

# ***Cd86 Cas9-KO Strategy***

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# Project Overview

**Project Name**

***Cd86***

**Project type**

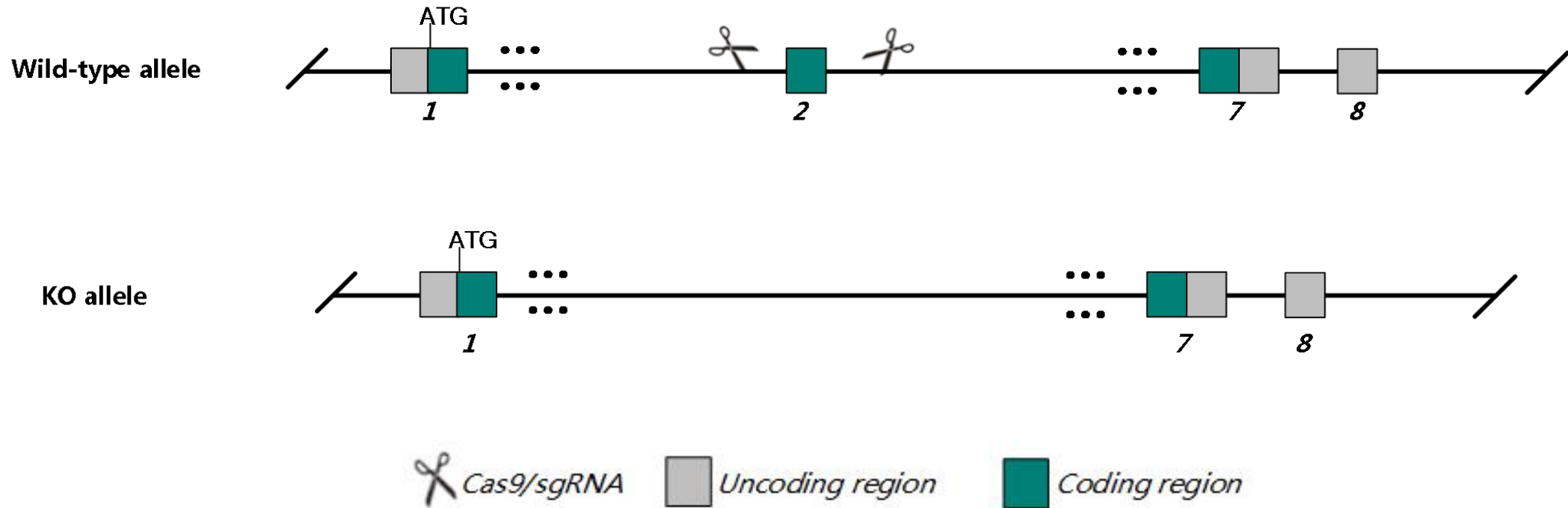
**Cas9-KO**

**Strain background**

**C57BL/6JGpt**

# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Cd86* gene. The schematic diagram is as follows:



- The *Cd86* gene has 4 transcripts. According to the structure of *Cd86* gene, exon2 of *Cd86-201* (ENSMUST00000089620.10) transcript is recommended as the knockout region. The region contains 50bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Cd86* gene. The brief process is as follows: CRISPR/Cas9 system

- According to the existing MGI data, Homozygous null mice on an NOD background display a phenotype similar to human Guillain-Barre Syndrome, exhibiting severe peripheral nervous system inflammation, sciatic nerve demyelination, elevated auto-antibodies to myelin protein zero, hindlimb paralysis, and weak forelimb grip.
- The *Cd86* gene is located on the Chr16. 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 the gene knockout on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.



