

# *Cwc27* Cas9-KO Strategy

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# Project Overview

**Project Name**

***Cwc27***

**Project type**

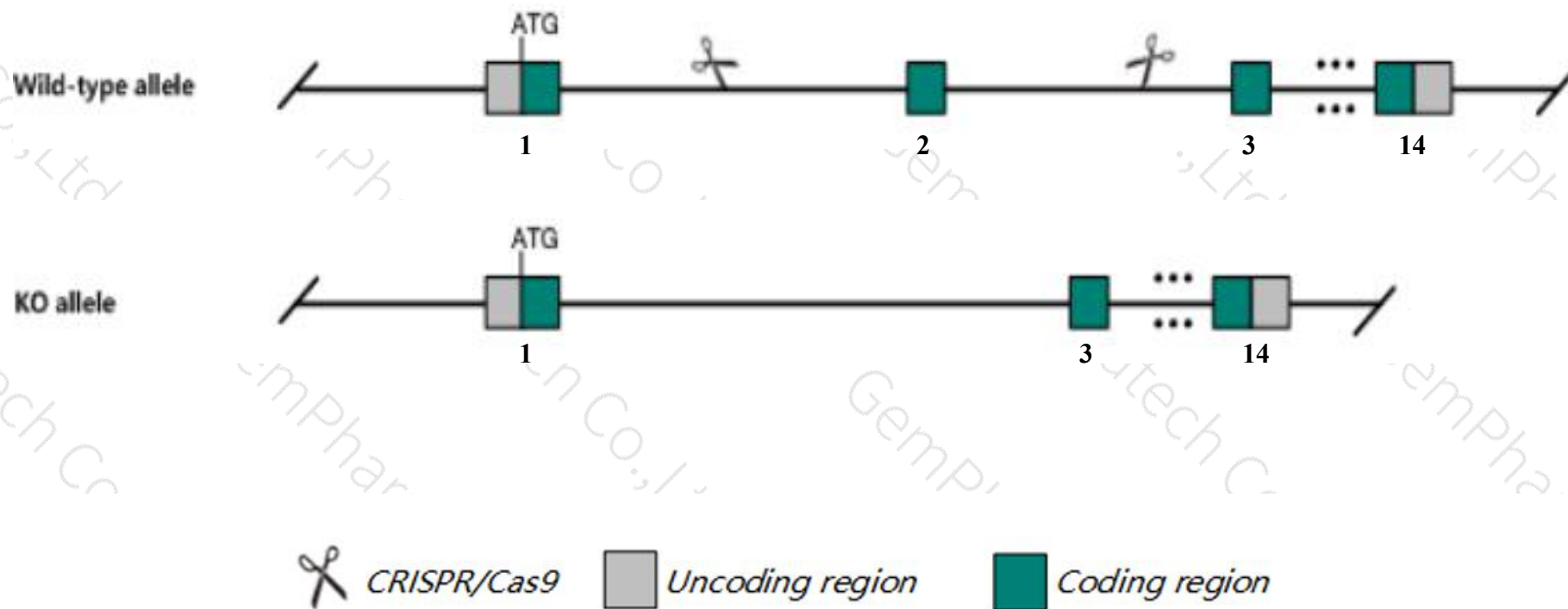
**Cas9-KO**

**Strain background**

**C57BL/6JGpt**

# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Cwc27* gene. The schematic diagram is as follows:



- The *Cwc27* gene has 4 transcripts. According to the structure of *Cwc27* gene, exon2 of *Cwc27-201* (ENSMUST00000022228.12) transcript is recommended as the knockout region. The region contains 97bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Cwc27* gene. The brief process is as follows: CRISPR/Cas9 system

- According to the existing MGI data, homozygous mutant mice exhibit reduced viability. Survivors after birth show signs of growth retardation and retinal depigmentation, along with numerous neurological, immunological, and blood chemistry abnormalities.
- The partial intron of transcript *Sreklip1*-203 will be deleted together in this strategy.
- The knockout region is near to the N-terminal of *Sreklip1* gene, this strategy may influence the regulatory function of the N-terminal of *Sreklip1* gene.
- Transcript *Cwc27*-202&203 may not be affected.
- The *Cwc27* 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)

