

# ***Kcnk2* Cas9-CKO Strategy**

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Date: 2019-11-13

# Project Overview

**Project Name**

***Kcnk2***

**Project type**

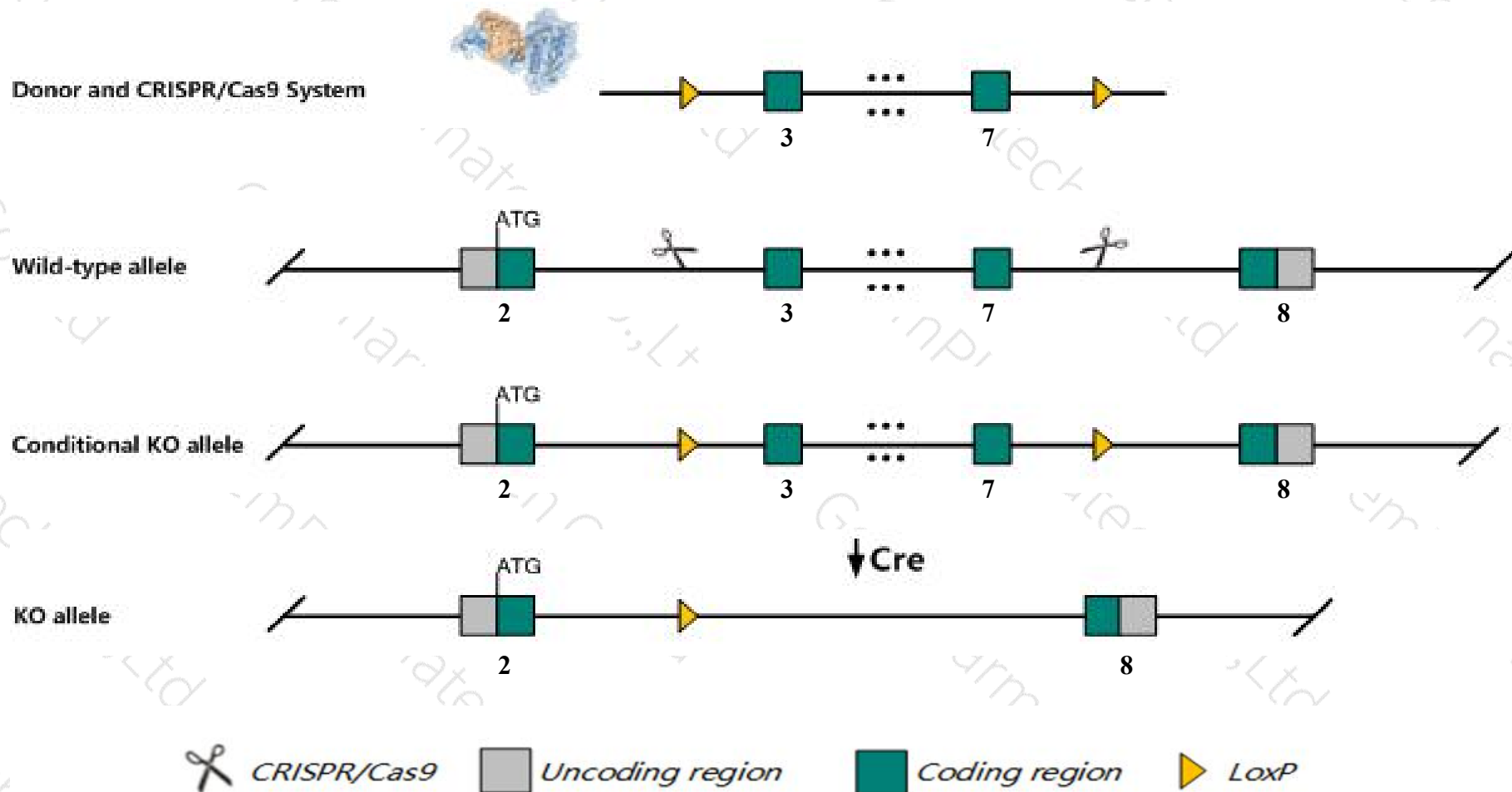
**Cas9-CKO**

**Strain background**

**C57BL/6JGpt**

# Conditional Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Kcnk2* gene. The schematic diagram is as follows:



- The *Kcnk2* gene has 10 transcripts. According to the structure of *Kcnk2* gene, exon3-exon7 of *Kcnk2*-202 (ENSMUST00000110920.6) transcript is recommended as the knockout region. The region contains 917bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Kcnk2* gene. The brief process is as follows: CRISPR/Cas9 system and Donor 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.
- The flox mice will be knocked out after mating with mice expressing Cre recombinase, resulting in the loss of function of the target gene in specific tissues and cell types.

- According to the existing MGI data, Homozygous null mice display increased sensitivity to pharmacologically induced seizures and ischemia.
- Transcript *Kcnk2*-207&210 may not be affected.
- The *Kcnk2* gene is located on the Chr1. 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 loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at existing technological level.



# Gene information (NCBI)

## Kcnk2 potassium channel, subfamily K, member 2 [ *Mus musculus* (house mouse) ]

Gene ID: 16526, updated on 16-Sep-2019

### Summary

|                    |                                                                                                                                                                           |
|--------------------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Official Symbol    | Kcnk2 provided by <a href="#">MGI</a>                                                                                                                                     |
| Official Full Name | potassium channel, subfamily K, member 2 provided by <a href="#">MGI</a>                                                                                                  |
| Primary source     | <a href="#">MGI:MGI:109366</a>                                                                                                                                            |
