

# *Nedd9* Cas9-KO Strategy

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**Reviewer:**

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**Design Date:**

**2020-2-10**

# Project Overview

**Project Name**

*Nedd9*

**Project type**

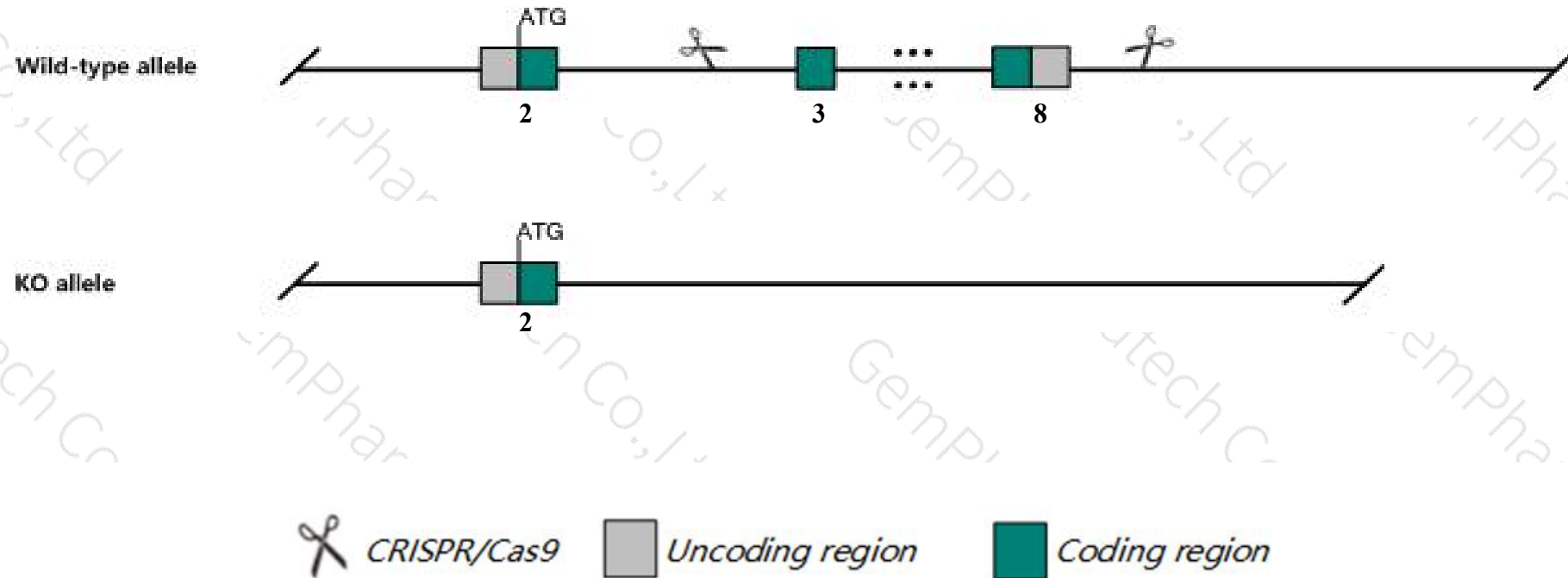
**Cas9-KO**

**Strain background**

**C57BL/6JGpt**

# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Nedd9* gene. The schematic diagram is as follows:



- The *Nedd9* gene has 6 transcripts. According to the structure of *Nedd9* gene, exon3-exon8 of *Nedd9-201* (ENSMUST00000021794.13) transcript is recommended as the knockout region. The region contains most of the coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Nedd9* gene. The brief process is as follows: CRISPR/Cas9 system

- According to the existing MGI data, Mice homozygous for one null allele exhibit impaired lymphocyte trafficking and a deficit of splenic marginal zone B cells. Mice homozygous for another null allele display impaired spatial learning and decreased hippocampal dendritic spine densities.
- The *Nedd9* gene is located on the Chr13. 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.



