# *Chn1* Cas9-CKO Strategy

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**Reviewer:** Huimin Su
**Design Date:** 2020-2-27

# Project Overview

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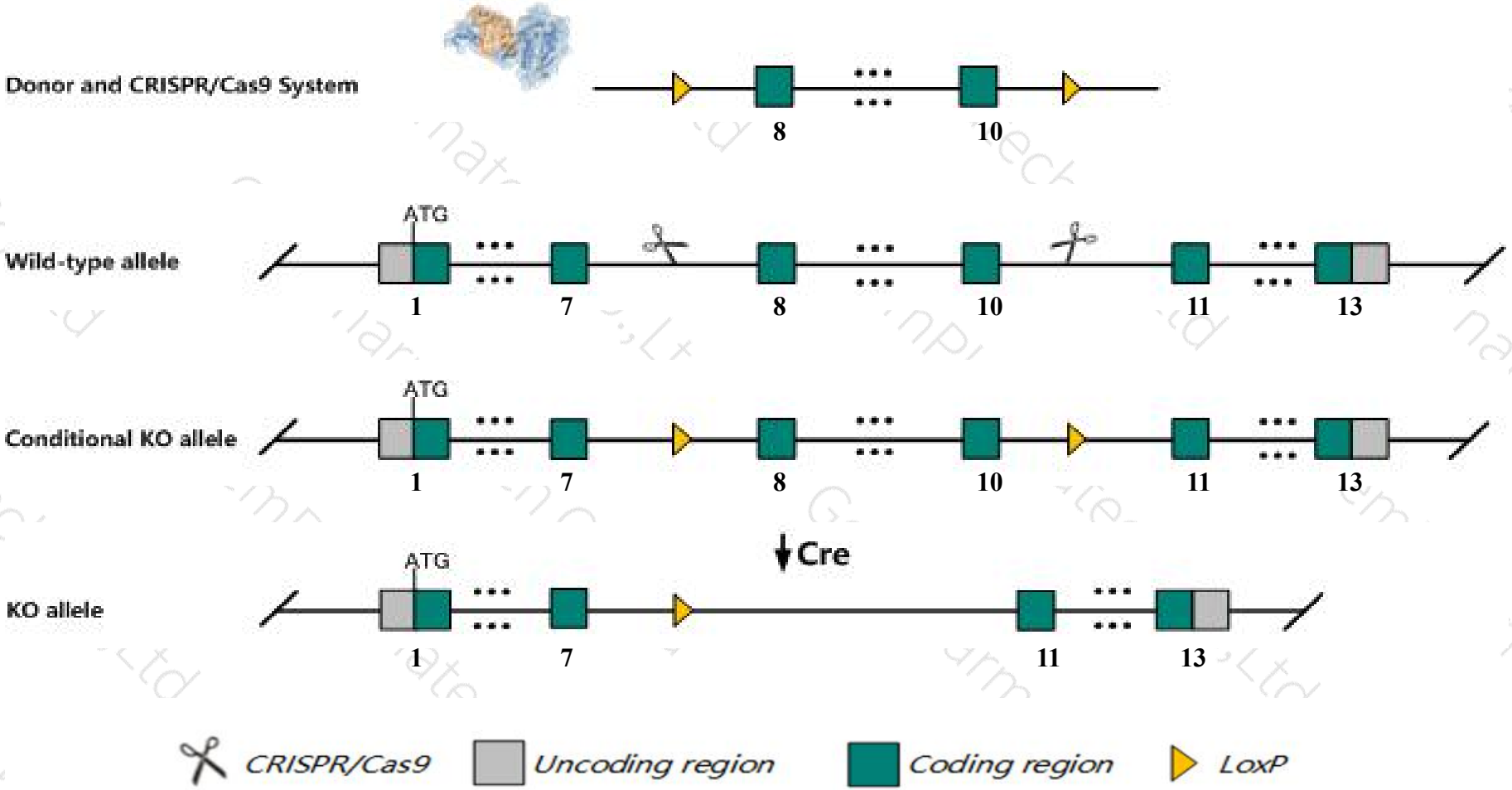
|---|---|
| **Project Name** | ***Chn1*** |
| **Project type** | **Cas9-CKO** |
| **Strain background** | **C57BL/6JGpt** |

# Conditional Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Chn1* gene. The schematic diagram is as follows:



# Technical routes

- The *Chn1* gene has 15 transcripts. According to the structure of *Chn1* gene, exon8-exon10 of *Chn1-203* (ENSMUST00000112024.9) transcript is recommended as the knockout region.The region contains 337bp coding sequence. Knock out the region will result in disruption of protein function.