| See related        | <a href="#">Ensembl:ENSMUSG00000037624</a>                                                                                                                                |
| Gene type          | protein coding                                                                                                                                                            |
| RefSeq status      | VALIDATED                                                                                                                                                                 |
| Organism           | <a href="#">Mus musculus</a>                                                                                                                                              |
| Lineage            | Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus |
| Also known as      | TREK-1; AI848635; A430027H14Rik                                                                                                                                           |
| Expression         | Broad expression in CNS E18 (RPKM 6.7), cortex adult (RPKM 5.7) and 19 other tissues <a href="#">See more</a>                                                             |
| Orthologs          | <a href="#">human</a> <a href="#">all</a>                                                                                                                                 |

### Genomic context

Location: 1; 1 H6

See Kcnk2 in [Genome Data Viewer](#)

Exon count: 9

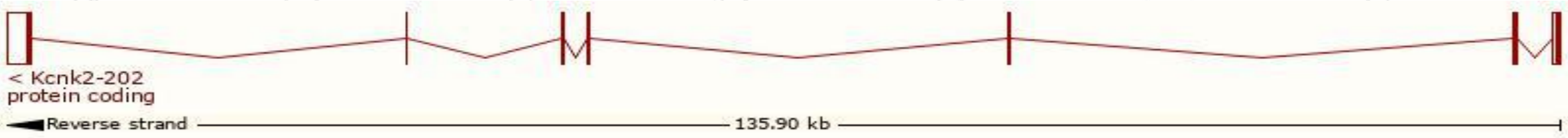
| Annotation release  | Status            | Assembly                                       | Chr | Location                                       |
|---------------------|-------------------|------------------------------------------------|-----|------------------------------------------------|
| <a href="#">108</a> | current           | GRCm38.p6 ( <a href="#">GCF_000001635.26</a> ) | 1   | NC_000067.6 (189207930..189402782, complement) |
| Build 37.2          | previous assembly | MGSCv37 ( <a href="#">GCF_000001635.18</a> )   | 1   | NC_000067.5 (191031809..191168071, complement) |

