

# *Isl1* Cas9-KO Strategy

**Designer: Jinling Wang**

**Reviewer: Miaomiao Cui**

**Design Date: 2019-1-16**

# Project Overview

**Project Name**

*Isl1*

**Project type**

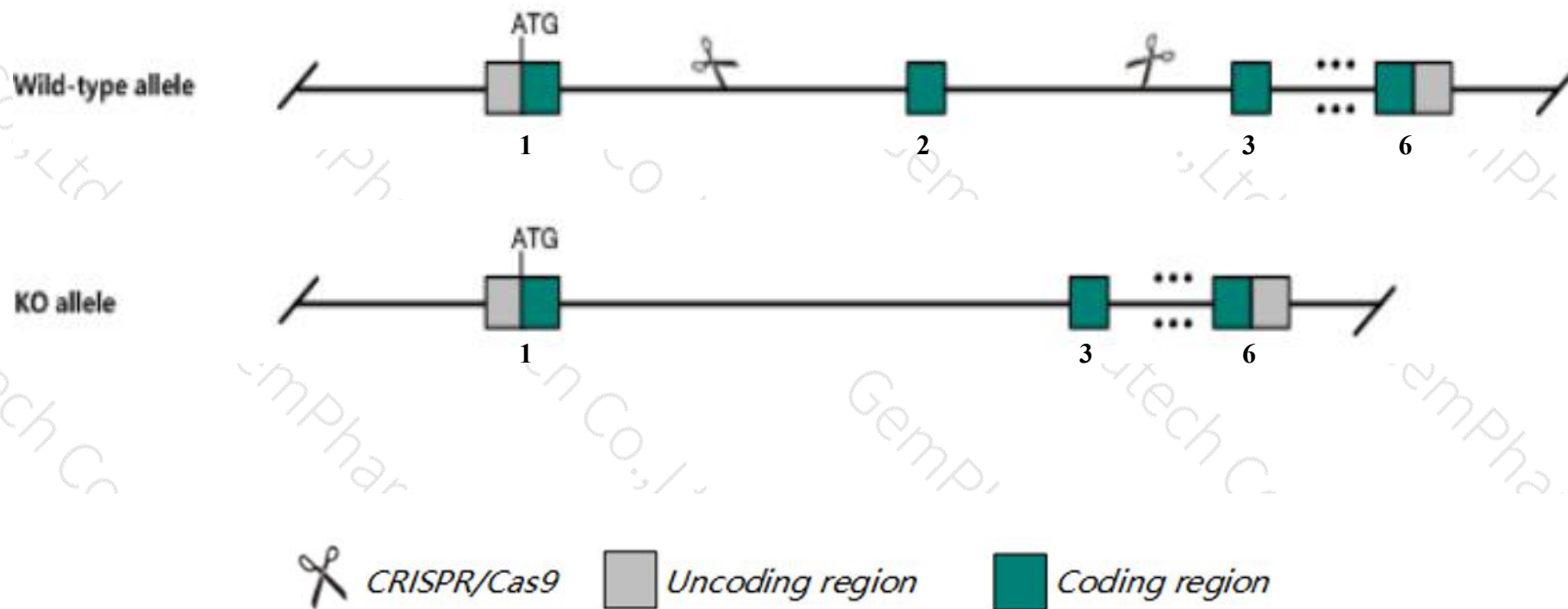
**Cas9-KO**

**Strain background**

**C57BL/6JGpt**

# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Isl1* gene. The schematic diagram is as follows:



- The *Isl1* gene has 5 transcripts. According to the structure of *Isl1* gene, exon2 of *Isl1-201*(ENSMUST00000036060.12) transcript is recommended as the knockout region. The region contains 190bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Isl1* 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, homozygotes for a targeted null mutation fail to develop motor neurons and die by embryonic day 11.5 with abnormal heart and pancreas development. Mice heterozygous for an ENU mutation exhibit chronic otitis media and hearing loss.
- The *Isl1* 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)