- In this project we use CRISPR/Cas9 technology to modify *Chn1* 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.

# Notice

- According to the existing MGI data,mice homologous for a null allele exhibit transient postnatal size reduction, abnormal gait and abnormal innervation of the spinal cord. Part of null homozygous show preweaning lethality.

- Transcript Chn1-205 may not be affected.

- The *Chn1* gene is located on the Chr2. 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）

集萃药康 GemPharmatech

## Chn1 chimerin 1 [ *Mus musculus* (house mouse) ]

Gene ID: 108699, updated on 19-Nov-2019

### Summary

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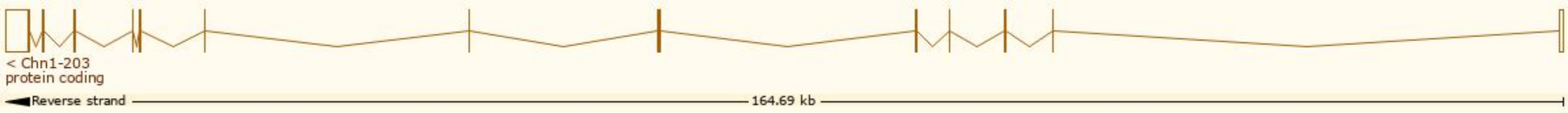
|---|---|
| **Official Symbol** | Chn1 provided by MGI |
| **Official Full Name** | chimerin 1 provided by MGI |
| **Primary source** | MGI:MGI:1915674 |
| **See related** | Ensembl:ENSMUSG00000056486 |
| **Gene type** | protein coding |
| **RefSeq status** | VALIDATED |
| **Organism** | Mus musculus |
| **Lineage** | Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus |
| **Also known as** | ARHGAP2; AI413815; 0610007I19Rik; 0710001E19Rik; 1700112L09Rik; 2900046J01Rik |
| **Expression** | Biased expression in cortex adult (RPKM 122.5), frontal lobe adult (RPKM 81.5) and 6 other tissues See more |
| **Orthologs** | human all |