# Transcript information (Ensembl)

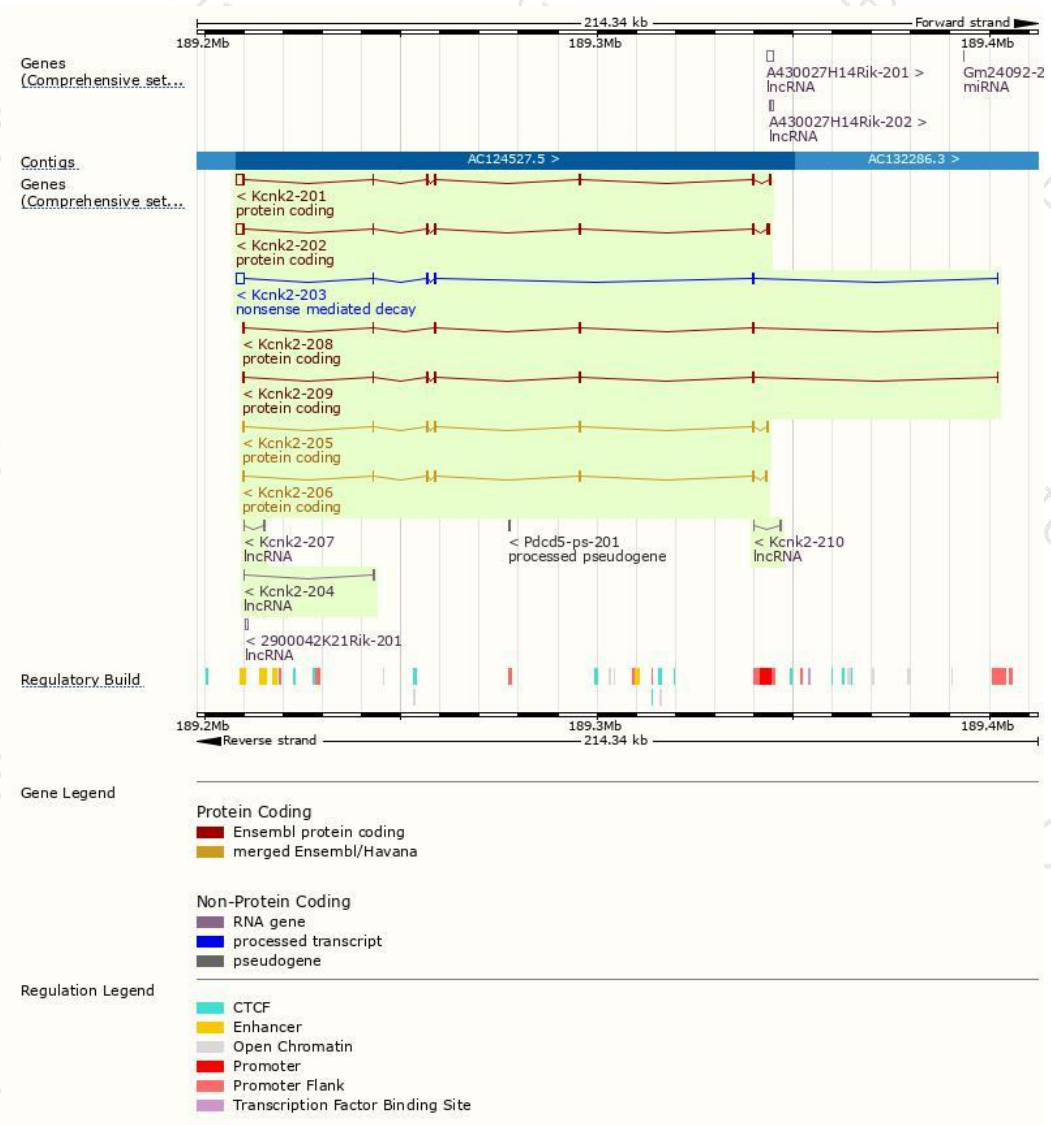
The gene has 10 transcripts,all transcripts are shown below:

