

# *Pex1* Cas9-KO Strategy

Designer: Xueting Zhang

Reviewer: Yanhua Shen

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# Project Overview

**Project Name**

*Pex1*

**Project type**

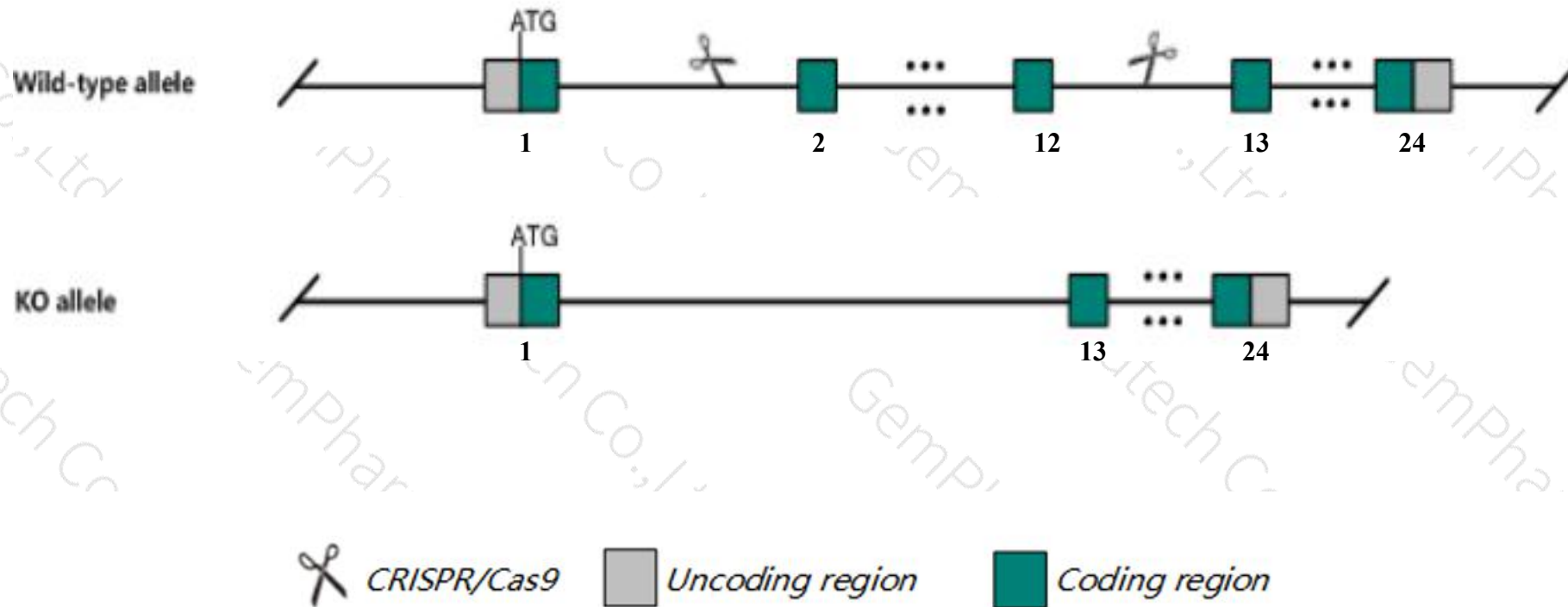
**Cas9-KO**

**Strain background**

**C57BL/6JGpt**

# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Pex1* gene. The schematic diagram is as follows:



- The *Pex1* gene has 14 transcripts. According to the structure of *Pex1* gene, exon2-exon12 of *Pex1-202* (ENSMUST00000121291.7) transcript is recommended as the knockout region. The region contains 1945bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Pex1* gene. The brief process is as follows: CRISPR/Cas9 system v