## Isl1 ISL1 transcription factor, LIM/homeodomain [Mus musculus (house mouse)]

Gene ID: 16392, updated on 13-Mar-2020

### Summary



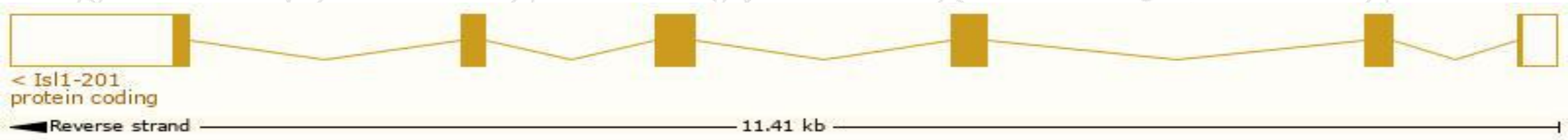
<b>Official Symbol</b>	Isl1 provided by <a href="#">MGI</a>
<b>Official Full Name</b>	ISL1 transcription factor, LIM/homeodomain provided by <a href="#">MGI</a>
<b>Primary source</b>	<a href="#">MGI:MGI:101791</a>
<b>See related</b>	<a href="#">Ensembl:ENSMUSG00000042258</a>
<b>Gene type</b>	protein 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>Expression</b>	Biased expression in whole brain E14.5 (RPKM 10.4), CNS E14 (RPKM 9.7) and 9 other tissues <a href="#">See more</a>
<b>Orthologs</b>	<a href="#">human</a> <a href="#">all</a>

# Transcript information (Ensembl)

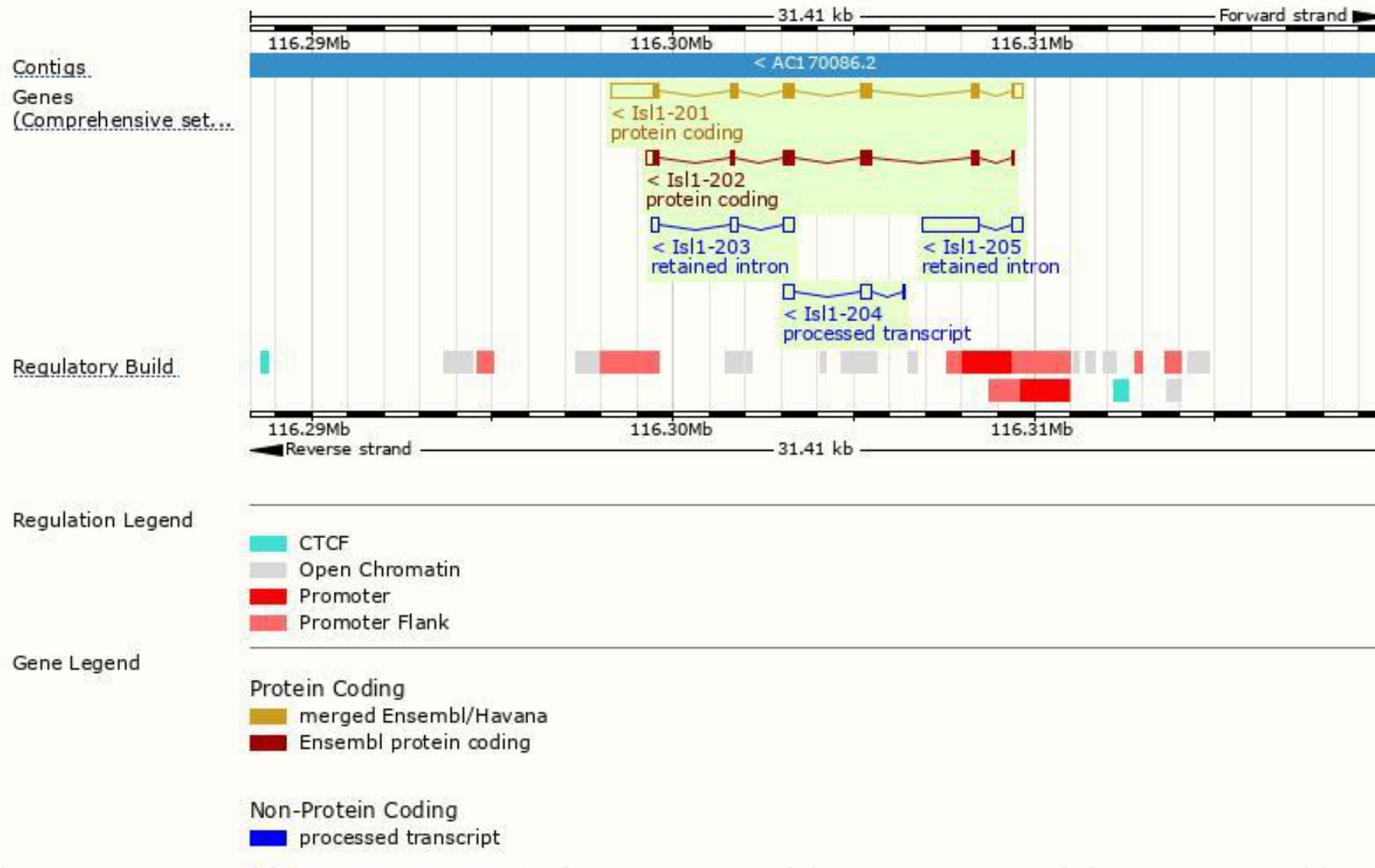
The gene has 5 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Isl1-201	<a href="#">ENSMUST00000036060.12</a>	2516	<a href="#">349aa</a>	Protein coding	<a href="#">CCDS26790</a>	<a href="#">A2RSV5 P61372</a>	TSL:1 GENCODE basic APPRIS P1
Isl1-202	<a href="#">ENSMUST00000176044.2</a>	1198	<a href="#">326aa</a>	Protein coding	-	<a href="#">P61372</a>	TSL:1 GENCODE basic
Isl1-204	<a href="#">ENSMUST00000176812.2</a>	588	No protein	Processed transcript	-	-	TSL:3
Isl1-205	<a href="#">ENSMUST00000177469.2</a>	1864	No protein	Retained intron	-	-	TSL:1
Isl1-203	<a href="#">ENSMUST00000176444.1</a>	676	No protein	Retained intron	-	-	TSL:1

