

# *Ap3m2* Cas9-KO Strategy

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**Design Date:** 2020-5-12

# Project Overview

**Project Name**

*Ap3m2*

**Project type**

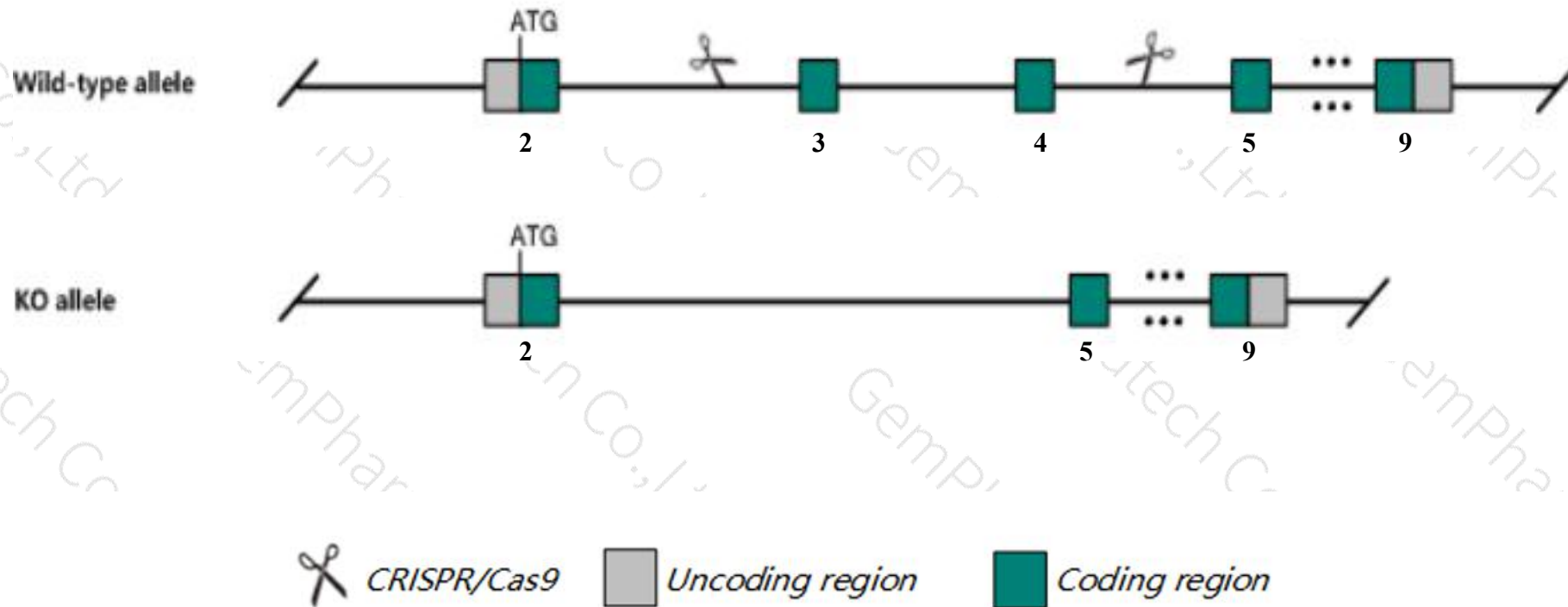
**Cas9-KO**

**Strain background**

**C57BL/6JGpt**

# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Ap3m2* gene. The schematic diagram is as follows:



- The *Ap3m2* gene has 4 transcripts. According to the structure of *Ap3m2* gene, exon3-exon4 of *Ap3m2-201* (ENSMUST00000163739.2) transcript is recommended as the knockout region. The region contains 310bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Ap3m2* gene. The brief process is as follows: CRISPR/Cas9 system

- According to the existing MGI data, homozygous null mice suffer from spontaneous recurrent epileptic seizures, are more susceptible to drug-induced seizures and show impaired gaba release, fewer synaptic vesicles, enhanced long-term potentiation, and abnormal propagation of neuronal excitability via the temporoammonic pathway.
- The *Ap3m2* gene is located on the Chr8. 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)