- According to the existing MGI data, Mice homozygous for a knock-in allele display premature death, postnatal growth retardation, fatty livers, a bile acid defect associated with intestinal lipid malabsorption and cholestasis, and a retinopathy associated with retinal cone cell degeneration and abnormal cone and rod electrophysiology.
- The knockout region is near to the N-terminal of *Rbm48* gene, this strategy may influence the regulatory function of the N-terminal of *Rbm48* gene.
- Transcript *Pex1*-2029&210&211&212&214 may not be affected.
- The *Pex1* gene is located on the Chr5. 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)

## Pex1 peroxisomal biogenesis factor 1 [ *Mus musculus* (house mouse) ]

Gene ID: 71382, updated on 25-Feb-2020

### Summary

- Official Symbol** Pex1 provided by [MGI](#)
- Official Full Name** peroxisomal biogenesis factor 1 provided by [MGI](#)
- Primary source** [MGI:MGI:1918632](#)
- See related** [Ensembl:ENSMUSG00000005907](#)
- 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** ZWS1; 5430414H02Rik; E330005K07Rik
- Expression** Ubiquitous expression in placenta adult (RPKM 4.3), CNS E14 (RPKM 3.6) and 28 other tissues [See more](#)
- Orthologs** [human](#) [all](#)

### Genomic context

**Location:** 5; 5 A1

See Pex1 in [Genome Data Viewer](#)

**Exon count:** 25

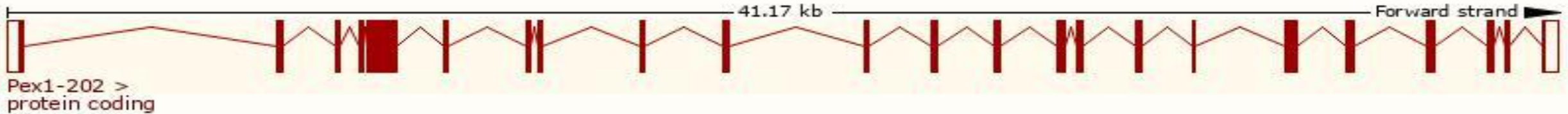
Annotation release	Status	Assembly	Chr	Location
<a href="#">108</a>	current	GRCm38.p6 ( <a href="#">GCF_000001635.26</a> )	5	NC_000071.6 (3596066..3637230)
Build 37.2	previous assembly	MGSCv37 ( <a href="#">GCF_000001635.18</a> )	5	NC_000071.5 (3596066..3637101)