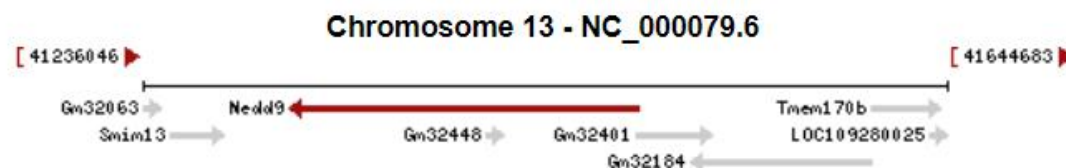
# Gene information (NCBI)

**Nedd9** neural precursor cell expressed, developmentally down-regulated gene 9 [ *Mus musculus* (house mouse) ]

Gene ID: 18003, updated on 12-Aug-2019

## Summary

<b>Official Symbol</b>	Nedd9 provided by <a href="#">MGI</a>
<b>Official Full Name</b>	neural precursor cell expressed, developmentally down-regulated gene 9 provided by <a href="#">MGI</a>
<b>Primary source</b>	<a href="#">MGI:MGI:97302</a>
<b>See related</b>	<a href="#">Ensembl:ENSMUSG00000021365</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>	CasL; HEF1; MEF1; p105; Cas-L; Nedd-9
<b>Expression</b>	Ubiquitous expression in lung adult (RPKM 23.8), spleen adult (RPKM 20.3) and 22 other tissues <a href="#">See more</a>
<b>Orthologs</b>	<a href="#">human</a> <a href="#">all</a>



# Transcript information (Ensembl)

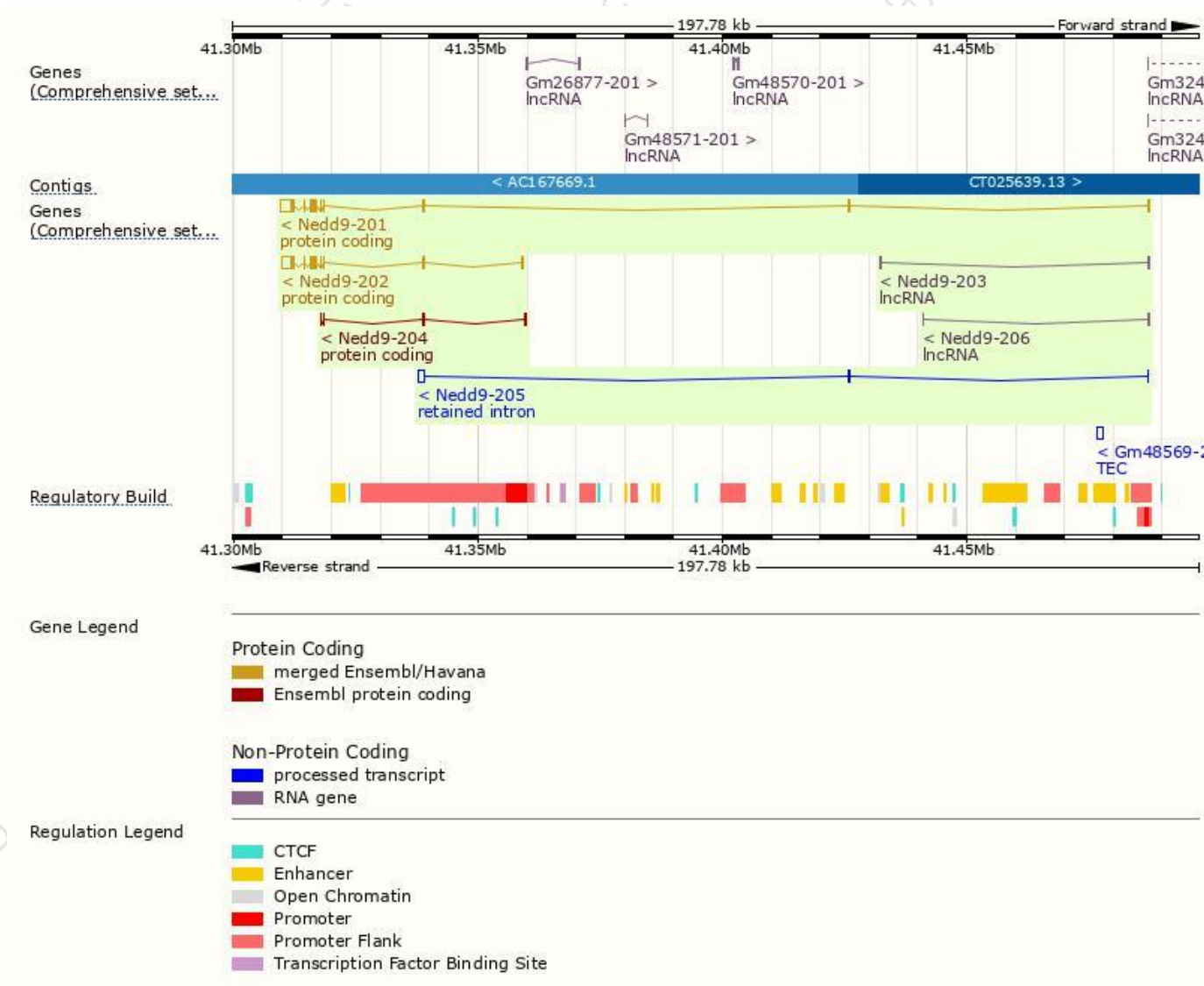
