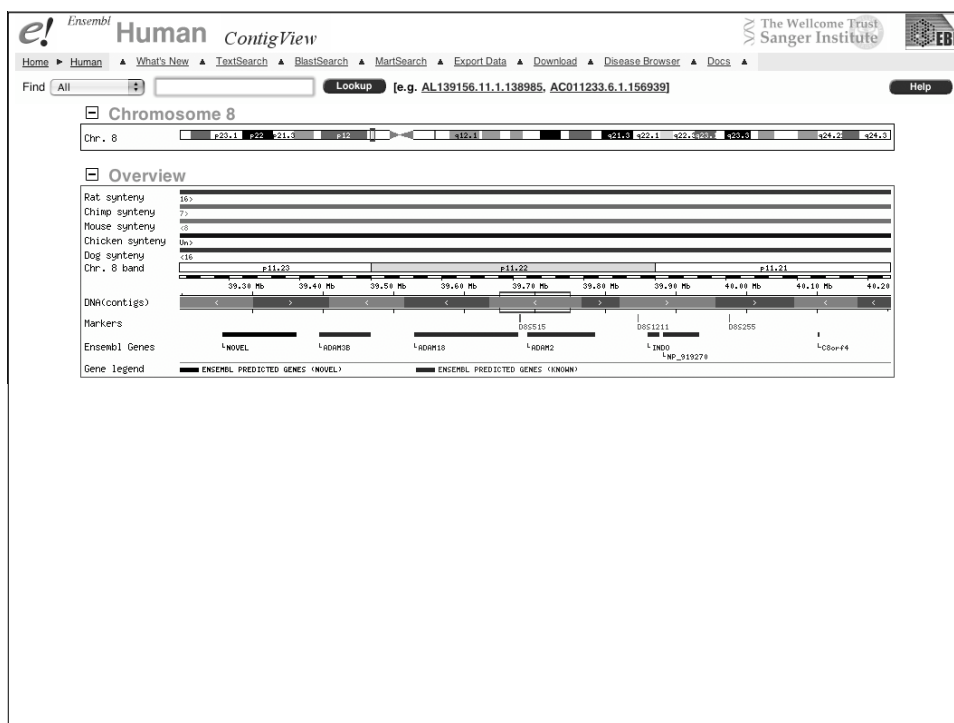
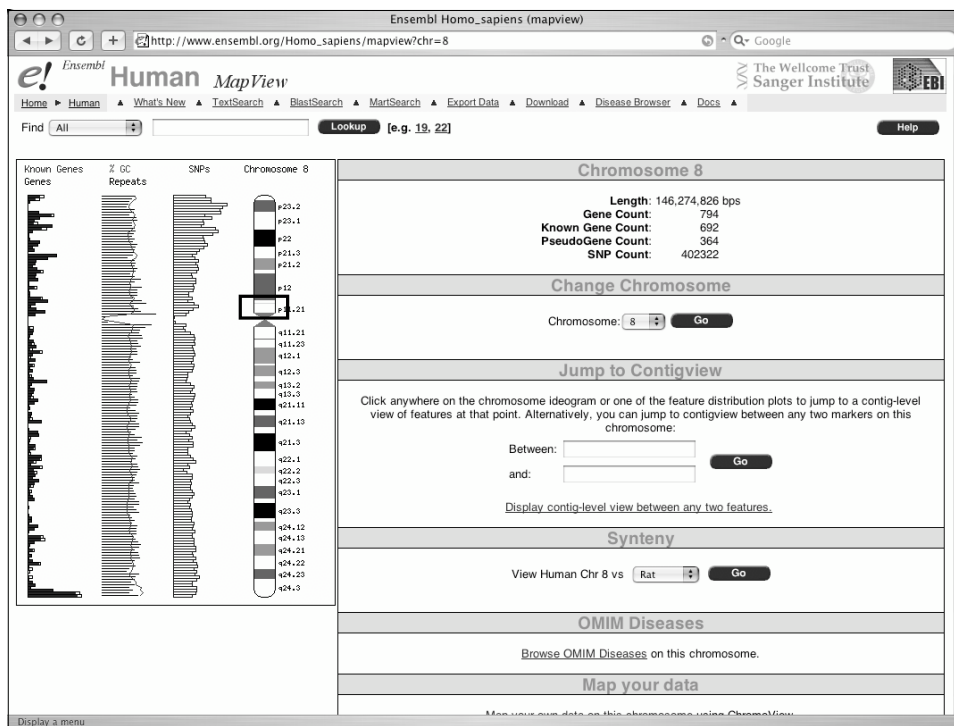
Documentation (includes tutorial on direct data access & instructions for installing Ensembl on your own site)