# Gene information (NCBI)

## Cd86 CD86 antigen [Mus musculus (house mouse)]

Gene ID: 12524, updated on 12-Feb-2019

### Summary



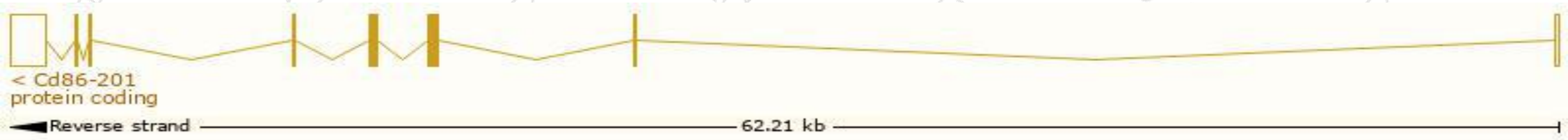
<b>Official Symbol</b>	Cd86 provided by <a href="#">MGI</a>
<b>Official Full Name</b>	CD86 antigen provided by <a href="#">MGI</a>
<b>Primary source</b>	<a href="#">MGI:MGI:101773</a>
<b>See related</b>	<a href="#">Ensembl:ENSMUSG00000022901</a>
<b>Gene type</b>	protein coding
<b>RefSeq status</b>	VALIDATED
<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>	B7, B7-2, B7.2, B70, CLS1, Cd28l2, ETC-1, Ly-58, Ly58, MB7, MB7-2, TS/A-2
<b>Expression</b>	Broad expression in spleen adult (RPKM 3.3), liver E18 (RPKM 2.8) and 21 other tissues <a href="#">See more</a>
<b>Orthologs</b>	<a href="#">human</a> <a href="#">all</a>

# Transcript information (Ensembl)

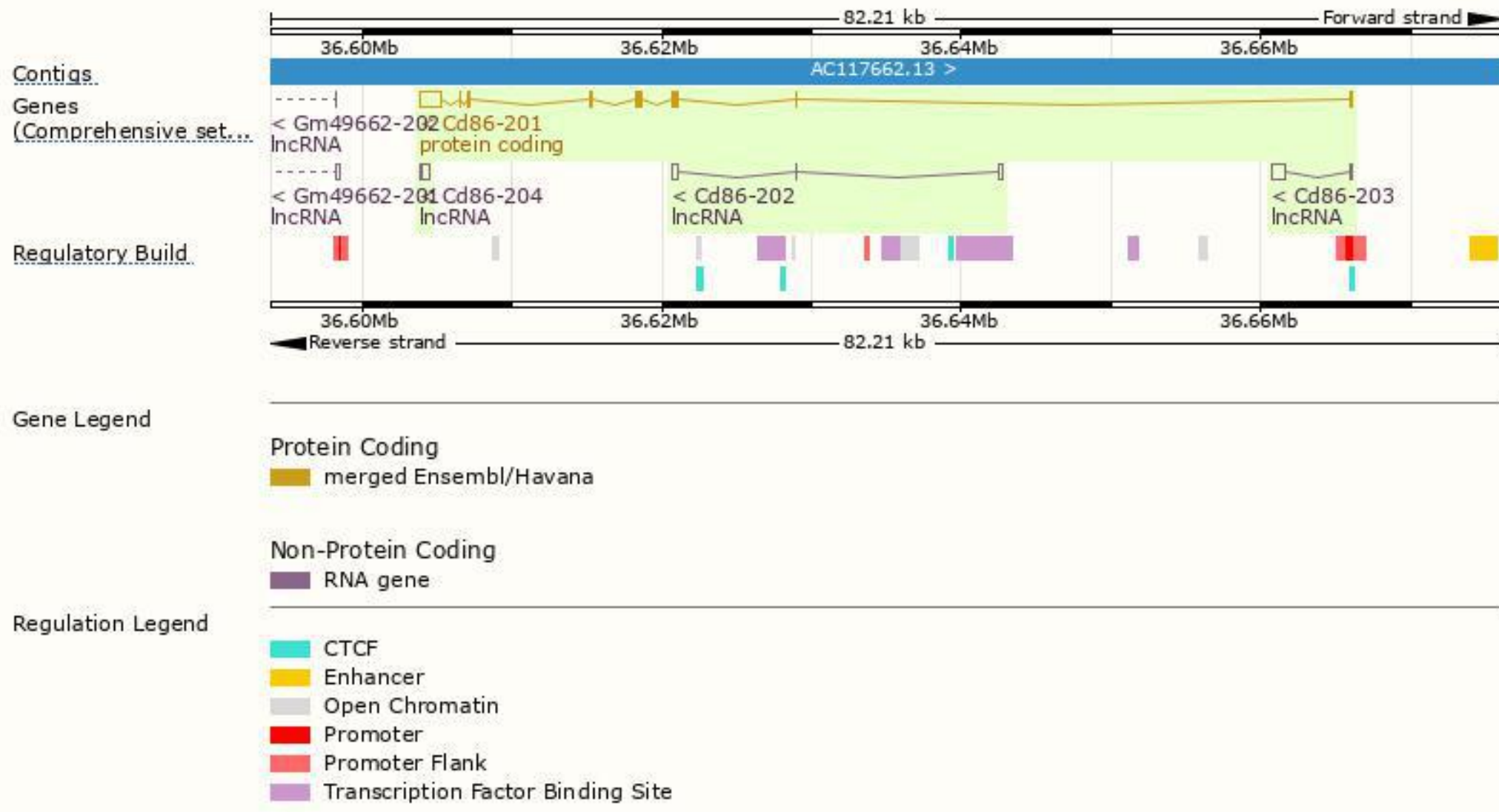
The gene has 4 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Cd86-201	<a href="#">ENSMUST00000089620.10</a>	2539	<a href="#">309aa</a>	Protein coding	<a href="#">CCDS28155</a>	<a href="#">P42082 Q549Q9</a>	TSL:1 GENCODE basic APPRIS P1
Cd86-203	<a href="#">ENSMUST00000145506.1</a>	975	No protein	lncRNA	-	-	TSL:1
Cd86-202	<a href="#">ENSMUST00000135280.1</a>	654	No protein	lncRNA	-	-	TSL:3
Cd86-204	<a href="#">ENSMUST00000154485.1</a>	548	No protein	lncRNA	-	-	TSL:3