# Transcript information（Ensembl）

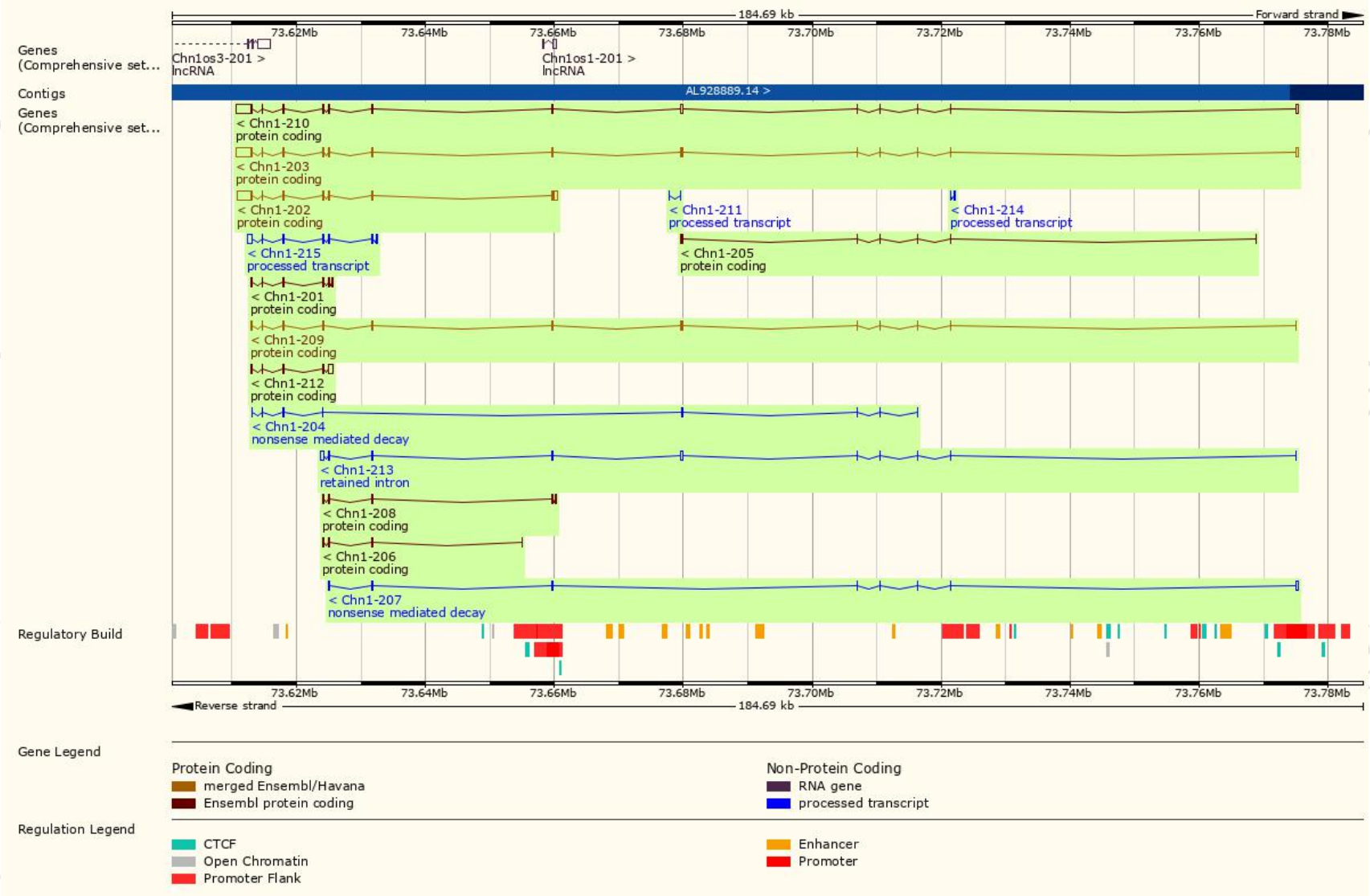
The gene has 15 transcripts,all transcripts are shown below:

| Name | Transcript ID | bp | Protein | Biotype | CCDS | UniProt | Flags |
|---|---|---|---|---|---|---|---|
| Chn1-203 | ENSMUST00000112024.9 | 4050 | 459aa | Protein coding | CCDS50608 | Q91V57 | TSL:1 GENCODE basic APPRIS P1 |
| Chn1-210 | ENSMUST00000180045.7 | 4050 | 210aa | Protein coding | CCDS38145 | A7VK14 Q91V57 | TSL:1 GENCODE basic |
| Chn1-202 | ENSMUST00000102677.10 | 3760 | 334aa | Protein coding | CCDS16133 | Q91V57 | TSL:1 GENCODE basic |
| Chn1-212 | ENSMUST00000229731.1 | 1403 | 210aa | Protein coding | CCDS38145 | A7VK14 Q91V57 | GENCODE basic |
| Chn1-209 | ENSMUST00000166199.8 | 1324 | 401aa | Protein coding | CCDS50607 | A7VK13 | TSL:1 GENCODE basic |
| Chn1-201 | ENSMUST00000070579.6 | 1023 | 210aa | Protein coding | CCDS38145 | A7VK14 Q91V57 | TSL:1 GENCODE basic |
| Chn1-208 | ENSMUST00000154258.7 | 712 | 95aa | Protein coding | - | B2FDI0 | CDS 3' incomplete TSL:5 |
| Chn1-205 | ENSMUST00000135904.1 | 631 | 128aa | Protein coding | - | B2FDI2 | CDS 3' incomplete TSL:2 |
| Chn1-206 | ENSMUST00000136953.1 | 413 | 81aa | Protein coding | - | B2FDI1 | CDS 3' incomplete TSL:3 |
| Chn1-207 | ENSMUST00000139252.1 | 881 | 87aa | Nonsense mediated decay | - | D6RCX8 | TSL:3 |
| Chn1-204 | ENSMUST00000124450.1 | 643 | 29aa | Nonsense mediated decay | - | F7C3N6 | CDS 5' incomplete TSL:3 |
| Chn1-215 | ENSMUST00000231013.1 | 1544 | No protein | Processed transcript | - | - | - |
| Chn1-214 | ENSMUST00000230959.1 | 205 | No protein | Processed transcript | - | - | - |
| Chn1-211 | ENSMUST00000229312.1 | 20 | No protein | Processed transcript | - | - | - |
| Chn1-213 | ENSMUST00000229987.1 | 1431 | No protein | Retained intron | - | - | - |