## Cwc27 CWC27 spliceosome-associated protein [Mus musculus (house mouse)]

Gene ID: 67285, updated on 13-Mar-2020

### Summary



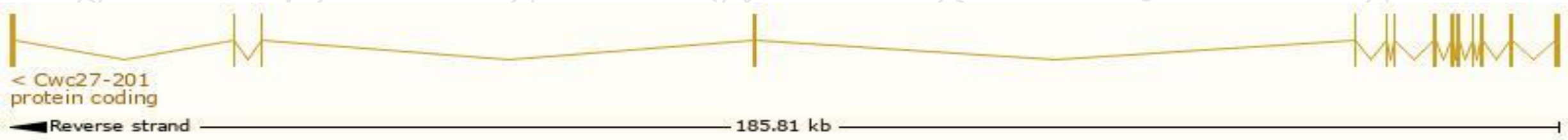
<b>Official Symbol</b>	Cwc27 provided by <a href="#">MGI</a>
<b>Official Full Name</b>	CWC27 spliceosome-associated protein provided by <a href="#">MGI</a>
<b>Primary source</b>	<a href="#">MGI:MGI:1914535</a>
<b>See related</b>	<a href="#">Ensembl:ENSMUSG000000021715</a>
<b>Gene type</b>	protein coding
<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>	3110009E13Rik, NY-CO-10, Sdccag10
<b>Expression</b>	Broad expression in CNS E11.5 (RPKM 1.3), CNS E14 (RPKM 0.7) and 21 other tissues <a href="#">See more</a>
<b>Orthologs</b>	<a href="#">human</a> <a href="#">all</a>

# Transcript information (Ensembl)

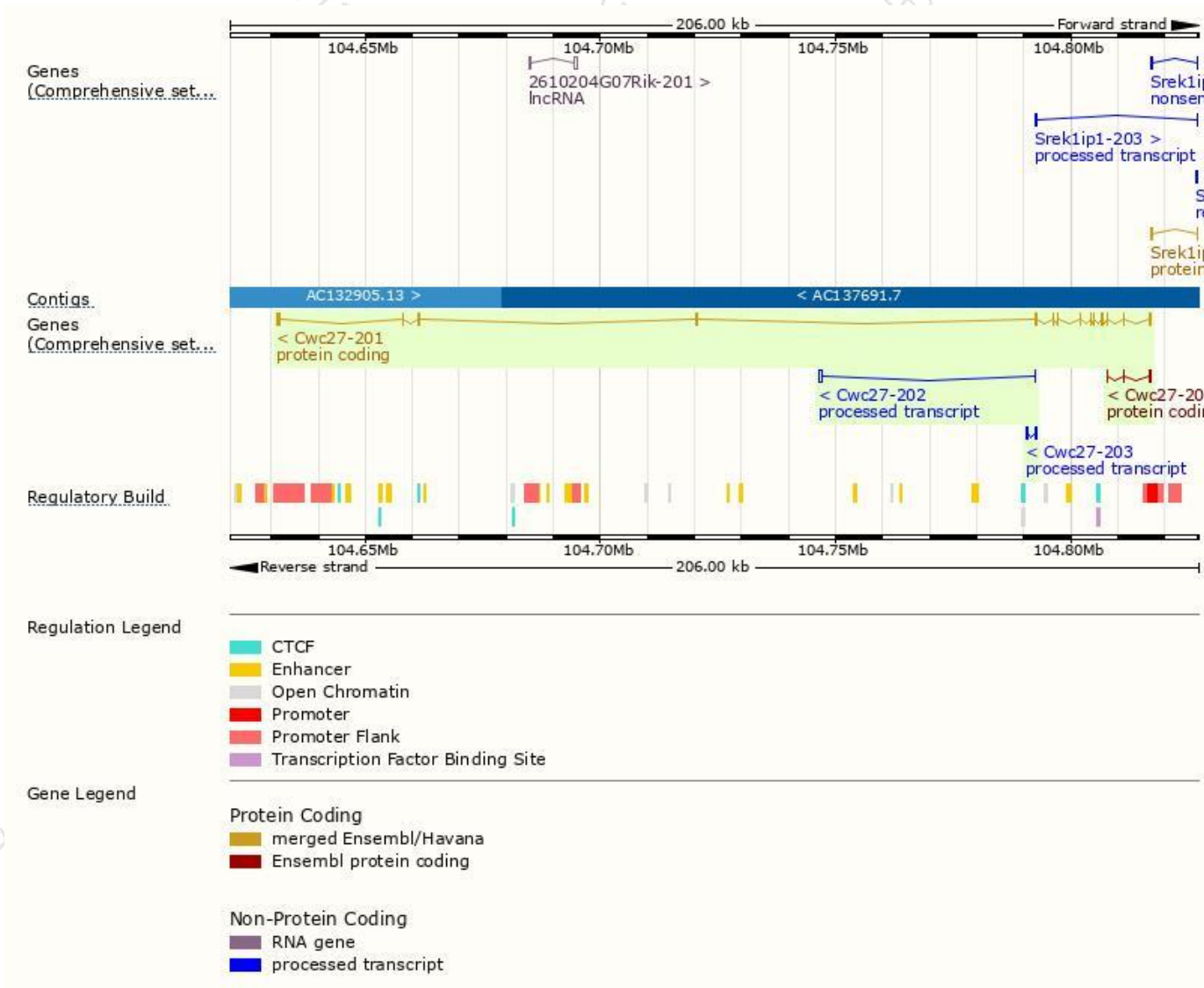
The gene has 4 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Cwc27-201	<a href="#">ENSMUST00000022228.12</a>	2071	<a href="#">469aa</a>	Protein coding	<a href="#">CCDS26752</a>	<a href="#">Q3TKY6</a>	TSL:1 GENCODE basic APPRIS is a system to annotate alternatively spliced transcripts based on a range of computational methods to identify the most functionally important transcript(s) of a gene. APPRIS P1
Cwc27-204	<a href="#">ENSMUST00000154165.1</a>	353	<a href="#">83aa</a>	Protein coding	-	<a href="#">D3Z2U3</a>	CDS 3' incomplete TSL:3
Cwc27-202	<a href="#">ENSMUST00000141495.1</a>	588	No protein	Processed transcript	-	-	TSL:3
Cwc27-203	<a href="#">ENSMUST00000147514.1</a>	451	No protein	Processed transcript	-	-	TSL:3