The strategy is based on the design of *Isl1-201* transcript,the transcription is shown below:

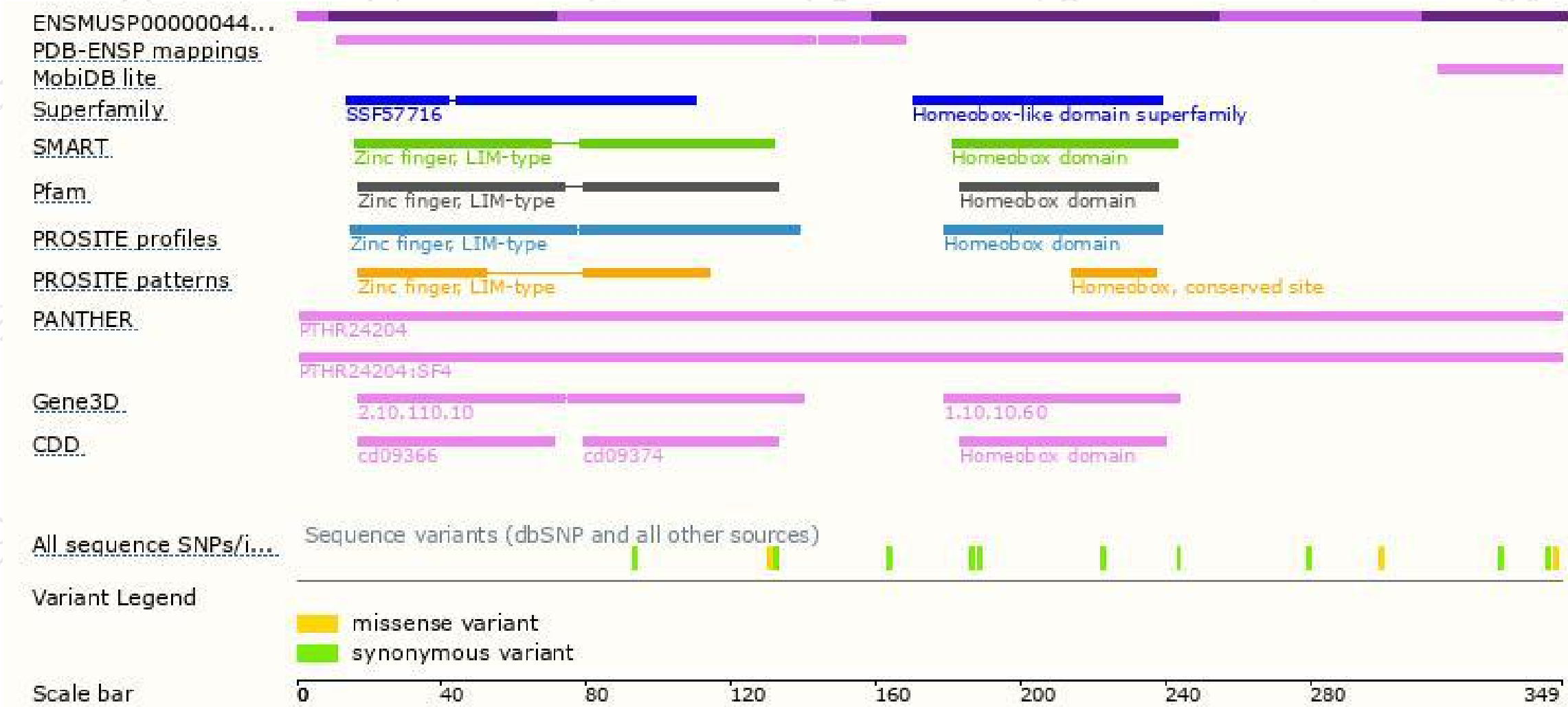


# Genomic location distribution

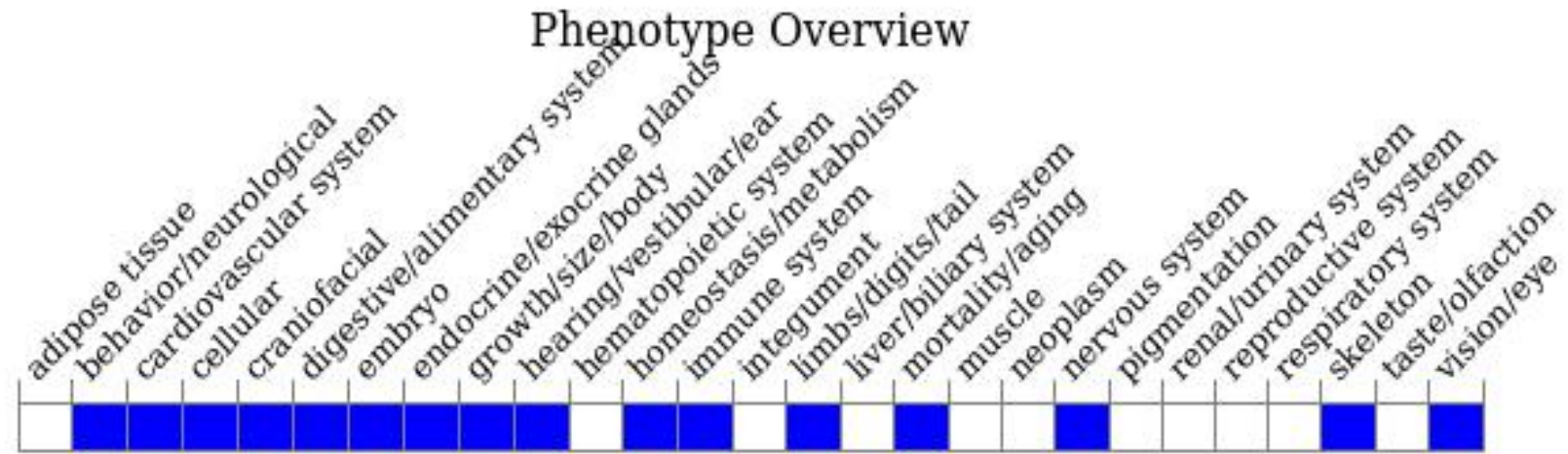




# Protein domain



# 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, homozygotes for a targeted null mutation fail to develop motor neurons and die by embryonic day 11.5 with abnormal heart and pancreas development. Mice heterozygous for an ENU mutation exhibit chronic otitis media and hearing loss.

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

Tel: 400-9660890

