

# *Hnrnpl* Cas9-CKO Strategy

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**Reviewer:**

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# Project Overview

**Project Name**

*Hnrnpl*

**Project type**

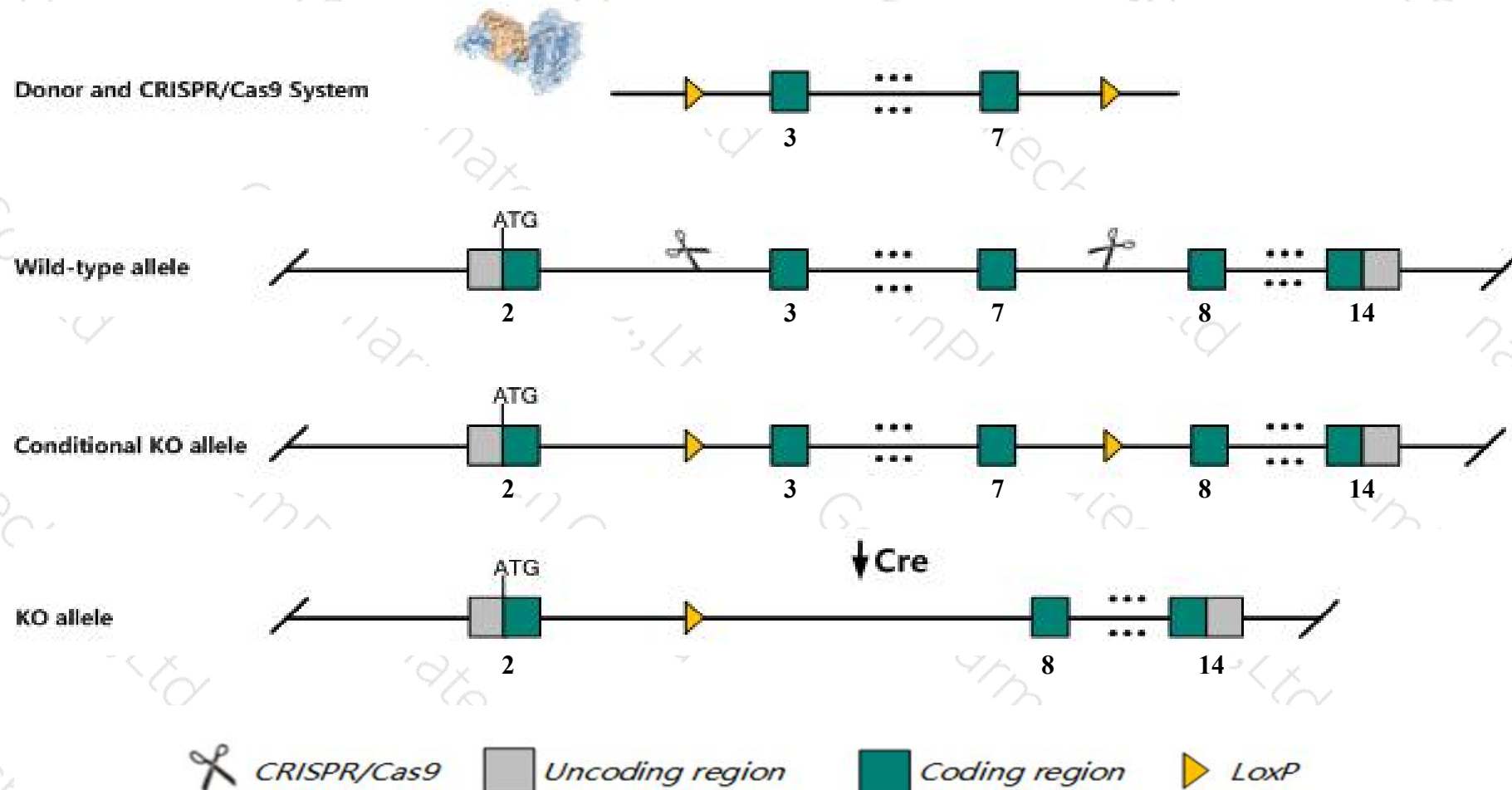
**Cas9-CKO**

**Strain background**

**C57BL/6JGpt**

# Conditional Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Hnrnp1* gene. The schematic diagram is as follows:



- The *Hnrnpl* gene has 13 transcripts. According to the structure of *Hnrnpl* gene, exon3-exon7 of *Hnrnpl*-211 (ENSMUST00000174548.7) transcript is recommended as the knockout region. The region contains 613bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Hnrnpl* gene. The brief process is as follows: CRISPR/Cas9 system and Donor were microinjected into the fertilized eggs of C57BL/6JGpt mice. Fertilized eggs were transplanted to obtain positive F0 mice which were confirmed by PCR and sequencing. A stable F1 generation mouse model was obtained by mating positive F0 generation mice with C57BL/6JGpt mice.
- The flox mice will be knocked out after mating with mice expressing Cre recombinase, resulting in the loss of function of the target gene in specific tissues and cell types.

- According to the existing MGI data, Mice homozygous for a targeted allele exhibit embryonic lethality after E3.5. Mice homozygous for a conditional allele activated in thymocytes exhibit decreased T cells in the periphery associated with impaired thymocyte chemotaxis.
