

# ***Kat2a Cas9-KO Strategy***

**Designer:**

**Daohua Xu**

**Reviewer :**

**Huimin Su**

**Design Date:**

**2019-10-12**

# Project Overview



---

**Project Name**

***Kat2a***

---

**Project type**

**Cas9-KO**

---

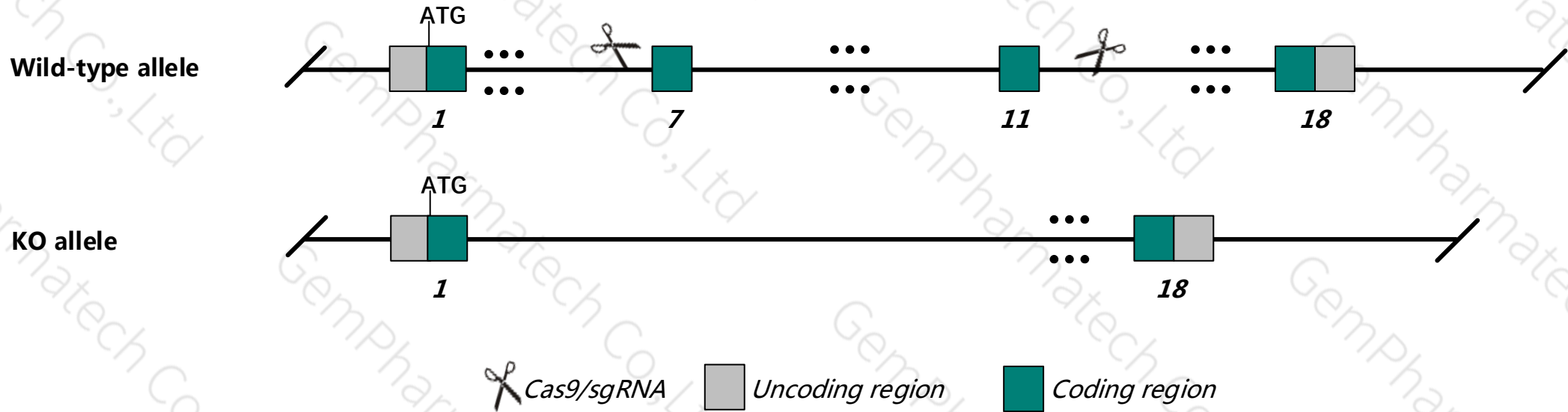
**Strain background**

**C57BL/6JGpt**

---

# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Kat2a* gene. The schematic diagram is as follows:



# Technical routes

- The *Kat2a* gene has 4 transcripts. According to the structure of *Kat2a* gene, exon7-exon11 of *Kat2a*-202 (ENSMUST00000103118.3) transcript is recommended as the knockout region. The region contains 685bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Kat2a* gene. The brief process is as follows: sgRNA was transcribed in vitro. Cas9, sgRNA 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 targeted null mutations exhibit poorly developed yolk sac blood vessels, retarded growth, absence of dorsal mesoderm lineages, failure to form somites, and lethality between embryonic days 9.5-11.5.
- The target gene of this strategy will retain 1/3 of the protein at the N-terminus.
- The *Kat2a* gene is located on the Chr11. 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 knockout on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.

# Gene information ( NCBI )

## Kat2a K(lysine) acetyltransferase 2A [ *Mus musculus* (house mouse) ]

Gene ID: 14534, updated on 11-Sep-2019

### Summary

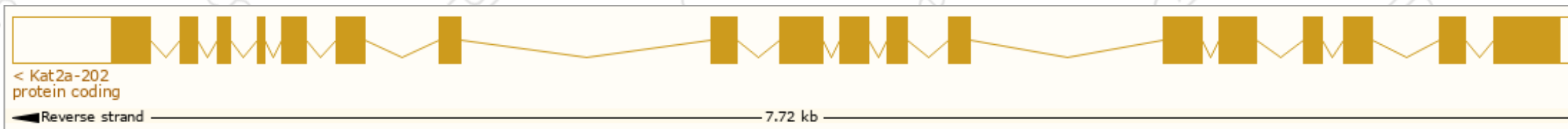
<b>Official Symbol</b>	Kat2a provided by <a href="#">MGI</a>
<b>Official Full Name</b>	K(lysine) acetyltransferase 2A provided by <a href="#">MGI</a>
<b>Primary source</b>	<a href="#">MGI:MGI:1343101</a>
<b>See related</b>	<a href="#">Ensembl:ENSMUSG00000020918</a>
<b>Gene type</b>	protein coding
<b>RefSeq 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>	Gcn5; Gcn5l2; mmGCN5; AW212720; 1110051E14Rik
<b>Expression</b>	Ubiquitous expression in adrenal adult (RPKM 33.5), ovary adult (RPKM 29.4) and 28 other tissues <a href="#">See more</a>
<b>Orthologs</b>	<a href="#">human</a> <a href="#">all</a>