## Ap3m2 adaptor-related protein complex 3, mu 2 subunit [Mus musculus (house mouse)]

Gene ID: 64933, updated on 13-Mar-2020

### Summary



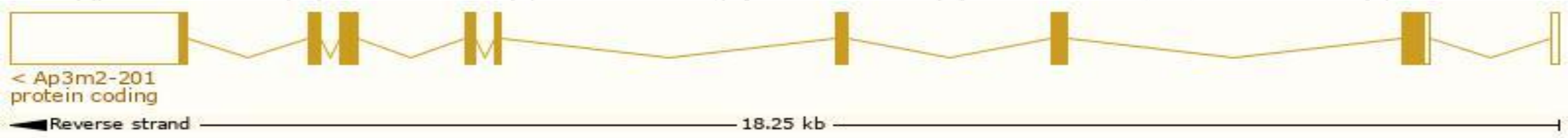
|                           |   |
|---------------------------|---|
| <b>Official Symbol</b>    | Ap3m2 provided by <a href="#">MGI</a>   |
| <b>Official Full Name</b> | adaptor-related protein complex 3, mu 2 subunit provided by <a href="#">MGI</a>   |
| <b>Primary source</b>     | <a href="#">MGI:MGI:1929214</a>   |
| <b>See related</b>        | <a href="#">Ensembl:ENSMUSG000000031539</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>      | 5830445E16Rik, AP-3B  |
| <b>Expression</b>         | Broad expression in cortex adult (RPKM 20.8), frontal lobe adult (RPKM 18.2) and 19 other tissues <a href="#">See more</a>  |
| <b>Orthologs</b>          | <a href="#">human</a> <a href="#">all</a>   |

# Transcript information（Ensembl）

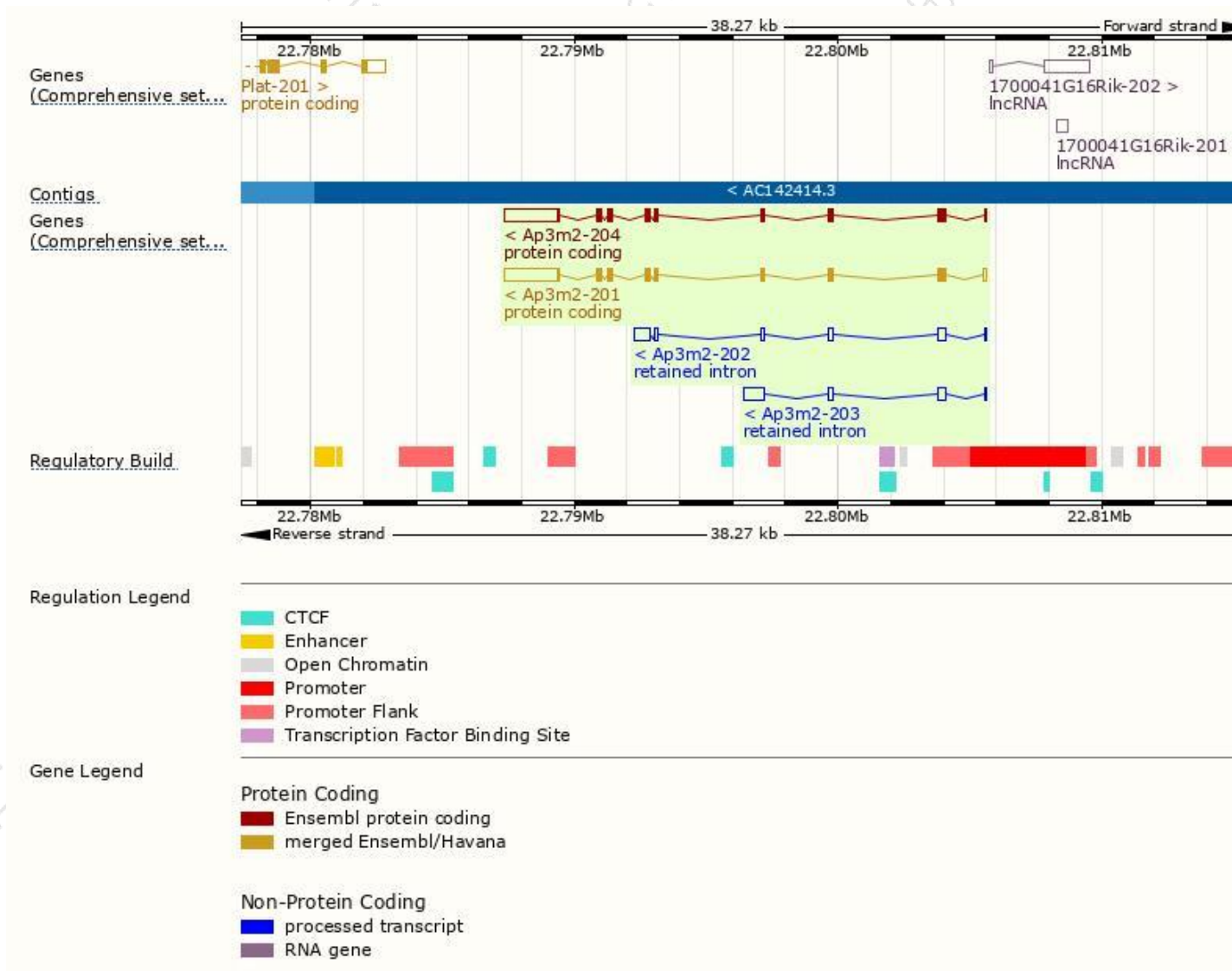
The gene has 4 transcripts,all transcripts are shown below:

| Name      | Transcript ID                        | bp   | Protein               | Biotype         | CCDS                      | UniProt                | Flags   |
|-----------|--------------------------------------|------|-----------------------|-----------------|---------------------------|------------------------|---|
| Ap3m2-201 | <a href="#">ENSMUST00000163739.2</a> | 3417 | <a href="#">418aa</a> | Protein coding  | <a href="#">CCDS22184</a> | <a href="#">Q8R2R9</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 |
| Ap3m2-204 | <a href="#">ENSMUST00000210656.1</a> | 3373 | <a href="#">418aa</a> | Protein coding  | <a href="#">CCDS22184</a> | <a href="#">Q8R2R9</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 |
| Ap3m2-202 | <a href="#">ENSMUST00000210148.1</a> | 1351 | No protein            | Retained intron | -                         | -                      | TSL:1   |
| Ap3m2-203 | <a href="#">ENSMUST00000210476.1</a> | 1339 | No protein            | Retained intron | -                         | -                      | TSL:1   |

The strategy is based on the design of *Ap3m2-201* transcript,The transcription is shown below

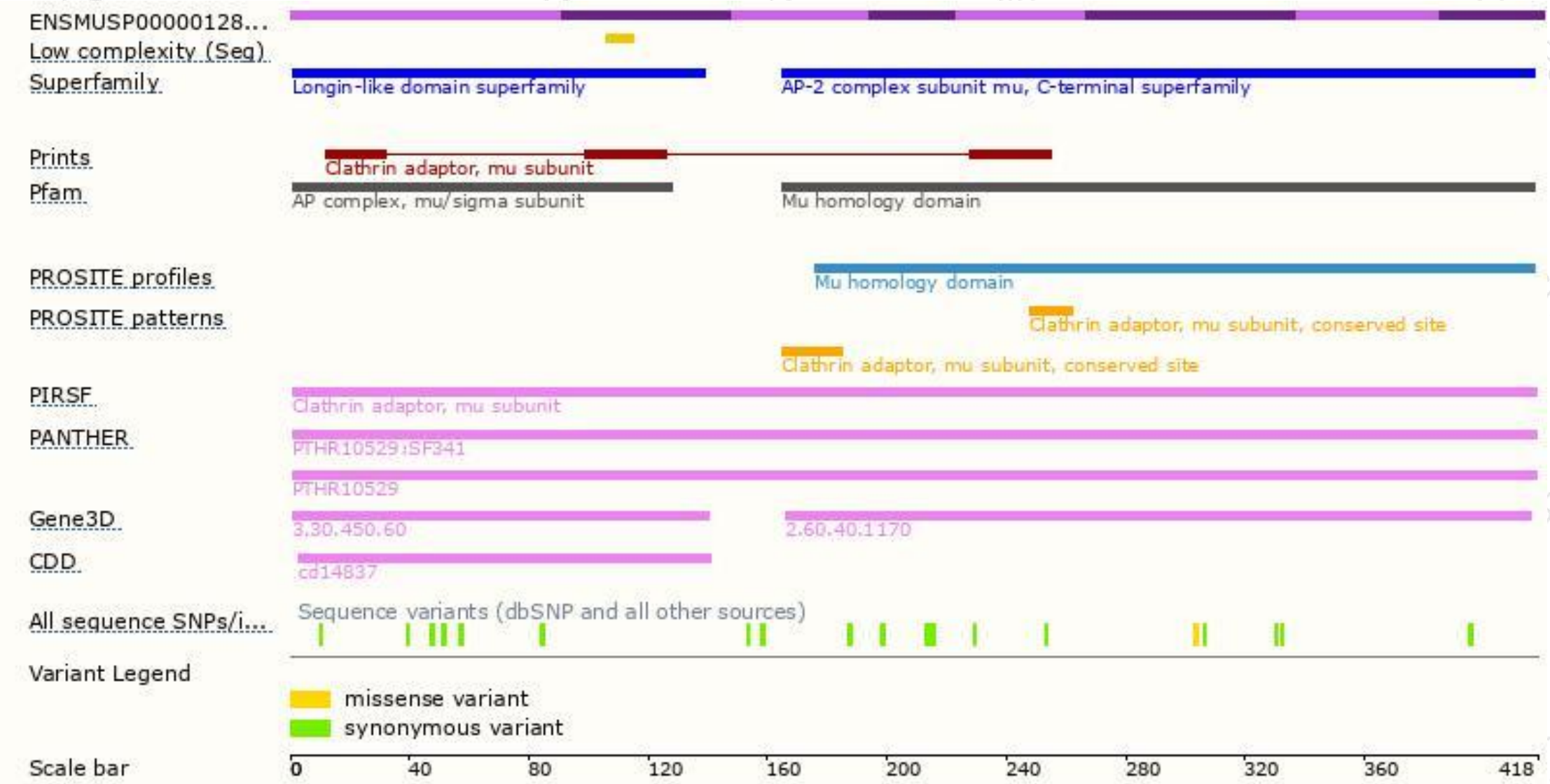


# Genomic location distribution

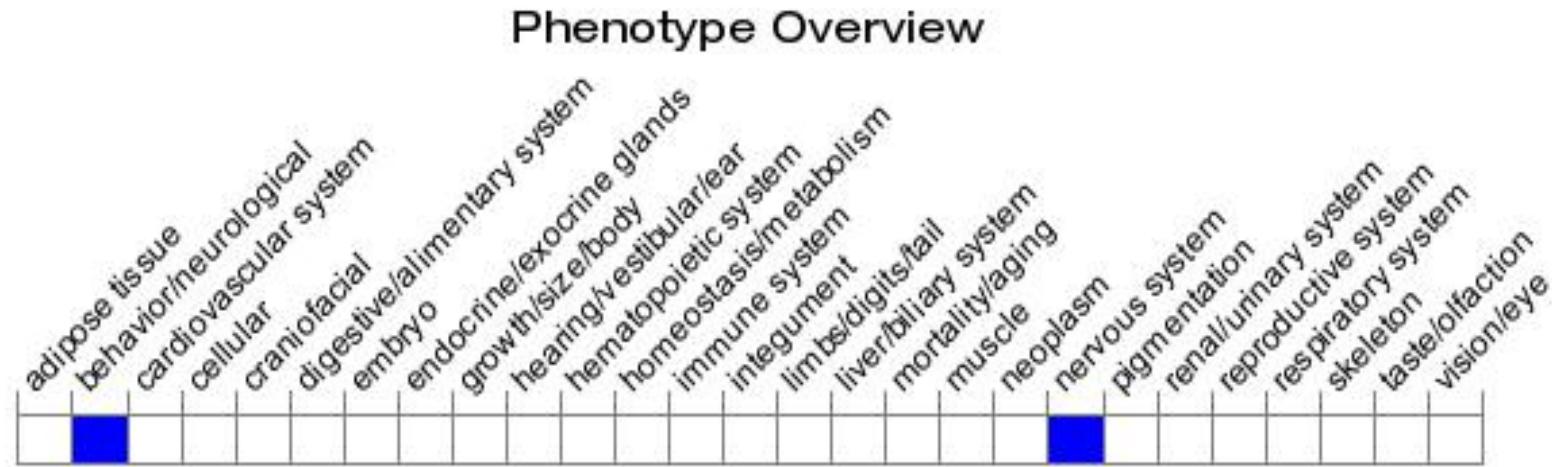




# Protein domain



# 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, Homozygous null mice suffer from spontaneous recurrent epileptic seizures, are more susceptible to drug-induced seizures and show impaired GABA release, fewer synaptic vesicles, enhanced long-term potentiation, and abnormal propagation of neuronal excitability via the temporoammonic pathway.

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

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