The strategy is based on the design of *Cwc27-201* transcript,the transcription is shown below

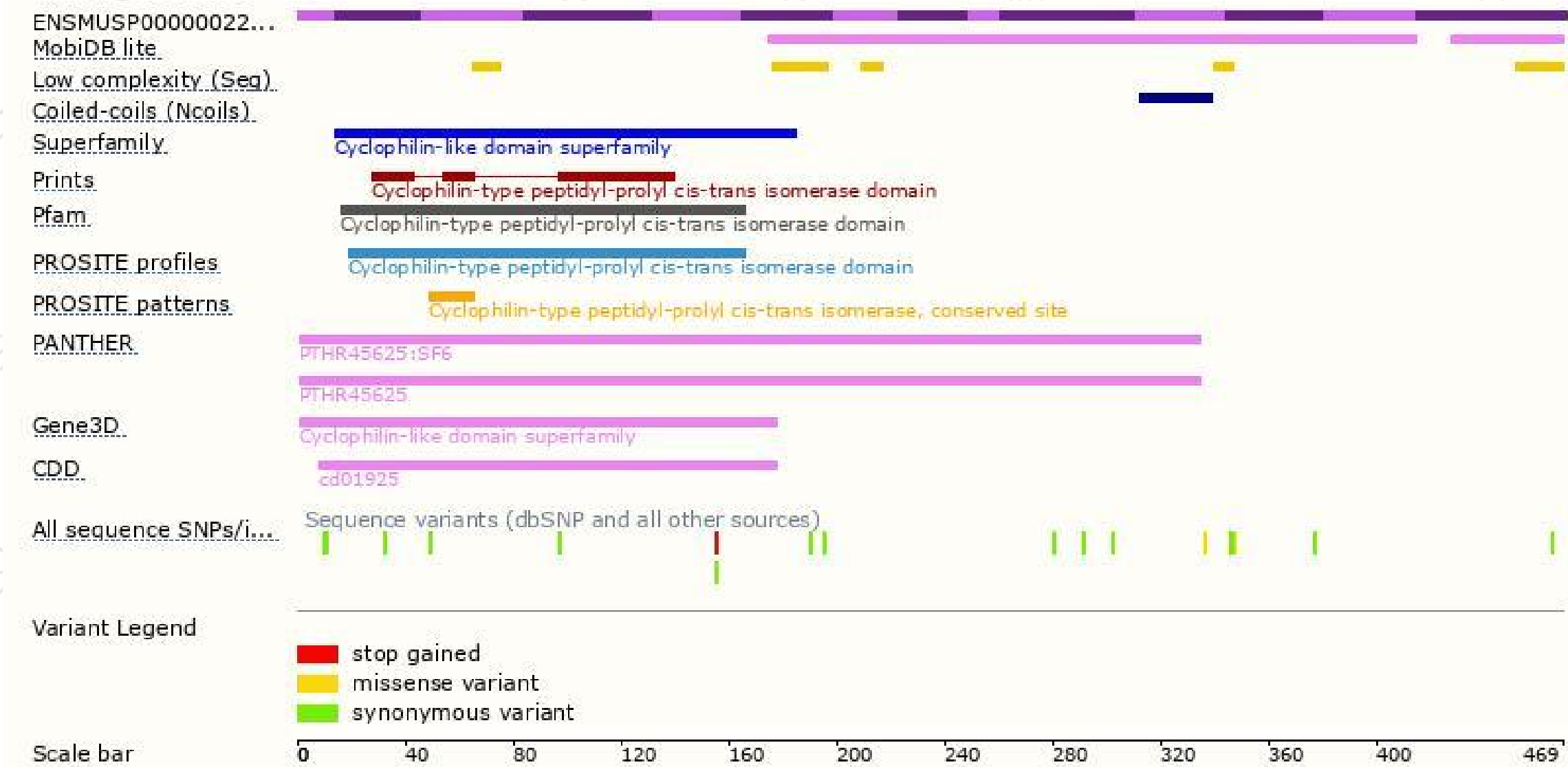


# Genomic location distribution

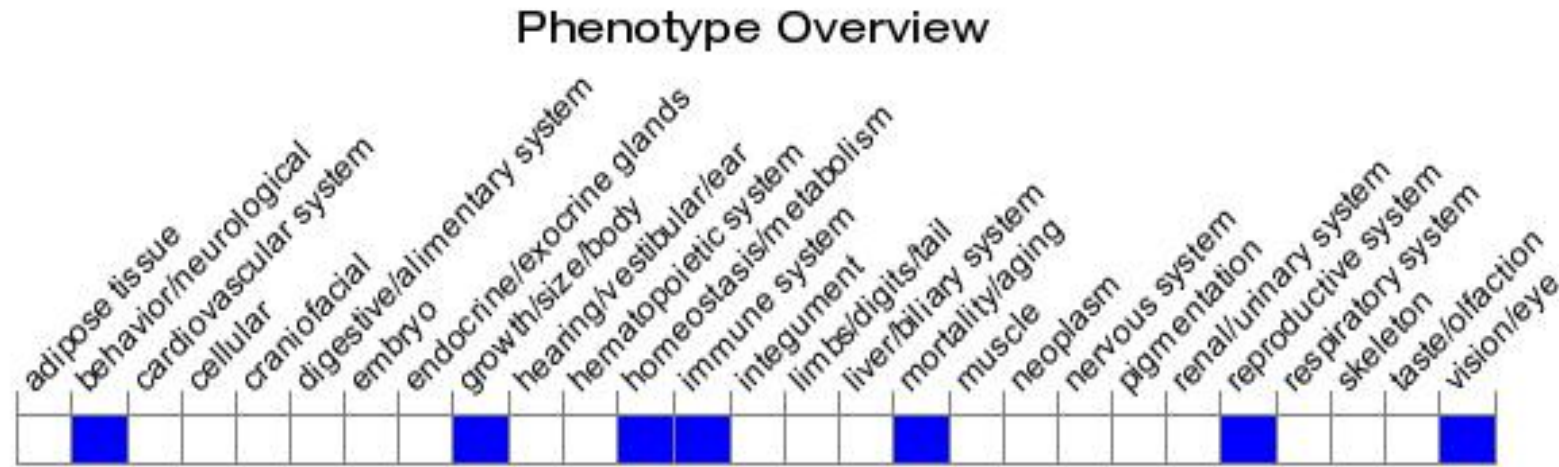




# Protein domain



# 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 mutant mice exhibit reduced viability. Survivors after birth show signs of growth retardation and retinal depigmentation, along with numerous neurological, immunological, and blood chemistry abnormalities.

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

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