

- Move from a **gene** to a **location** display with tabs
- BLAST** a sequence
- Mine data with **BioMart**
- Query using our **Perl API**

Try our **tools**:

- Assembly Converter
- ID History Converter
- Variant Effect Predictor

- View whole genome **alignments**
- Configure this page to customise your view

- Conserved sequence from analysis of multi-species alignments

- Browse **more organisms** at [www.ensemblgenomes.org](http://www.ensemblgenomes.org)

- Search for a gene, location, variation, clone, probeset, or phenotype

- Current **location** is indicated

- Alternate **haplotypes** are available

- Browse genes & annotation for **over 50 species** in our chordate site

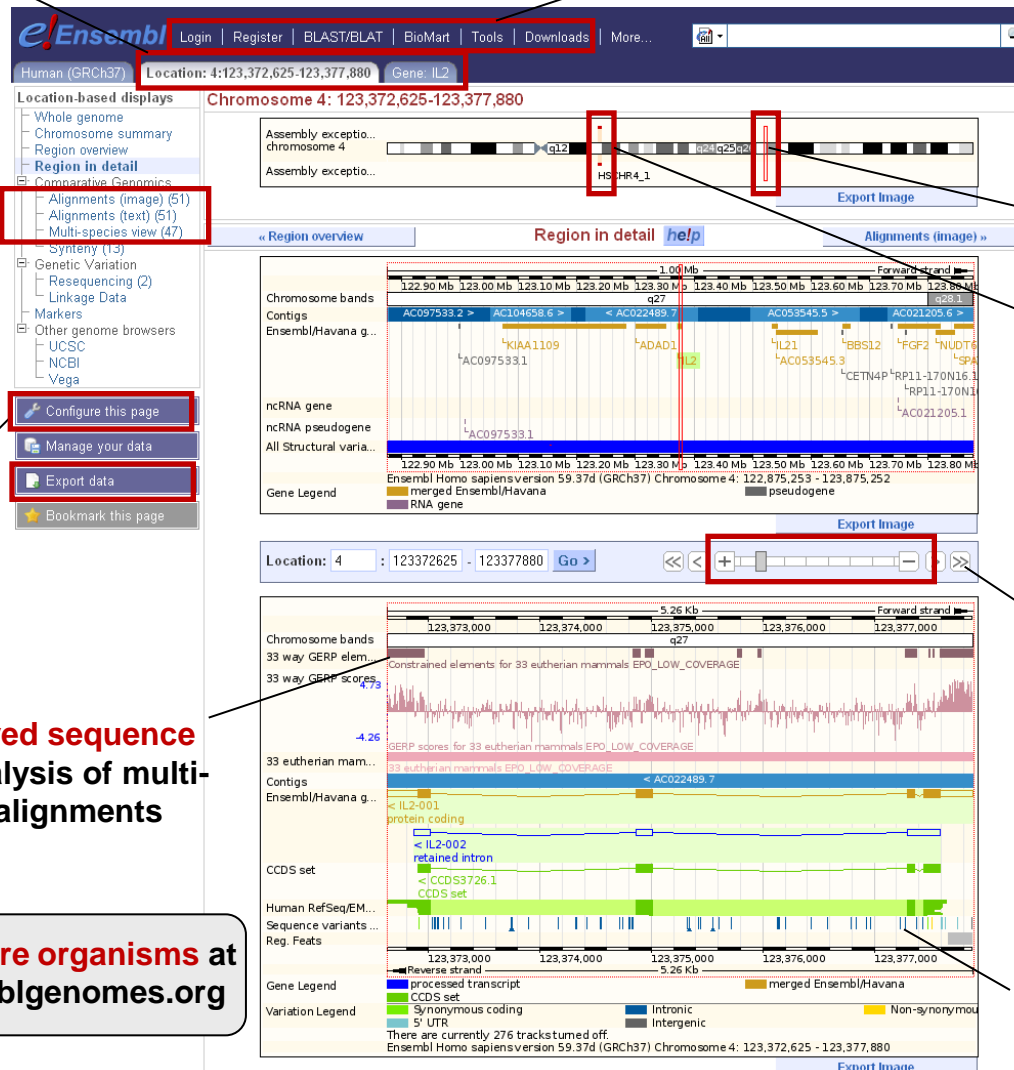
- Zoom into a region

Variation: rs3087209

**Variation Properties**

bp	123377483
Status	cluster, freq, hapmap
Class	snp
Ambiguity	M
code	
Mapweight	1
Alleles	A/C
Source	dbSNP, Illumina_Human1M-duoV3
Type	NON_SYNONYMOUS_CODING

- Click a colour-coded **sequence variant** for a pop-up box of information



Reference: Flicek, P. *et al.* Ensembl 2011 (2011) Nucleic Acids Research 39:D800-D806

Reference: BMC Bioinformatics – Ensembl Update 2010 <http://www.biomedcentral.com/series/ENSEMBL2010>