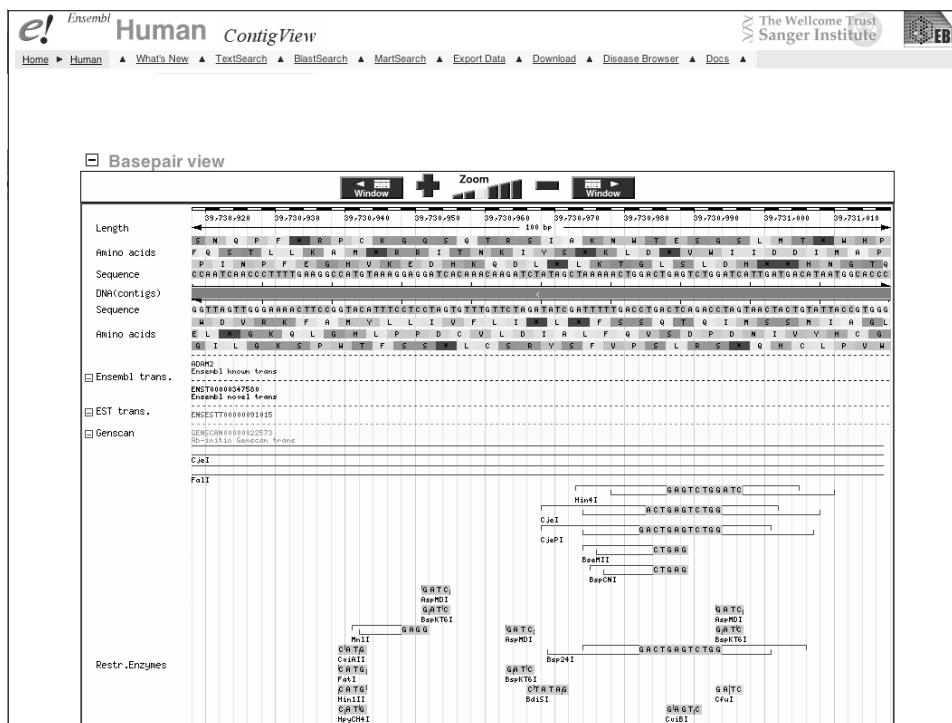
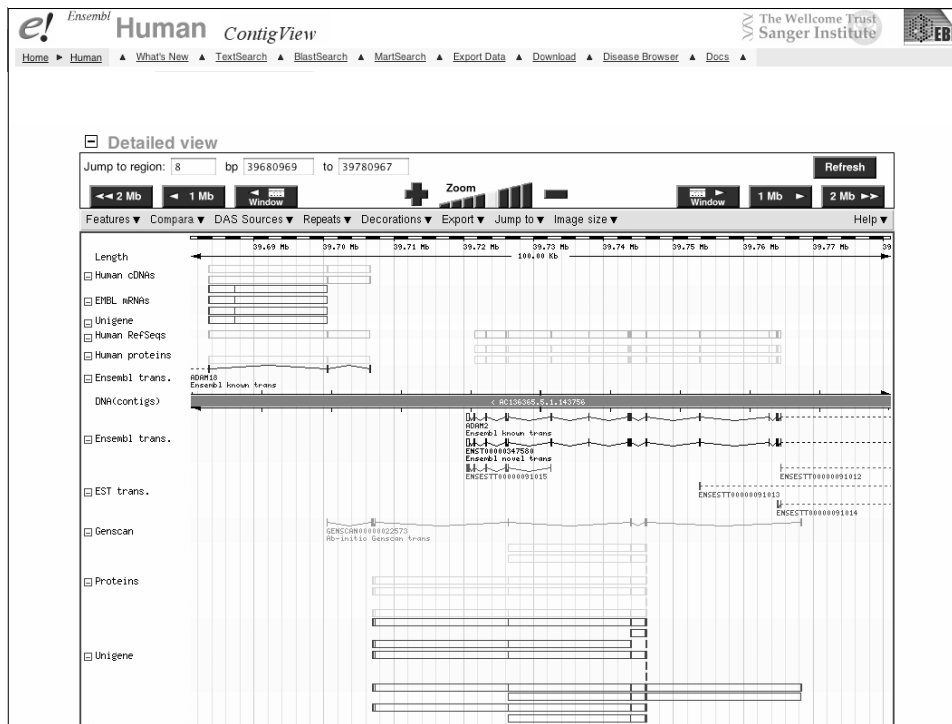
Ensembl Links and Site Map