The strategy is based on the design of *Cd86-201* transcript,The transcription is shown below

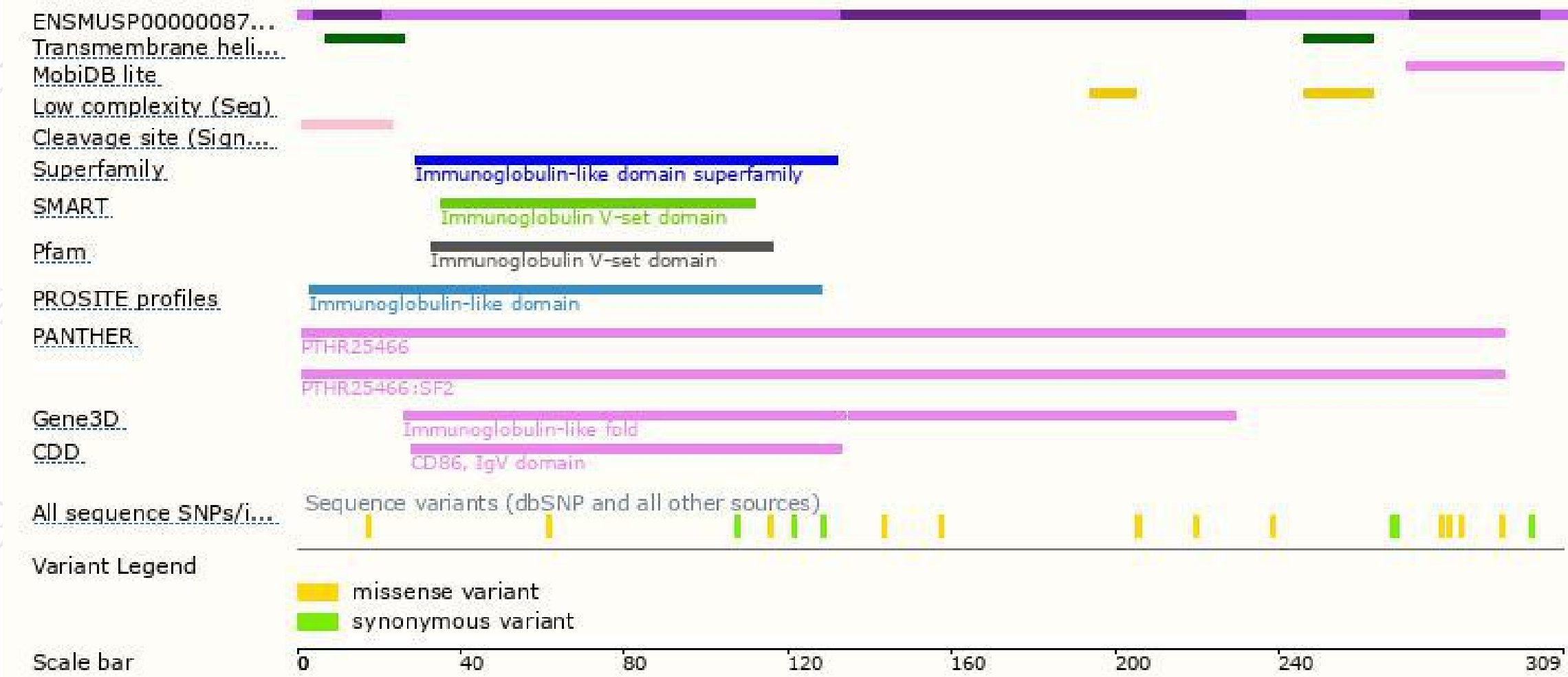


# Genomic location distribution

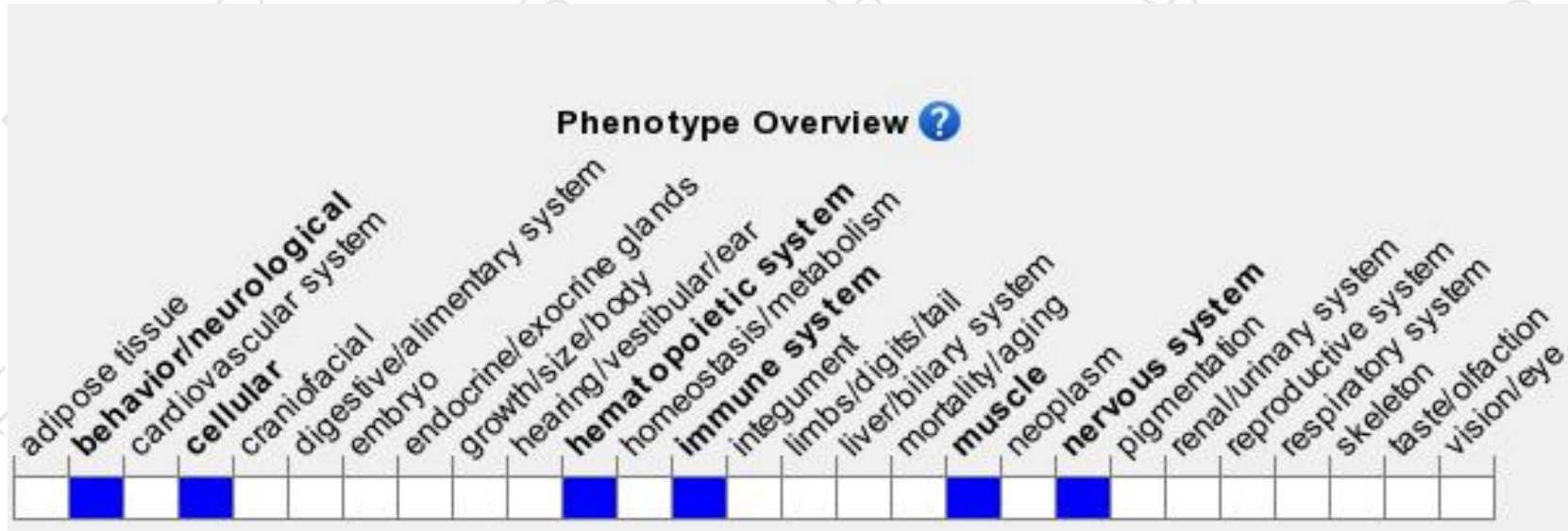




# Protein domain



# 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, Homozygous null mice on an NOD background display a phenotype similar to human Guillain-Barre Syndrome, exhibiting severe peripheral nervous system inflammation, sciatic nerve demyelination, elevated auto-antibodies to myelin protein zero, hindlimb paralysis, and weak forelimb grip.

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

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