# Transcript information ( Ensembl )

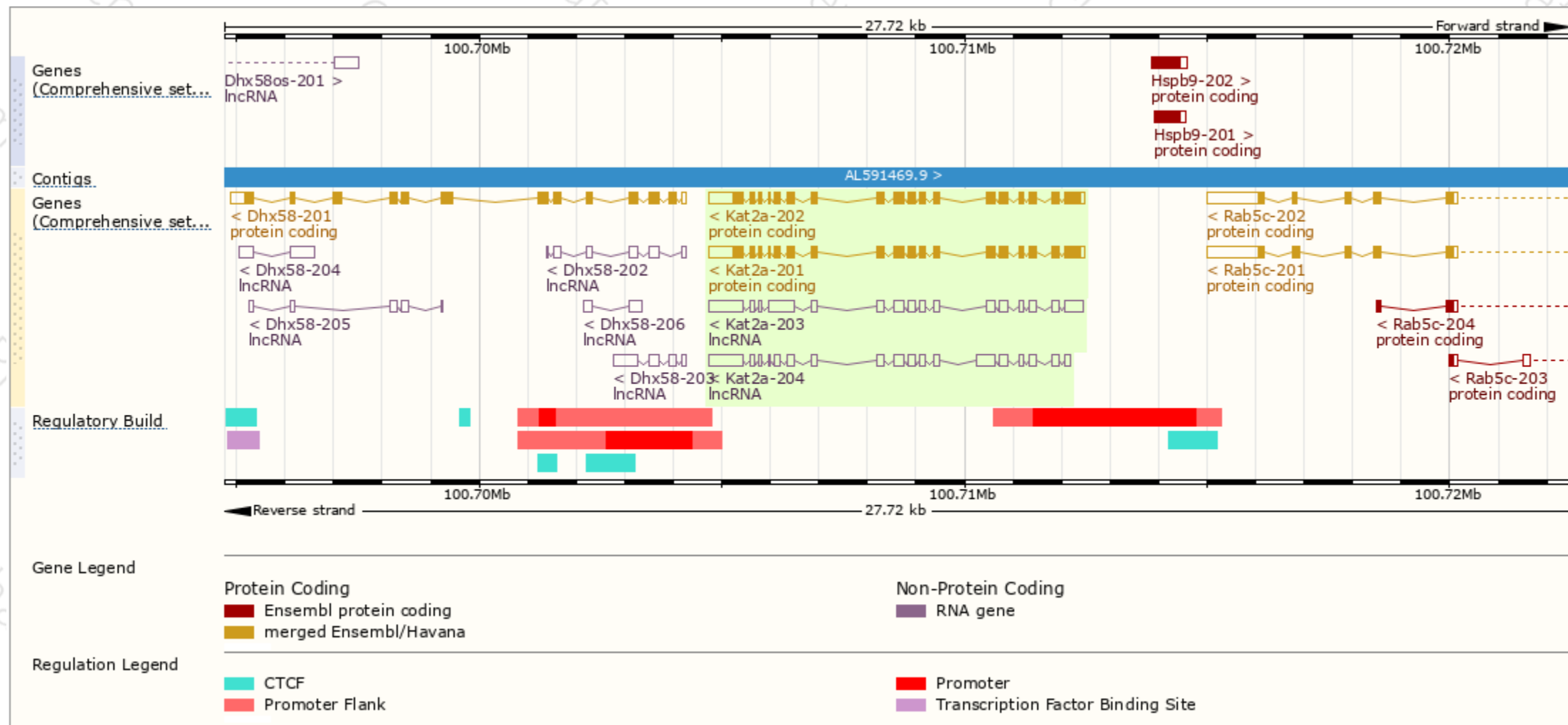
The gene has 4 transcripts, and all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Kat2a-202	<a href="#">ENSMUST00000103118.3</a>	3046	<a href="#">830aa</a>	Protein coding	<a href="#">CCDS25433</a>	<a href="#">Q9JHD2</a>	TSL:1 GENCODE basic APPRIS P4
Kat2a-201	<a href="#">ENSMUST00000006973.11</a>	3043	<a href="#">829aa</a>	Protein coding	<a href="#">CCDS25432</a>	<a href="#">Q6P3Z8</a>	TSL:1 GENCODE basic APPRIS ALT2
Kat2a-203	<a href="#">ENSMUST00000126299.1</a>	3269	No protein	lncRNA	-	-	TSL:1
Kat2a-204	<a href="#">ENSMUST00000153526.7</a>	2946	No protein	lncRNA	-	-	TSL:2