Other Species

Mosquito	Honeybee	C. elegans
Dog	Zebrafish	Fruittly
Fugu	Chicken	Mouse
Chimp	Rat	Tetraodon

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Ensembl

Change the tracks displayed on the ContigView

The screenshot displays the Ensembl Human Genome Browser interface. At the top, the browser title is "Ensembl Human Genome Browser (ContigView)". The address bar shows the URL: http://www.ensembl.org/Homo_sapiens/contigview?c=8:39730968&w=99999&panel_zoom=off. The navigation bar includes links for Home, Human, What's New, TextSearch, BlastSearch, MartSearch, Export Data, Download, Disease Browser, and Docs. A search bar contains the text "Find [All] [e.g. AL139156.11.1.138985, AC011233.6.1.156939]".

The main content area is titled "Chromosome 8" and shows a genomic track. Below this, the "Overview" section is visible. The "Detailed view" section is active, showing a region from 39,680,969 bp to 39,780,967 bp. The interface includes navigation controls for zooming and jumping to different regions. A list of features is shown on the left, with checkboxes for various tracks. The main track displays genomic data, including SNPs, genes, and other features. A specific SNP, rs408088, is highlighted, and its properties are shown in a pop-up window:

SNP: rs408088
SNP properties
View in LDView
bp: 39720624
status: -
SNP type: snp
ambiguity code: R
alleles: A/G
dbSNP: rs408088
Type: 3PRIME_UTR

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Ensembl Homo_sapiens (snpview)

http://www.ensembl.org/Homo_sapiens/snpview?snp=rs408088&source=dbSNP&c=8.39720624

Ensembl SNP Report

SNP	rs408088 (dbSNP123)
Synonyms	HGVbase SNP001766036
Validation Status	Unknown
Alleles	A/G (ambiguity code: R)

This SNP is currently mapped to the following genomic locations:

Genomic location (strand)	Transcript: start-end	Translation: start-end	Peptide allele	Consequence
8: 39720624-39720624 (-1)	ENST00000265708: 2426-2426	ENSP00000265708: n/a		3PRIME_UTR
	ENST00000347580: 2322-2322	ENSP00000343854: n/a		3PRIME_UTR

SNP neighbourhood

Ensembl Homo_sapiens (geneview)

http://www.ensembl.org/Homo_sapiens/geneview?gene=ENSG00000104755

Ensembl Human GeneView

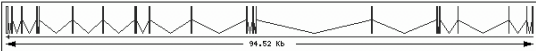
Find (All) [e.g. ENSG00000139618, BRCA2]

