

# *Rad23b* Cas9-KO Strategy

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**Reviewer:**

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**2019-9-20**

# Project Overview

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**Project Name**

*Rad23b*

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**Project type**

**Cas9-KO**

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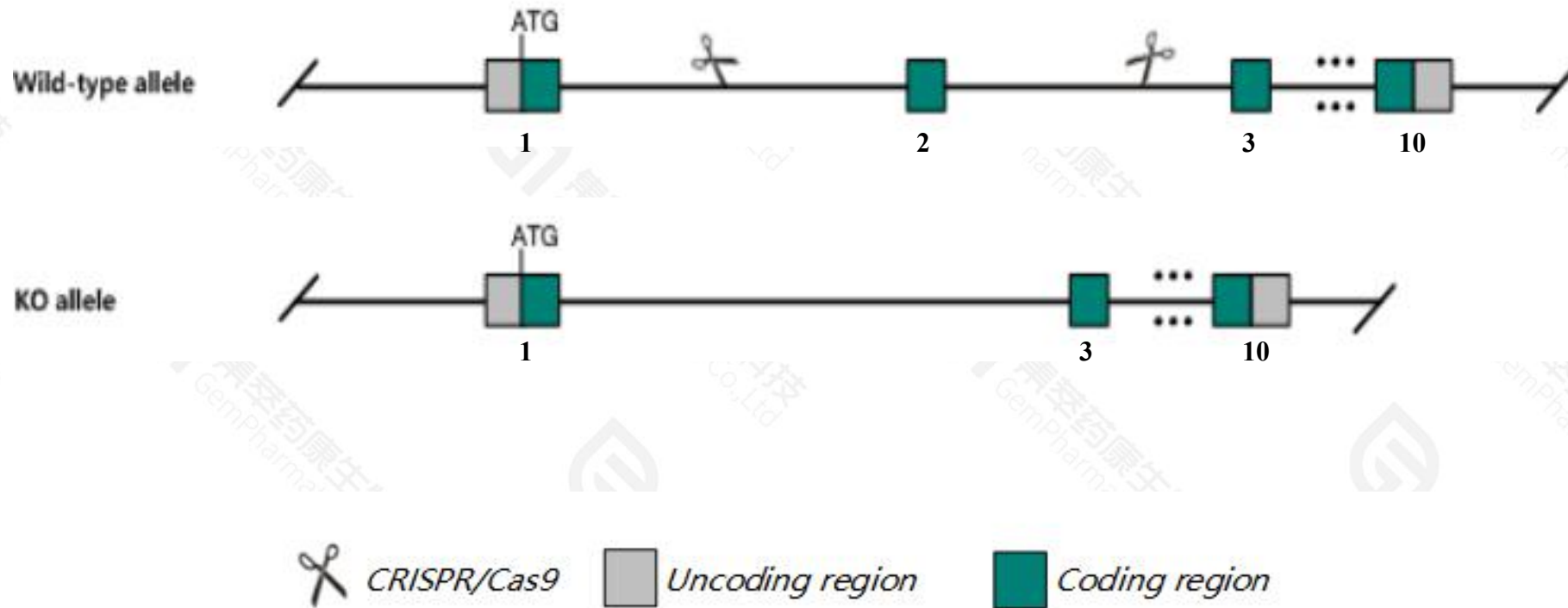
**Strain background**

**C57BL/6JGpt**

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# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Rad23b* gene. The schematic diagram is as follows:



# Technical routes

- The *Rad23b* gene has 4 transcripts. According to the structure of *Rad23b* gene, exon2 of *Rad23b-201*(ENSMUST00000030134.9) transcript is recommended as the knockout region. The region contains 82bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Rad23b* gene. The brief process is as follows: CRISPR/Cas9 system 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.

- According to the existing MGI data, mice homozygous for a disruption in this gene usually die around the time of birth. Those that survive show growth retardation, eye, reproductive, behavioral, and digestive system abnormalities. They usually die within 1 year of birth.
- The *Rad23b* gene is located on the Chr4. 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.