# Transcript information (Ensembl)

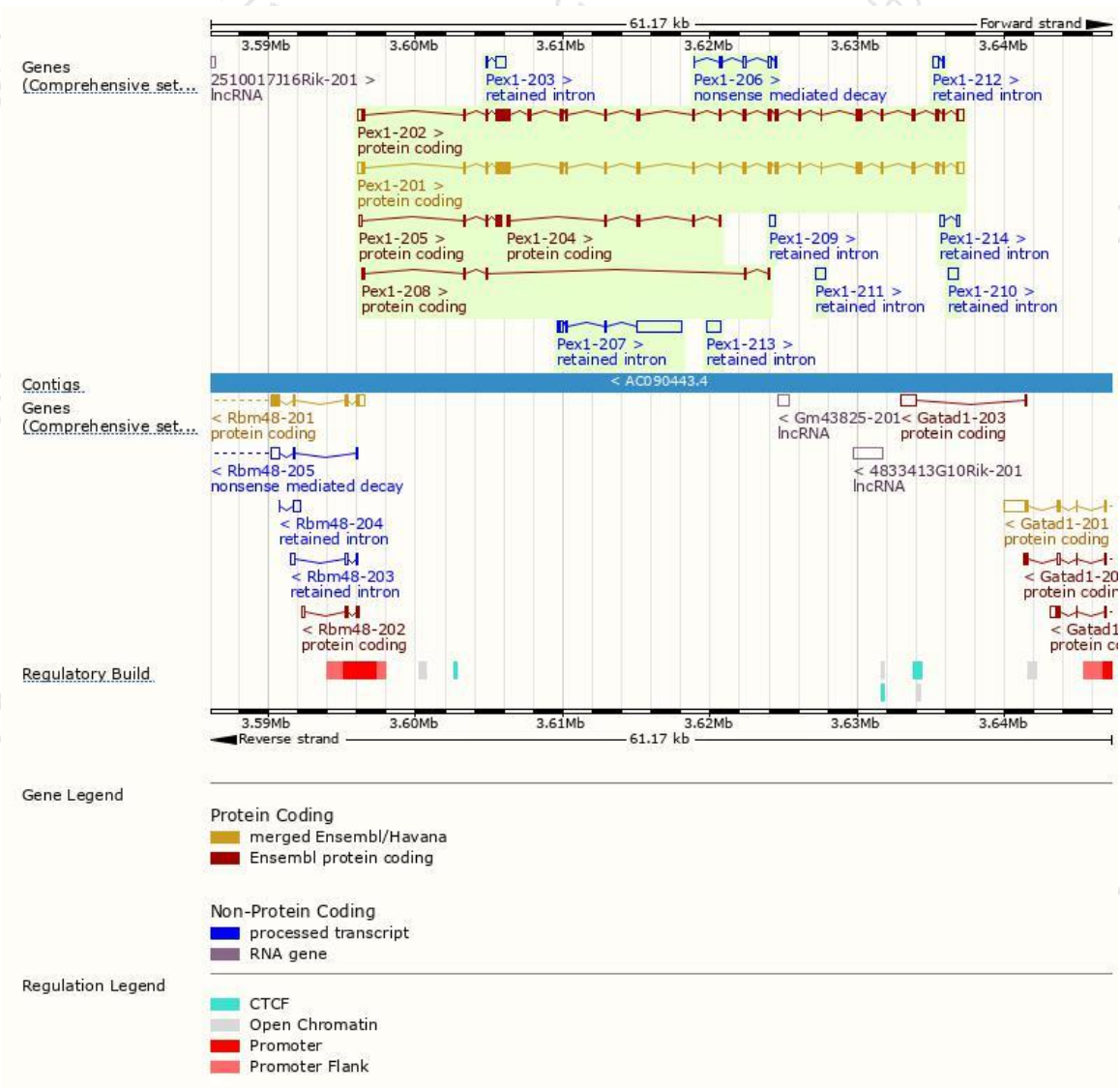
The gene has 14 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Pex1-202	<a href="#">ENSMUST00000121291.7</a>	4555	<a href="#">1284aa</a>	Protein coding	<a href="#">CCDS80201</a>	<a href="#">Q5BL07</a>	TSL:5 GENCODE basic APPRIS ALT2
Pex1-201	<a href="#">ENSMUST0000006061.12</a>	4433	<a href="#">1244aa</a>	Protein coding	<a href="#">CCDS19065</a>	<a href="#">Q5BL07</a>	TSL:1 GENCODE basic APPRIS P3
Pex1-205	<a href="#">ENSMUST00000142516.1</a>	727	<a href="#">69aa</a>	Protein coding	-	<a href="#">D3Z5A7</a>	CDS 3' incomplete TSL:3
Pex1-204	<a href="#">ENSMUST00000126545.1</a>	639	<a href="#">213aa</a>	Protein coding	-	<a href="#">F6RUH9</a>	5' and 3' truncations in transcript evidence prevent annotation of the start and the end of the CDS. CDS 5' and 3' incomplete TSL:3
Pex1-208	<a href="#">ENSMUST00000195894.1</a>	528	<a href="#">143aa</a>	Protein coding	-	<a href="#">A0A0G2JE39</a>	TSL:3 GENCODE basic
Pex1-206	<a href="#">ENSMUST00000143132.1</a>	664	<a href="#">76aa</a>	Nonsense mediated decay	-	<a href="#">F7CF88</a>	CDS 5' incomplete TSL:5
Pex1-207	<a href="#">ENSMUST00000143959.1</a>	3491	No protein	Retained intron	-	-	TSL:1
Pex1-213	<a href="#">ENSMUST00000199035.1</a>	987	No protein	Retained intron	-	-	TSL:NA
Pex1-203	<a href="#">ENSMUST00000123268.1</a>	746	No protein	Retained intron	-	-	TSL:3
Pex1-210	<a href="#">ENSMUST00000196432.1</a>	668	No protein	Retained intron	-	-	TSL:NA
Pex1-211	<a href="#">ENSMUST00000196692.1</a>	638	No protein	Retained intron	-	-	TSL:NA
Pex1-214	<a href="#">ENSMUST00000199213.1</a>	487	No protein	Retained intron	-	-	TSL:2
Pex1-212	<a href="#">ENSMUST00000197167.1</a>	474	No protein	Retained intron	-	-	TSL:2
Pex1-209	<a href="#">ENSMUST00000196124.1</a>	403	No protein	Retained intron	-	-	TSL:NA