The gene has 6 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Translation ID	Biotype	CCDS	UniProt	Flags
Nedd9-201	<a href="#">ENSMUST00000021794.13</a>	4871	<a href="#">833aa</a>	<a href="#">ENSMUSP00000021794.6</a>	Protein coding	<a href="#">CCDS49247</a>	<a href="#">Q35177</a>	TSL:1 GENCODE basic APPRIS ALT1
Nedd9-202	<a href="#">ENSMUST000000163623.2</a>	4626	<a href="#">833aa</a>	<a href="#">ENSMUSP000000125773.1</a>	Protein coding	<a href="#">CCDS36640</a>	<a href="#">A0A0R4J212</a>	TSL:1 GENCODE basic APPRIS P3
Nedd9-204	<a href="#">ENSMUST000000224803.1</a>	711	<a href="#">194aa</a>	<a href="#">ENSMUSP000000152937.1</a>	Protein coding	-	<a href="#">A0A286YCE4</a>	CDS 3' incomplete
Nedd9-205	<a href="#">ENSMUST000000225053.1</a>	1455	No protein	-	Retained intron	-	-	-
Nedd9-206	<a href="#">ENSMUST000000225297.1</a>	302	No protein	-	lncRNA	-	-	-
Nedd9-203	<a href="#">ENSMUST000000224083.1</a>	267	No protein	-	lncRNA	-	-	-