Ensembl Gene Report

Gene	ADAM2 (HUGO ID) (to view all Ensembl genes linked to the name click here)												
Ensembl Gene ID	ENSG00000104755												
Genomic Location	View gene in genomic location: 39720414 - 39814932 bp (39.7 Mb) on chromosome 8 This gene is located in sequence: AP005902.2.1.149577												
Description	ADAM 2 precursor (A disintegrin and metalloproteinase domain 2) (Fertilin beta subunit) (PH-30) (PH30). (Source:Uniprot/SWISSPROT;Acc:Q99965)												
Prediction Method	Genes were annotated by the Ensembl automatic analysis pipeline using either a GeneWise model from a human/vertebrate protein, a set of aligned human cDNAs followed by GenomeWise for ORF prediction or from Genscan exons supported by protein, cDNA and EST evidence. GeneWise models are further combined with available aligned cDNAs to annotate UTRs.												
Sequence Markup	View genomic sequence for this gene with exons highlighted												
Export Data	Export gene data in EMBL, GenBank or FASTA												
SNP information	View information about variations on this gene.												
Transcript Structure	<p>1: ADAM2 (ENST00000265708) [Transcript information] [Exon information] [Protein information]</p> <p>2: ENST00000347580 [Transcript information] [Exon information] [Protein information]</p>												
	<p>The following gene(s) have been identified as putative orthologues by reciprocal BLAST analysis:</p> <table border="1"> <thead> <tr> <th>Species</th> <th>Type</th> <th>dNdS</th> <th>Gene identifier</th> </tr> </thead> <tbody> <tr> <td><i>Canis familiaris</i></td> <td>UBRH</td> <td>---</td> <td>ENSCEF00000005797 (Novel Ensembl prediction) [MultiContigView] [Align] No description</td> </tr> <tr> <td><i>Danio rerio</i></td> <td>RHS</td> <td>---</td> <td>ENSDARG000000010070 (Novel Ensembl prediction) [MultiContigView] [Align] No description</td> </tr> </tbody> </table>	Species	Type	dNdS	Gene identifier	<i>Canis familiaris</i>	UBRH	---	ENSCEF00000005797 (Novel Ensembl prediction) [MultiContigView] [Align] No description	<i>Danio rerio</i>	RHS	---	ENSDARG000000010070 (Novel Ensembl prediction) [MultiContigView] [Align] No description
Species	Type	dNdS	Gene identifier										
<i>Canis familiaris</i>	UBRH	---	ENSCEF00000005797 (Novel Ensembl prediction) [MultiContigView] [Align] No description										
<i>Danio rerio</i>	RHS	---	ENSDARG000000010070 (Novel Ensembl prediction) [MultiContigView] [Align] No description										

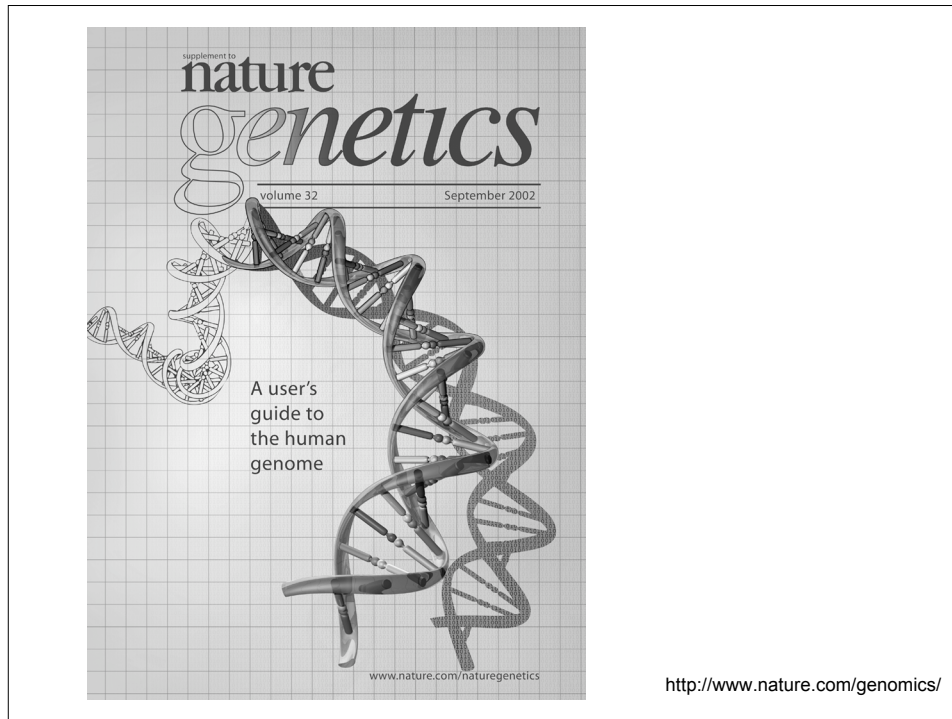
Ensembl Homo_sapiens (geneview)
http://www.ensembl.org/Homo_sapiens/geneview?gene=ENSG00000104755