- The flox region is about 3 kb away from the 5' end of the Gm44702 gene, which may affect the regulation of this gene.
- The *Hnrnp1* gene is located on the Chr7. If the knockout mice are crossed with other mice strains to obtain double gene positive homozygous mouse offspring, please avoid the two genes on the same chromosome.
- This Strategy is designed based on genetic information in existing databases. Due to the complexity of biological processes, all risk of loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at existing technological level.



# Gene information (NCBI)

## Hnrnp1 heterogeneous nuclear ribonucleoprotein L [Mus musculus (house mouse)]

Gene ID: 15388, updated on 31-Jan-2019

### Summary



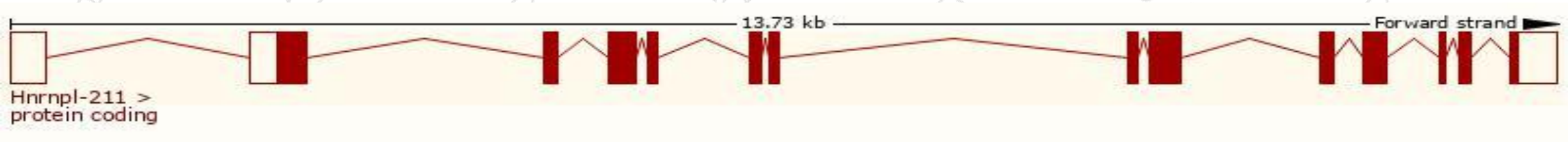
<b>Official Symbol</b>	Hnrnp1 provided by <a href="#">MGI</a>
<b>Official Full Name</b>	heterogeneous nuclear ribonucleoprotein L provided by <a href="#">MGI</a>
<b>Primary source</b>	<a href="#">MGI:MGI:104816</a>
<b>See related</b>	<a href="#">Ensembl:ENSMUSG00000015165</a>
<b>Gene type</b>	protein coding
<b>RefSeq status</b>	VALIDATED
<b>Organism</b>	<a href="#">Mus musculus</a>
<b>Lineage</b>	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
<b>Also known as</b>	C79783, D830027H13Rik, Hnrpl
<b>Expression</b>	Ubiquitous expression in CNS E11.5 (RPKM 136.9), limb E14.5 (RPKM 90.8) and 28 other tissues <a href="#">See more</a>
<b>Orthologs</b>	<a href="#">human</a> <a href="#">all</a>