The strategy is based on the design of *Kat2a-202* transcript, The transcription is shown below

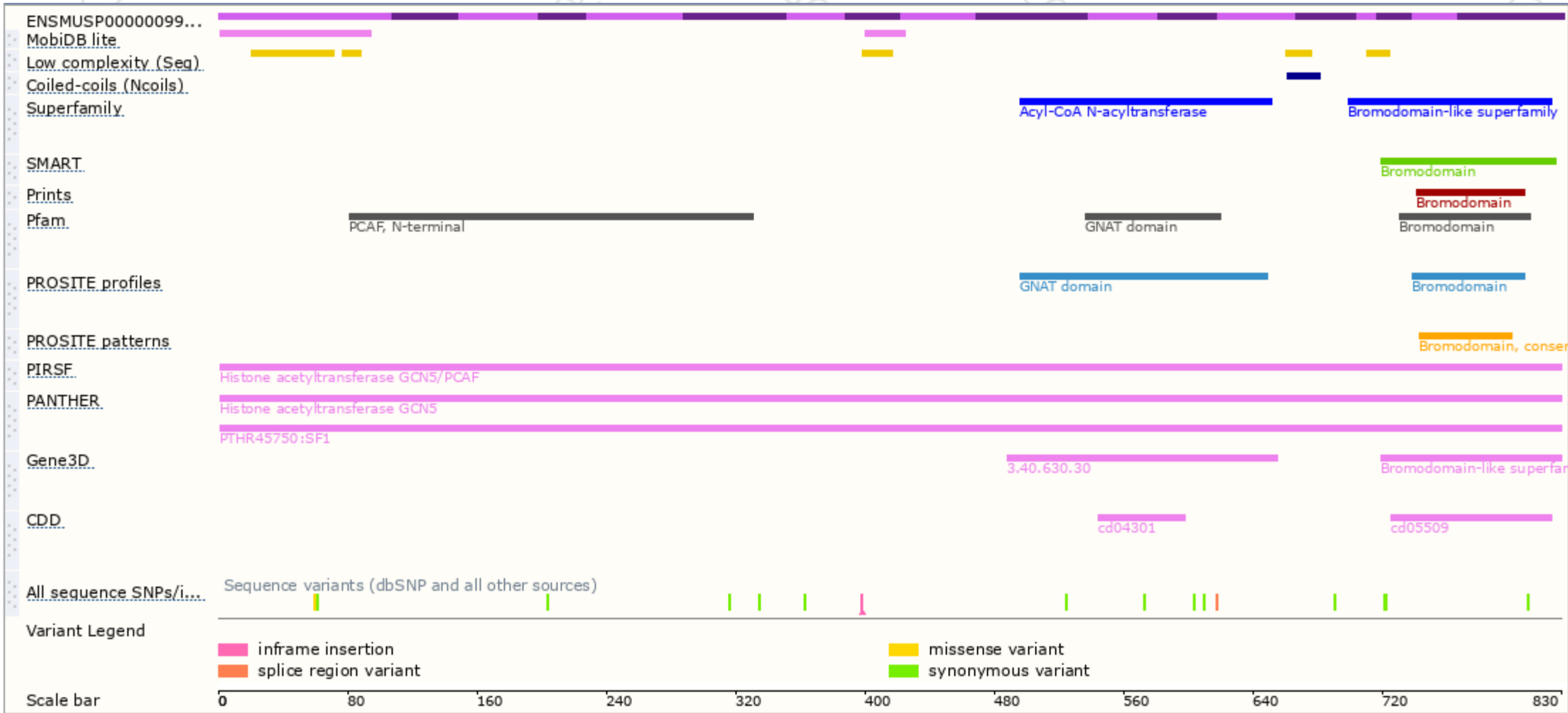


# Genomic location distribution

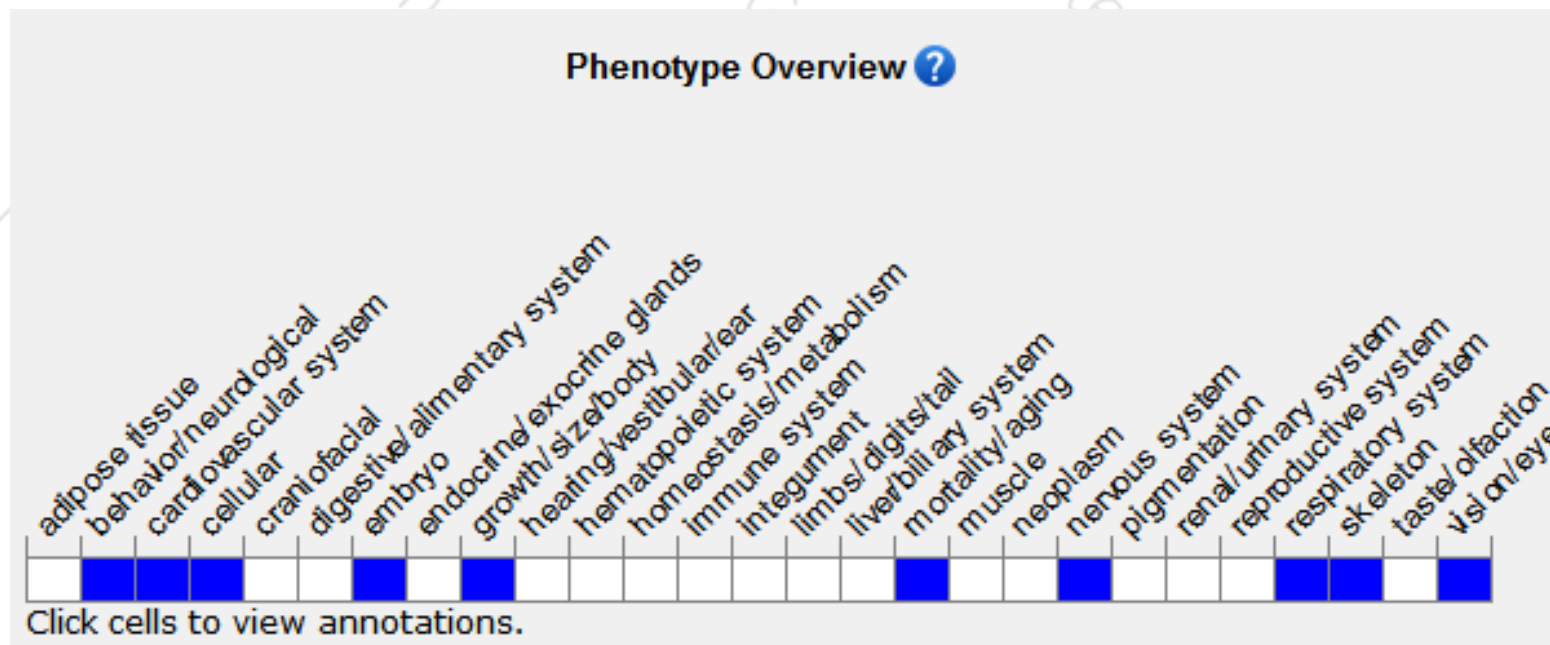




# Protein domain



# 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, Homozygotes for targeted null mutations exhibit poorly developed yolk sac blood vessels, retarded growth, absence of dorsal mesoderm lineages, failure to form somites, and lethality between embryonic days 9.5-11.5.

If you have any questions, you are welcome to inquire.  
Tel: 025-5864 1534