| Name      | Transcript ID                         | bp   | Protein               | Biotype                 | CCDS                      | UniProt                    | Flags                           |
|-----------|---------------------------------------|------|-----------------------|-------------------------|---------------------------|----------------------------|---------------------------------|
| Kcnk2-202 | <a href="#">ENSMUST00000110920.6</a>  | 3572 | <a href="#">411aa</a> | Protein coding          | <a href="#">CCDS48481</a> | <a href="#">P97438</a>     | TSL:5 GENCODE basic APPRIS ALT1 |
| Kcnk2-201 | <a href="#">ENSMUST00000079451.12</a> | 3297 | <a href="#">414aa</a> | Protein coding          | <a href="#">CCDS78773</a> | <a href="#">Q6P6P9</a>     | TSL:1 GENCODE basic APPRIS ALT1 |
| Kcnk2-205 | <a href="#">ENSMUST00000192723.1</a>  | 1642 | <a href="#">411aa</a> | Protein coding          | <a href="#">CCDS48481</a> | <a href="#">P97438</a>     | TSL:1 GENCODE basic APPRIS ALT1 |
| Kcnk2-206 | <a href="#">ENSMUST00000193319.5</a>  | 1548 | <a href="#">426aa</a> | Protein coding          | <a href="#">CCDS48480</a> | <a href="#">P97438</a>     | TSL:1 GENCODE basic APPRIS P4   |
| Kcnk2-209 | <a href="#">ENSMUST00000194402.5</a>  | 1373 | <a href="#">422aa</a> | Protein coding          | -                         | <a href="#">A0A0A6YXK1</a> | TSL:5 GENCODE basic APPRIS ALT1 |
| Kcnk2-208 | <a href="#">ENSMUST00000194172.5</a>  | 1200 | <a href="#">237aa</a> | Protein coding          | -                         | <a href="#">A0A0A6YXX0</a> | TSL:5 GENCODE basic             |
| Kcnk2-203 | <a href="#">ENSMUST00000180044.7</a>  | 3131 | <a href="#">152aa</a> | Nonsense mediated decay | -                         | <a href="#">A0A0R4J279</a> | TSL:1                           |
| Kcnk2-210 | <a href="#">ENSMUST00000195016.1</a>  | 493  | No protein            | lncRNA                  | -                         | -                          | TSL:3                           |
| Kcnk2-204 | <a href="#">ENSMUST00000192529.1</a>  | 384  | No protein            | lncRNA                  | -                         | -                          | TSL:5                           |
| Kcnk2-207 | <a href="#">ENSMUST00000193686.1</a>  | 334  | No protein            | lncRNA                  | -                         | -                          | TSL:3                           |

The strategy is based on the design of *Kcnk2-202* transcript,The transcription is shown below

