

- 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

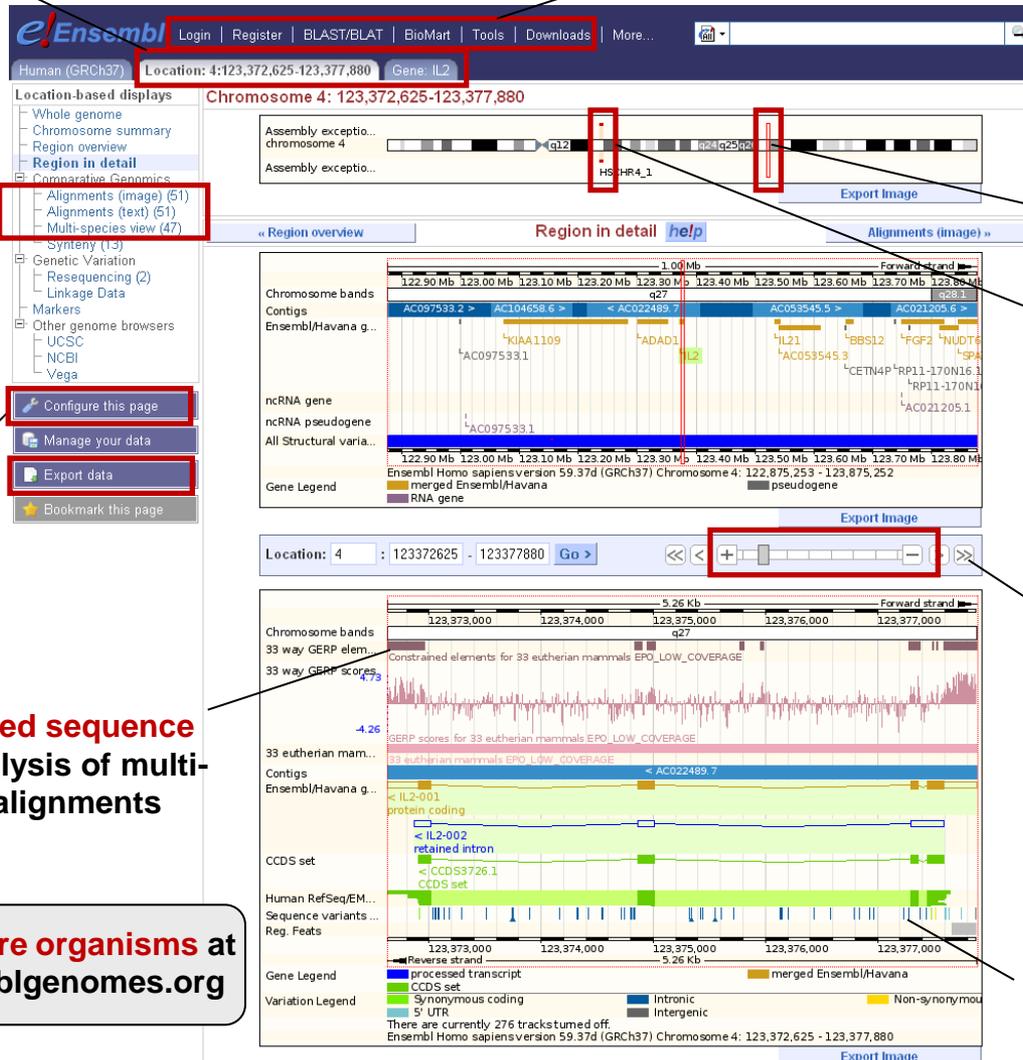
Mapweight 1

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>

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

## 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

Show/hide columns Search: \_\_\_\_\_

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 IL2 [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: [QTTTHUMG00000133075](#) [view all locations]

Contigs  
Ensembl/Havana g...  
AC022485.7  
IL2-001 protein coding  
IL2-002 retained intron

Export Image

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

- **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

- **Contextual navigation** for a **gene**
- **Comparative and variation** displays are available

- **Export** sequence

- **Page-specific help** is available

## The Transcript Tab: Exons View

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

Key

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

Show/hide columns Search: \_\_\_\_\_

No.	Exon / Intron	Start	End	Start Phase	End Phase	Length	Sequence
	5' upstream sequence						.....gacaaagaaaatttctgagttacttttgatcccaaccccttaagaagaaggagaaaactgtttcacaagaaggcttaattgcatgaattagag
1	<a href="#">ENSE000001298054</a>	123,377,449	123,377,880	-	0	432	CTATCACCTAAGTGTGGGCTAATGTAACAAGAGAGGATTCACCTACATCATTCAAGTCA GTCCTTTGGGGGTTTAAAGAAATTCGAAGAGCTATCAGAAGAGGAAAAATGAAGGTAATG TCTTTTCAGACGGTAAAGCTTTGAAAAATGTGTAAATATGTAACAACTTTGACACCC CCATAATATTTTCCAGAATTAACAGTATAAATGTCATCTCTGTTGTCAGAGTTCCCTAT CACTCTTTTAATCACTACTCACAGTAACTCAACTCTGGCCACAATGACAGGATGCAAA CTCCTGTCTTGGATTGCACTAAGTCTTGGACTTTGTCACAAACAGTGCACCTACTCA&GT TTTAAAGAAAACACAGCTACAACTGGAGACTTTCCTGGATTACAGATGATTTTG AATGGAAATTAAT
	Intron 1:2	123,377,359	123,377,448			90	gtaagtatattcttcttactaaaattattacatttagtaacttagctggagatcatt tcttaataacaatgcaattatactttcttag
2	<a href="#">ENSE000000935280</a>	123,377,299	123,377,380	0	0	60	AATTACAAGAATCCCAAACTCACAGGATGCTCACATTTAAGTTTACATGCCAAAGAAG

- **Protein signatures**
- **Download** to open in **MS Word**

- **Click a variation** for population frequencies, phenotypes, and risk alleles
- **UTR**
- **Coding sequence**
- **Intron**
- **Upstream sequence**

Ensembl  
Contact us at helpdesk@ensembl.org