The strategy is based on the design of *Nedd9-201* transcript,The transcription is shown below

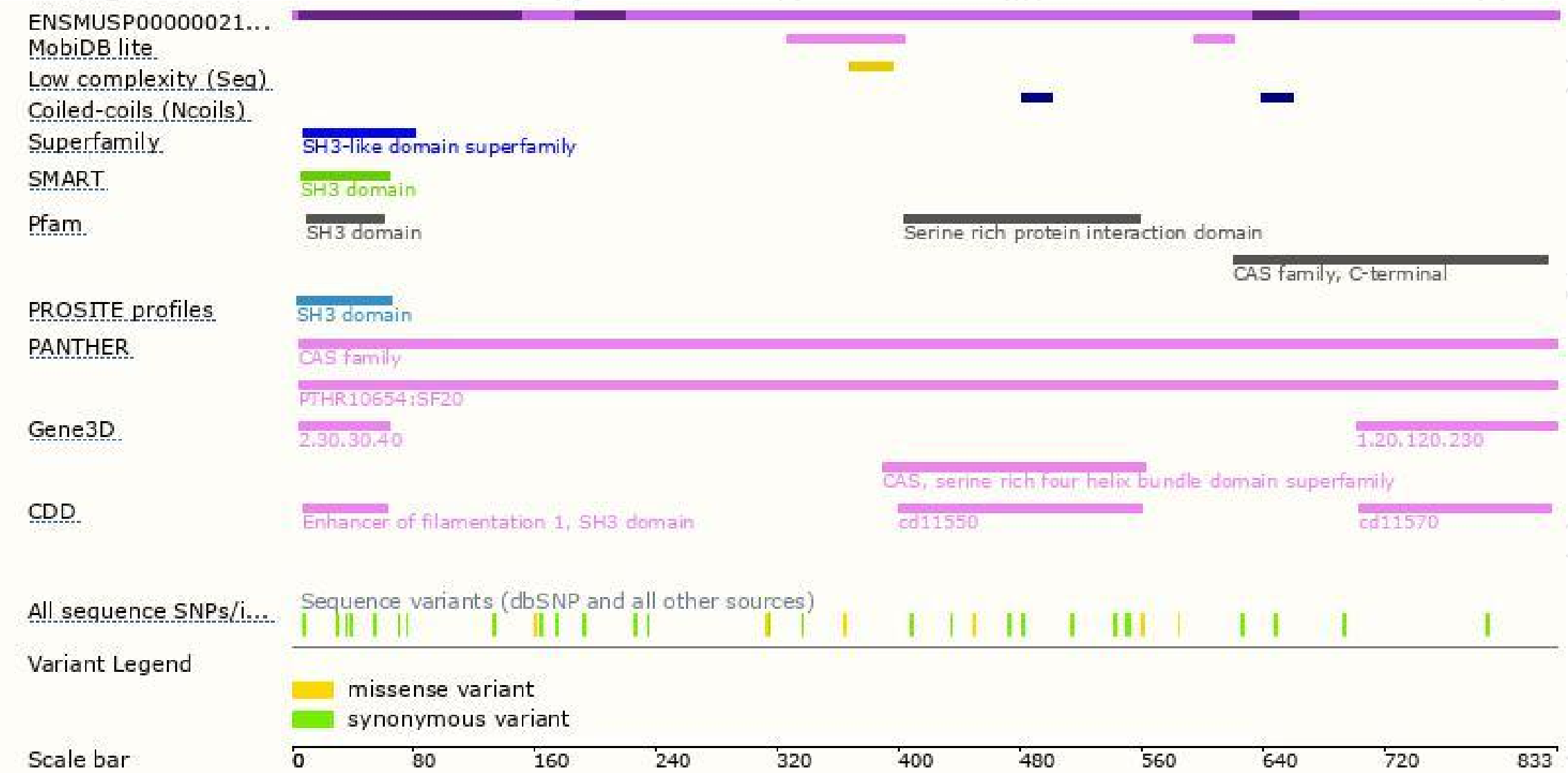


# Genomic location distribution



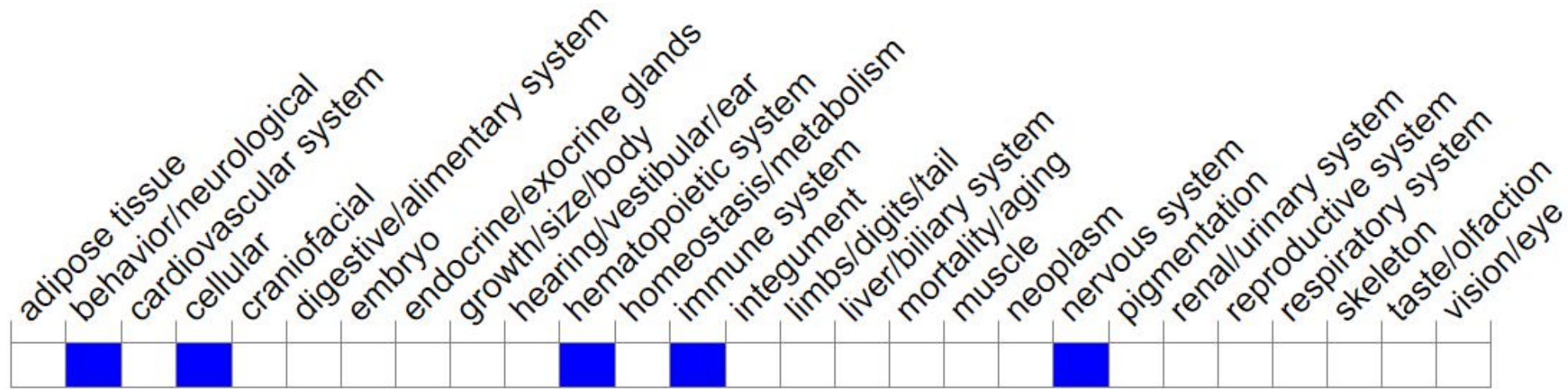


# Protein domain



# Mouse phenotype description(MGI)

## Phenotype Overview ?



*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 one null allele exhibit impaired lymphocyte trafficking and a deficit of splenic marginal zone B cells. Mice homozygous for another null allele display impaired spatial learning and decreased hippocampal dendritic spine densities.

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

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