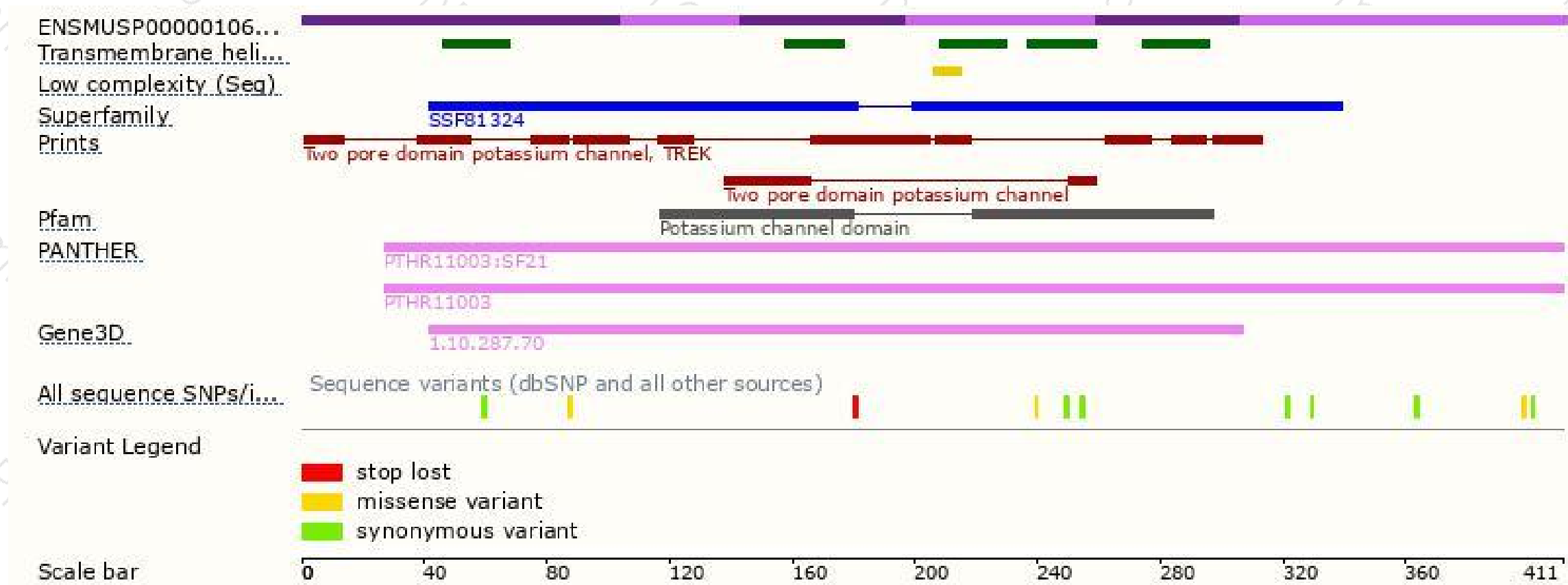


# Genomic location distribution

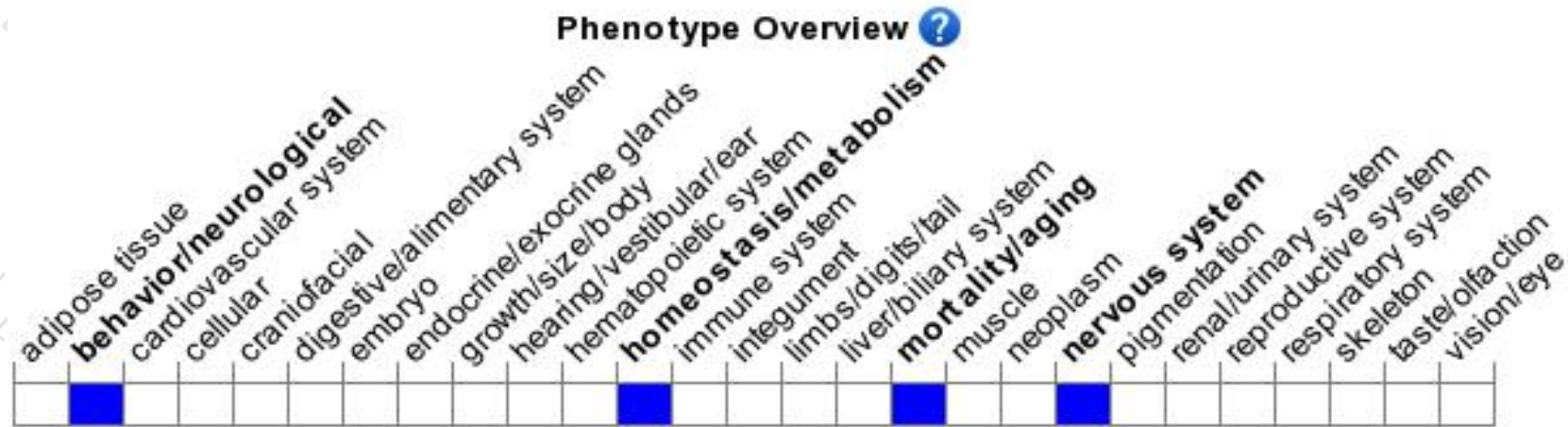




# Protein domain



# 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, Homozygous null mice display increased sensitivity to pharmacologically induced seizures and ischemia.

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

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