## Rad23b RAD23 homolog B, nucleotide excision repair protein [Mus musculus (house mouse)]

Gene ID: 19359, updated on 13-Mar-2020

### Summary

**Official Symbol** Rad23b provided by [MGI](#)

**Official Full Name** RAD23 homolog B, nucleotide excision repair protein provided by [MGI](#)

**Primary source** [MGI:MGI:105128](#)

**See related** [Ensembl:ENSMUSG00000028426](#)

**Gene type** protein coding

**RefSeq status** VALIDATED

**Organism** [Mus musculus](#)

**Lineage** Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus

**Also known as** 0610007D13Rik, AV001138, HR23B, mHR23B, p58

**Expression** Ubiquitous expression in testis adult (RPKM 58.0), CNS E11.5 (RPKM 42.9) and 28 other tissues [See more](#)

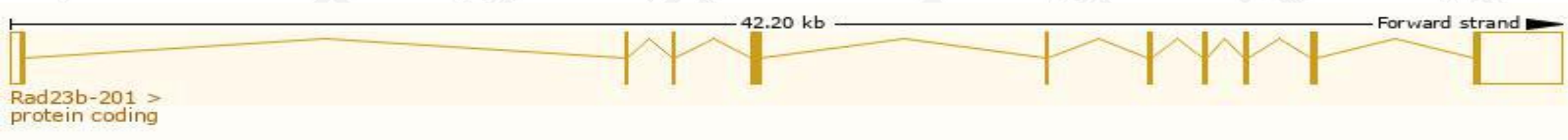
**Orthologs** [human](#) [all](#)

# Transcript information (Ensembl)

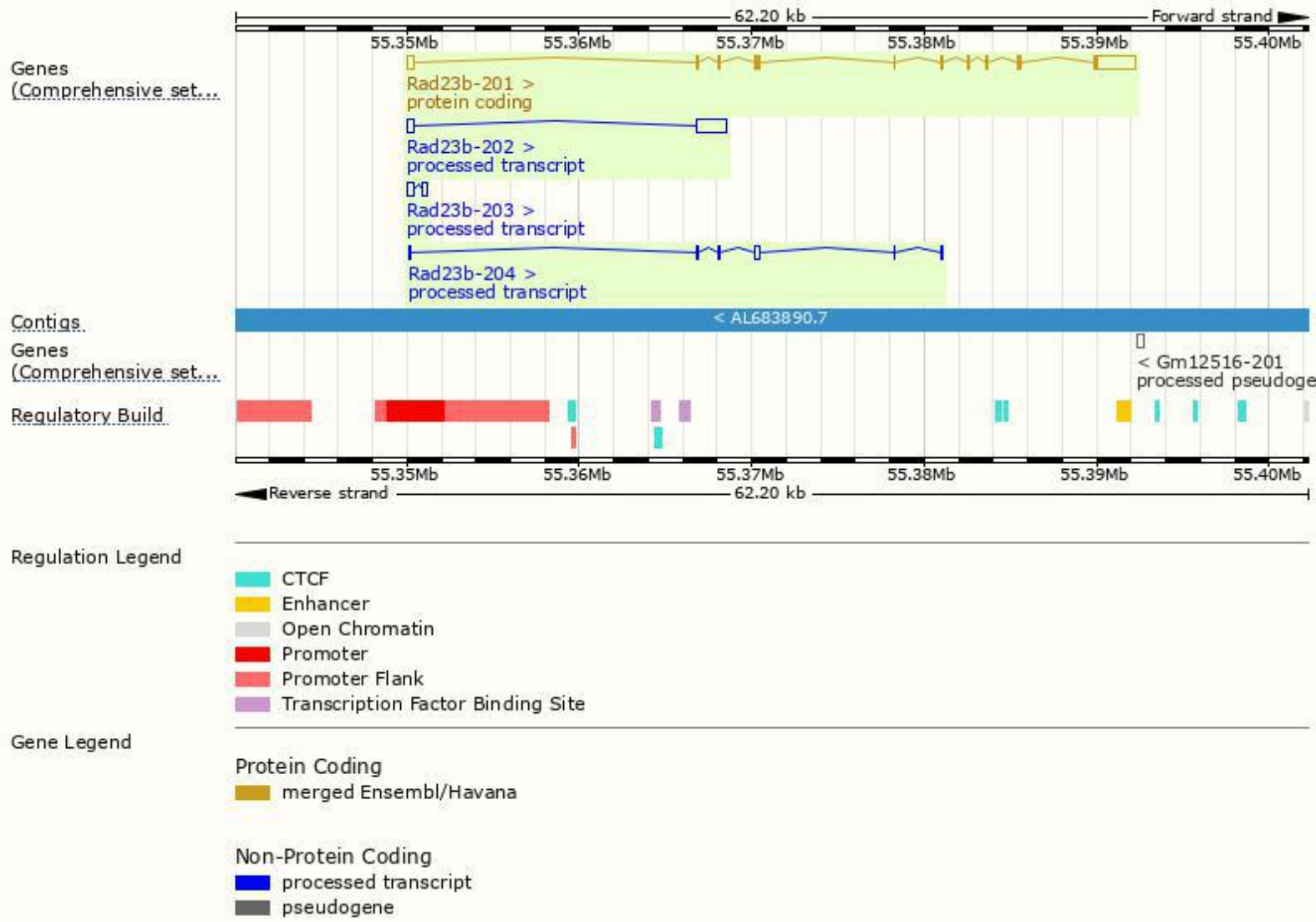
The gene has 4 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Rad23b-201	<a href="#">ENSMUST00000030134.8</a>	3810	<a href="#">416aa</a>	Protein coding	<a href="#">CCDS18194</a>	<a href="#">P54728</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
Rad23b-202	<a href="#">ENSMUST00000127266.1</a>	2176	No protein	Processed transcript	-	-	TSL:1
Rad23b-204	<a href="#">ENSMUST00000156263.1</a>	776	No protein	Processed transcript	-	-	TSL:3
Rad23b-203	<a href="#">ENSMUST00000148719.1</a>	663	No protein	Processed transcript	-	-	TSL:1