# Transcript information (Ensembl)

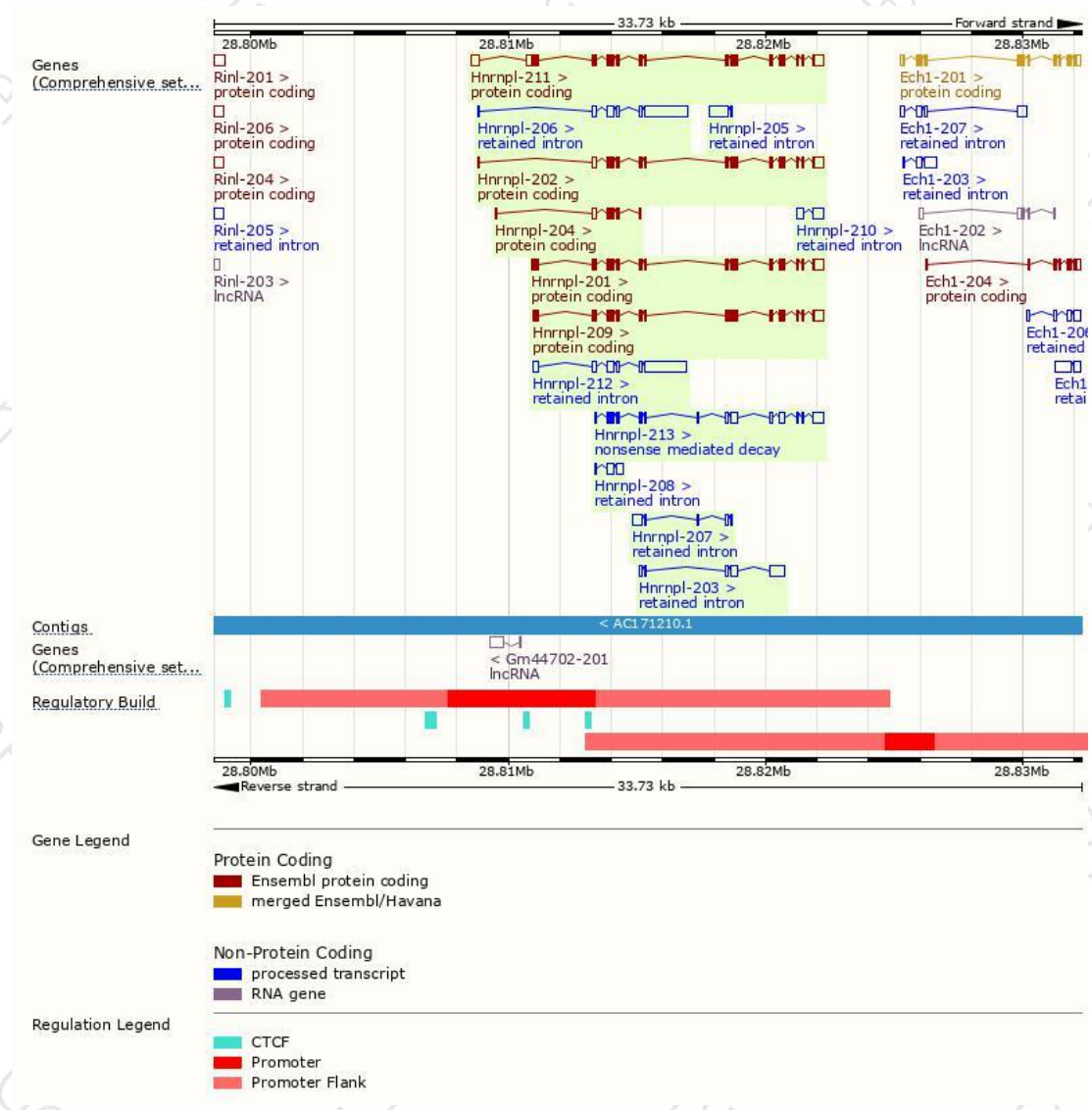
The gene has 13 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Hnrnpl-211	<a href="#">ENSMUST00000174548.7</a>	2679	<a href="#">586aa</a>	Protein coding	<a href="#">CCDS39864</a>	<a href="#">Q8R081</a>	TSL:5 GENCODE basic APPRIS P1
Hnrnpl-201	<a href="#">ENSMUST00000038572.14</a>	2142	<a href="#">586aa</a>	Protein coding	<a href="#">CCDS39864</a>	<a href="#">Q8R081</a>	TSL:1 GENCODE basic APPRIS P1
Hnrnpl-209	<a href="#">ENSMUST00000174477.7</a>	2180	<a href="#">615aa</a>	Protein coding	-	<a href="#">G5E924</a>	CDS 5' incomplete TSL:1
Hnrnpl-202	<a href="#">ENSMUST00000172529.7</a>	1898	<a href="#">456aa</a>	Protein coding	-	<a href="#">G3UY38</a>	TSL:5 GENCODE basic
Hnrnpl-204	<a href="#">ENSMUST00000172884.7</a>	502	<a href="#">112aa</a>	Protein coding	-	<a href="#">G3UYY3</a>	CDS 3' incomplete TSL:5
Hnrnpl-213	<a href="#">ENSMUST00000174882.7</a>	1850	<a href="#">201aa</a>	Nonsense mediated decay	-	<a href="#">G3UY56</a>	CDS 5' incomplete TSL:5
Hnrnpl-212	<a href="#">ENSMUST00000174755.7</a>	2426	No protein	Retained intron	-	-	TSL:1
Hnrnpl-206	<a href="#">ENSMUST00000173750.7</a>	2275	No protein	Retained intron	-	-	TSL:1
Hnrnpl-203	<a href="#">ENSMUST00000172841.1</a>	1087	No protein	Retained intron	-	-	TSL:3
Hnrnpl-205	<a href="#">ENSMUST00000173578.1</a>	768	No protein	Retained intron	-	-	TSL:2
Hnrnpl-210	<a href="#">ENSMUST00000174526.1</a>	672	No protein	Retained intron	-	-	TSL:2
Hnrnpl-207	<a href="#">ENSMUST00000173818.7</a>	644	No protein	Retained intron	-	-	TSL:3
Hnrnpl-208	<a href="#">ENSMUST00000174396.1</a>	504	No protein	Retained intron	-	-	TSL:2