Tutorials: <http://www.ensembl.org/info/website/tutorials>

## The Gene Tab

Human (GRCh37) Location: 4,123,372,625-123,377,880 **Gene: IL2** Transcript: IL2-001 Variation: rs3087209

**Gene-based displays**

- Gene summary
- Splice variants (2)
- Supporting evidence
- Sequence
- External references (2)
- Regulation
- Comparative Genomics
  - Genomic alignments (51)
  - Gene Tree (image)
  - Gene Tree (text)
  - Gene Tree (alignment)
  - Orthologues (31)
  - Paralogues
  - Protein families (1)
- Genetic Variation
  - Variation Table
  - Variation Image
- External Data
  - Personal annotation
  - ID History
  - Gene history

**Gene: IL2 (ENSG00000109471)**

interleukin 2 [Source:HGNC Symbol;Acc:6001]

**Location** [Chromosome 4: 123,372,625-123,377,880](#) reverse strand.

**Transcripts** There are 2 transcripts in this gene

Name	Transcript ID	Length (bp)	Protein ID	Length (aa)	Biotype	CCDS
IL2-001	ENST00000226730	1029	ENSP00000226730	153	Protein coding	CCDS3726
IL2-002	ENST00000477645	586	No protein product	-	Retained intron	-

**Gene summary** [help](#)

**Name** IL2 (HGNC Symbol)

**Synonyms** IL-2 [To view all Ensembl genes linked to the name [click here](#).]

**CCDS** This gene is a member of the Human CCDS set: [CCDS3726](#)

**Gene type** Known protein coding

**Prediction Method** Gene containing both Ensembl genebuild transcripts and [Havana](#) manual curation, see [article](#).

**Alternative genes** This gene corresponds to the following database identifiers:  
Havana gene: [QTTHUMG00000133075](#) [view all locations](#)

Contigs  
Ensembl/Havana g...

25.26 Kb  
123.36 Mb 123.37 Mb 123.38 Mb 123.39 Mb  
Forward strand  
Reverse strand

AC022485.7  
IL2-001  
protein coding  
IL2-002  
retained intron

[Export Image](#)

- Tabs allow navigation through location, **gene**, transcript, variation, and gene regulation information

- Contextual navigation for a **gene**

- Comparative and **variation** displays are available

- **Export** sequence

- The **transcript table** lists splice variants and has **sortable columns**

- Page-specific **help** is available

- **Gold** transcripts are protein coding and are agreed on by Ensembl and Vega/Havana

- **Red** transcripts are protein coding and are from Ensembl or VEGA/Havana

- **Blue, Grey, or Pink** transcripts are non-coding

- Human protein coding transcripts make the **GENCODE** set

## The Transcript Tab: Exons View

« Supporting evidence » **Exons** [help](#) « cDNA sequence »

**Key**

**Colour key for variations**

- Variations
- 5' UTR
- Downstream
- Intronic
- Non-synonymous coding
- Regulatory region
- Synonymous coding
- Upstream

**Show/hide columns** Search:

No.	Exon / Intron	Start	End	Start Phase	End Phase	Length	Sequence
1	5' upstream sequence	123,377,449	123,377,880	-	0	432	.....Ggacaaagaaatttctgagttacttttgcacccagc cccttaagaagaaggaggaataactgtttcctacagaaggcgttaattgcatgaattagag CTATCACCTAAGTGTGGCTAATGTAACAAGAGGGATTTCACCTACATCATTCACTGCA GTCCTTGGGGGTTTAAAGAAATTCAGAGGCTCATCAGAGGAAATGAAGGTAATG ATTTTCAGACGGTAAGTCTTTGAAAAATGTGTAAATGTAACAACTTTTGACACCC CCATAATTTTCCAGAATTAAACAGTATAAATGTCATCTCTGTGTTCAAGAGTTCCTAT CACTCTTTTAATCACTACTCAGTAACCTCAACCTGCGCACAAATGACAGGATGCAA CTCCTGTCTTGCAATGCACTAAGTCTTGCACTTTGCAACAACAGTGCACCTACTCAAGT TTTCAAGAAACACAGCTACAACCTGGAGCATTTTCCTGGATTACAGATGATTTTG AATGGAAATTAAT
2	Intron 1-2	123,377,359	123,377,448	-	0	90	gtaagtatacttcttcttactaataattatcatttagtaattctagctggagatcatt tcttaataacaatgcaattatactttcttag
3	ENSE000000935280	123,377,299	123,377,358	0	0	60	AATTACAAGATCCCAAACTCACCGAGGATGCTCACATTAAGTTTACATGCCCAAGAAG

**5' UTR**

**Coding sequence**

**Upstream sequence**

**Intron**

- Protein signatures

- Download to open in **MS Word**

- Click a **variation** for population frequencies, phenotypes, and risk alleles

- Upstream sequence

- Intron