The strategy is based on the design of *Rad23b-201* transcript,the transcription is shown below:

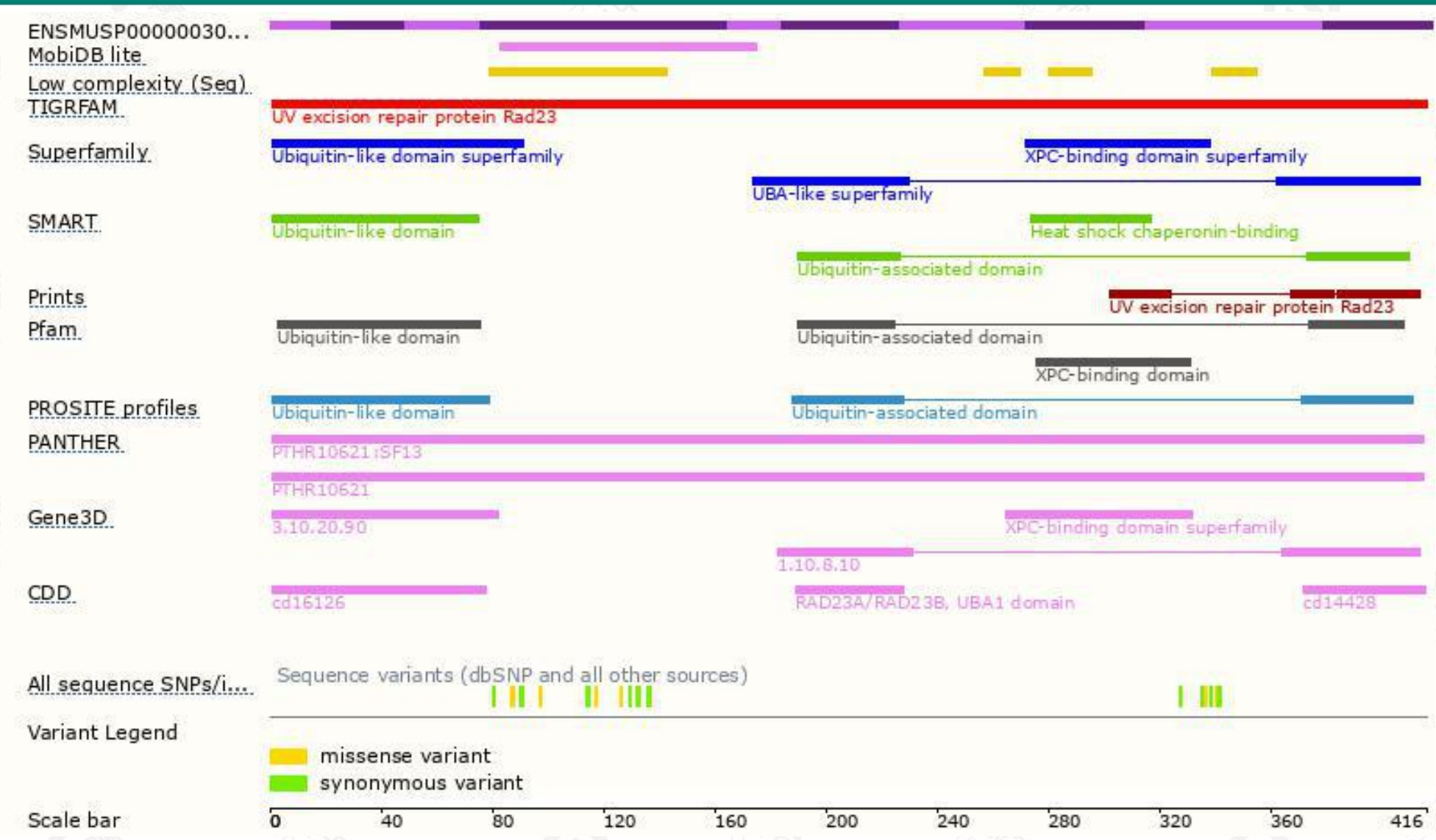


# Genomic location distribution

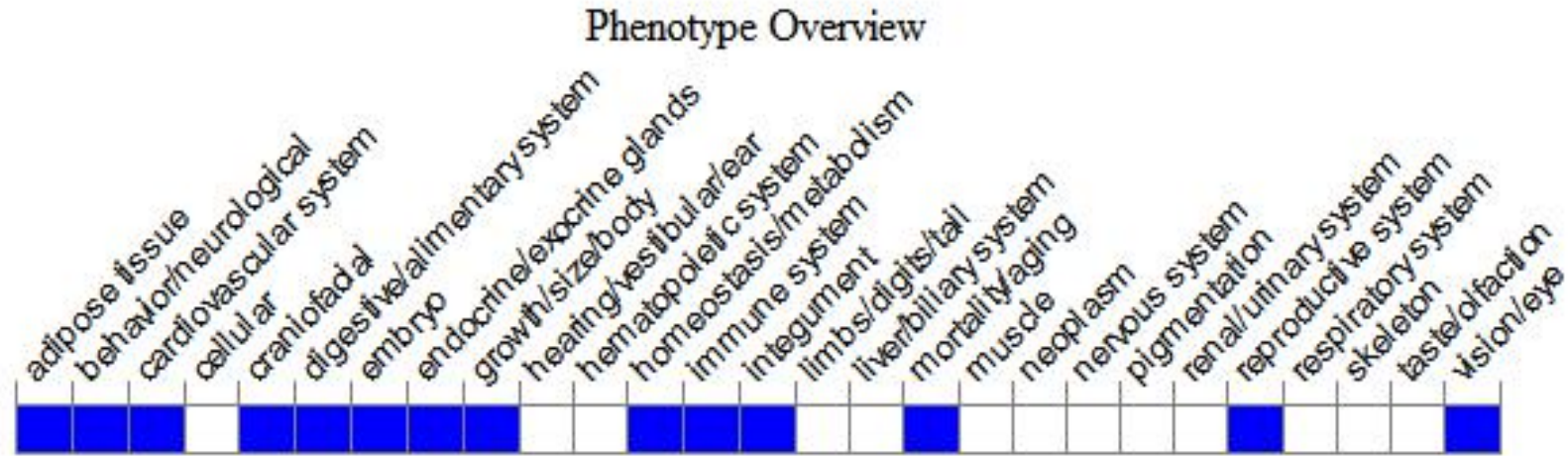




# Protein domain



# 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, mice homozygous for a disruption in this gene usually die around the time of birth. Those that survive show growth retardation, eye, reproductive, behavioral, and digestive system abnormalities. They usually die within 1 year of birth.

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