The strategy is based on the design of *Hnrnpl-211* transcript,The transcription is shown below

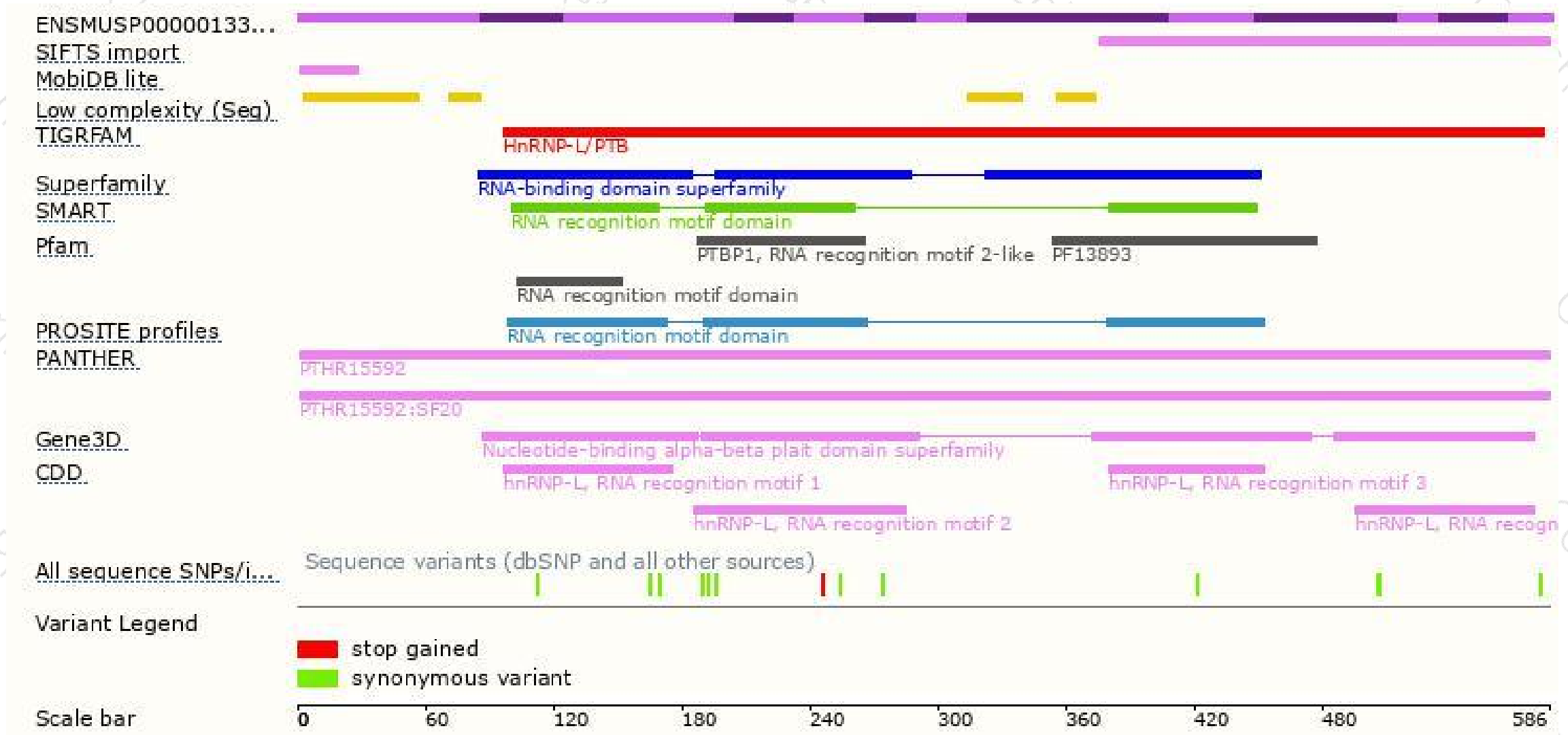


# Genomic location distribution

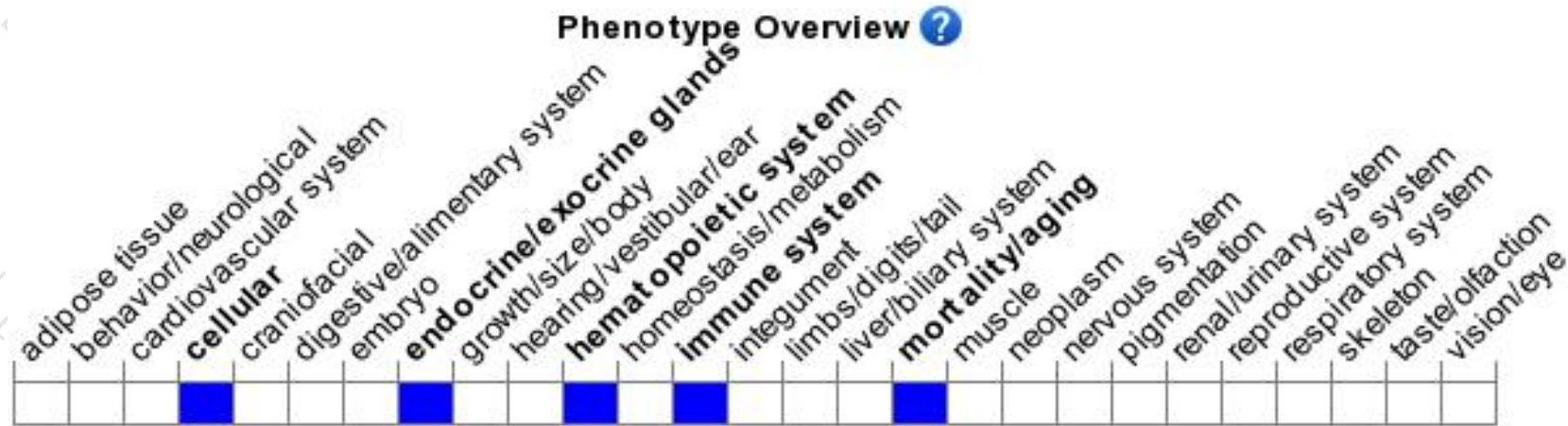




# Protein domain



# Mouse phenotype description(MGI)



*Phenotypes affected by the gene are marked in blue. Data quoted from MGI database(<http://www.informatics.jax.org/>).*

According to the existing MGI data, Mice homozygous for a targeted allele exhibit embryonic lethality after E3.5. Mice homozygous for a conditional allele activated in thymocytes exhibit decreased T cells in the periphery associated with impaired thymocyte chemotaxis.

If you have any questions, you are welcome to inquire.

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