Transcript/Translation Summary

ADAM2	Stable ID: ENSG00000265708 Exons: 21 Transcript length: 2636 bp Translation length: 735 residues [Transcript information] [Exon information] [Protein information]
Similarity Matches	This Ensembl entry corresponds to the following database identifiers: AFFY HG Focus: 207664_at AFFY HG U133 PLUS 2: 207664_at AFFY HG U133A: 207664_at AFFY HG U133A 2: 207664_at AFFY HG U95Av2: 32298_at AFFY U133 X3P: g11497606_3p_at EMBL: AJ133005 [align] BC034957 [align] U38805 [align] U52370 [align] X99374 [align] HUGO: Search GeneCards for ADAM2 LocustLink: 2515 [align] MIM: 601533 Protein ID: AAC51110.1 [align] AAD04206.1 [align] AAH34957.1 [align] CAA67753.1 [align] CAB40813.1 [align] RefSeq: NM_001464 [Target %id: 99; Query %id: 99] [align] NP_001455 [Target %id: 99; Query %id: 99] [align] Uniprot/SWISSPROT: A002_HUMAN [Target %id: 100; Query %id: 100] [align]
GO	The following GO terms have been mapped to this entry via UniProt: GO:0004222 [metalloendopeptidase activity] IEA GO:0005178 [integrin binding] TAS GO:0005515 [protein binding] IEA GO:0005887 [integral to plasma membrane] TAS GO:0006508 [proteolysis and peptidolysis] IEA GO:0007155 [cell adhesion] IEA GO:0007342 [fusion of sperm to egg plasma membrane] TAS
InterPro	IPR002870 Metalloendopeptidase M12B - [View other EnsEMBL genes with this domain] IPR001762 Disintegrin - [View other EnsEMBL genes with this domain] IPR001590 Metalloprotease ADAM/reprolysin M12B - [View other EnsEMBL genes with this domain]
Protein Family	ENSF00000000082 : ADAM PRECURSOR A DISINTEGRIN AND METALLOPROTEINASE DOMAIN This cluster contains 23 Ensembl gene member(s)
Transcript Structure	

Online 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 Tour
<http://www.ensembl.org/Docs/enstour/>
- The NCBI Handbook
<http://www.ncbi.nlm.nih.gov/books/bv.fcgi?call=bv.View..ShowSection&rid=handbook>



References

- Current Protocols in Bioinformatics
UNIT 1.4: The UCSC Genome Browser
UNIT 1.5: Using the NCBI Map Viewer to Browse Genomic Sequence Data
Access through <http://nihlibrary.nih.gov/ResearchTools/OnlineJournals.htm>
- UCSC
Hsu F *et al.* The UCSC Proteome Browser. *Nucleic Acids Res.* 2005. 33:D454-8.
Karolchik D *et al.* The UCSC Table Browser data retrieval tool. *Nucleic Acids Res.* 2004. 32:D493-6.
Karolchik D *et al.* The UCSC Genome Browser Database. *Nucleic Acids Res.* 2003. 31:51-4.
- NCBI
Wheeler DL *et al.* Database resources of the National Center for Biotechnology Information. *Nucleic Acids Res.* 2005:D39-45.
- Ensembl
Hubbard T *et al.* Ensembl 2005. *Nucleic Acids Res.* 2005. 33:D447-53.
Hammond MP, and Birney E. Genome information resources - developments at Ensembl. 2004. *Trends Genet.* 20:268-72.
Birney E *et al.* An overview of Ensembl. 2004. *Genome Res.* 14:925-8.