The strategy is based on the design of *Pex1-202* transcript,The transcription is shown below

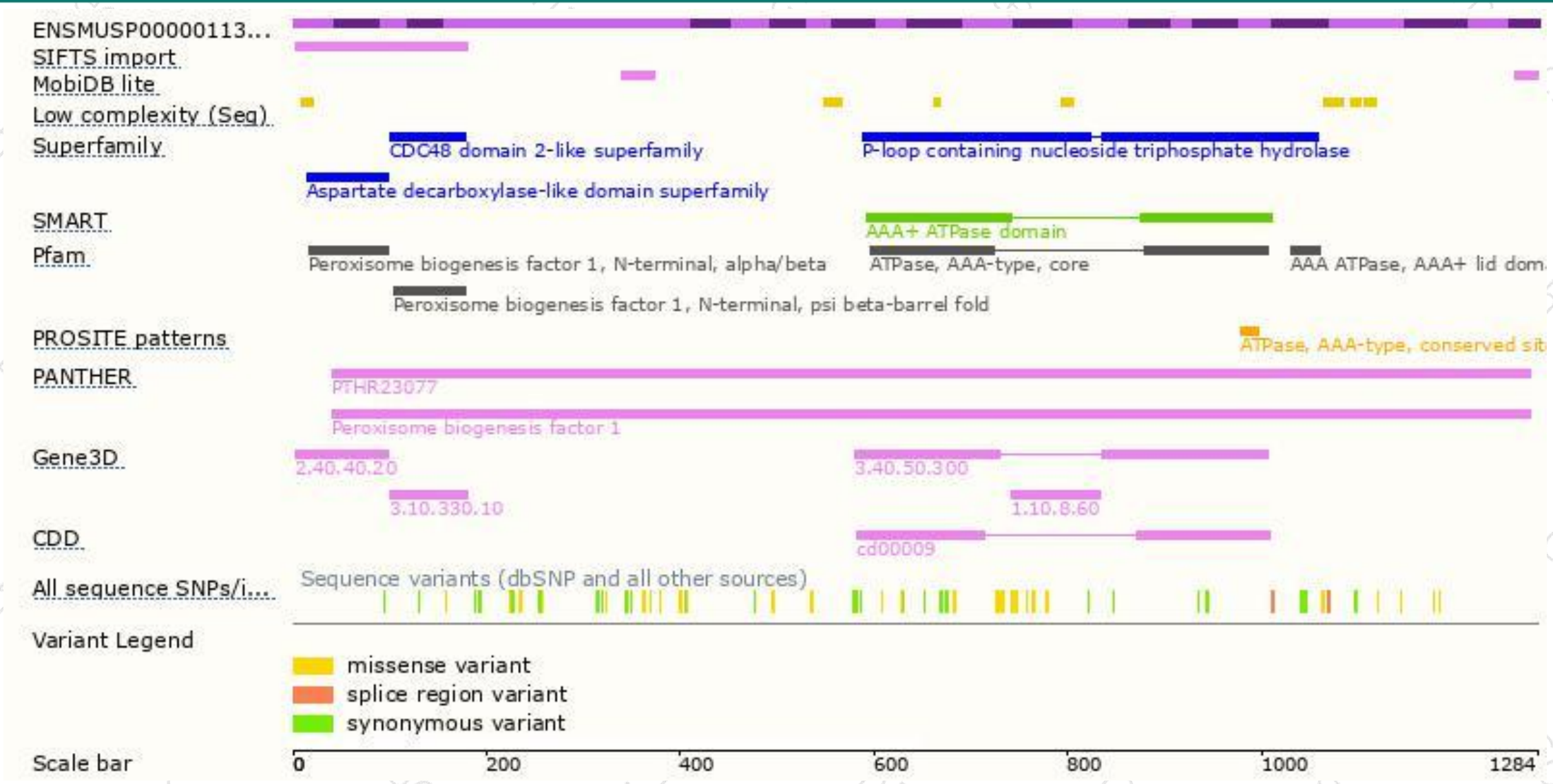


# Genomic location distribution

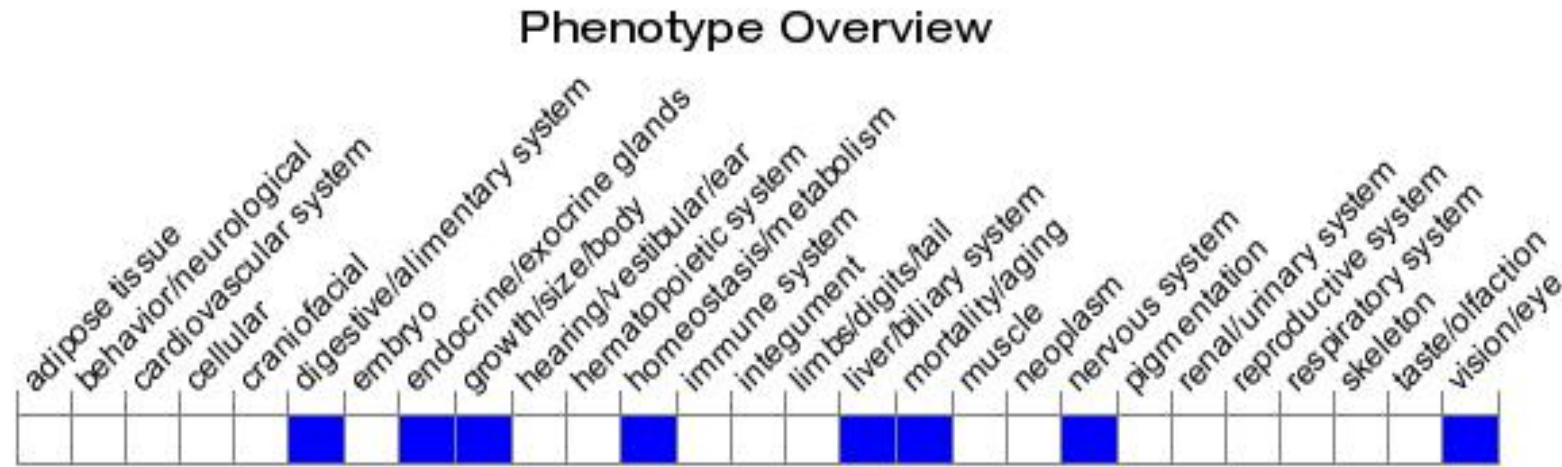




# Protein domain



# 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 knock-in allele display premature death, postnatal growth retardation, fatty livers, a bile acid defect associated with intestinal lipid malabsorption and cholestasis, and a retinopathy associated with retinal cone cell degeneration and abnormal cone and rod electrophysiology.

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

Tel: 400-9660890