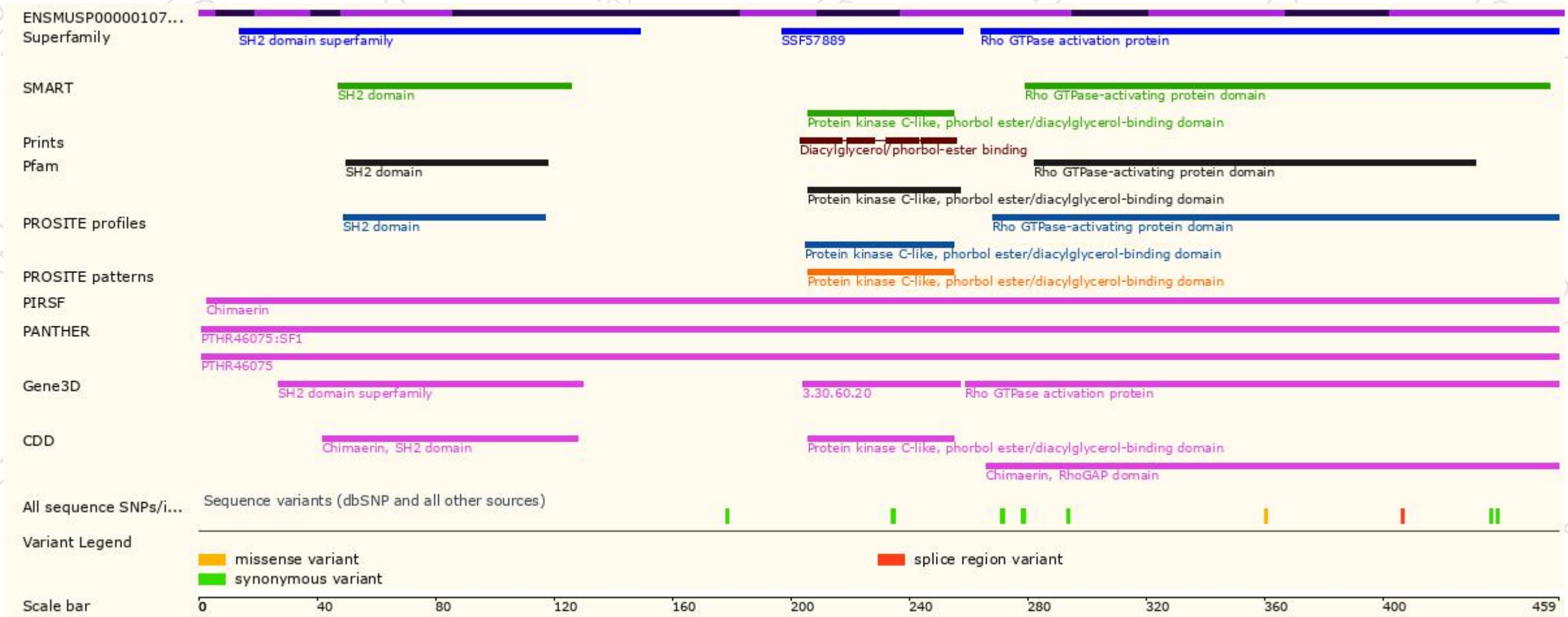
The strategy is based on the design of *Chn1-203* transcript, the transcription is shown below:



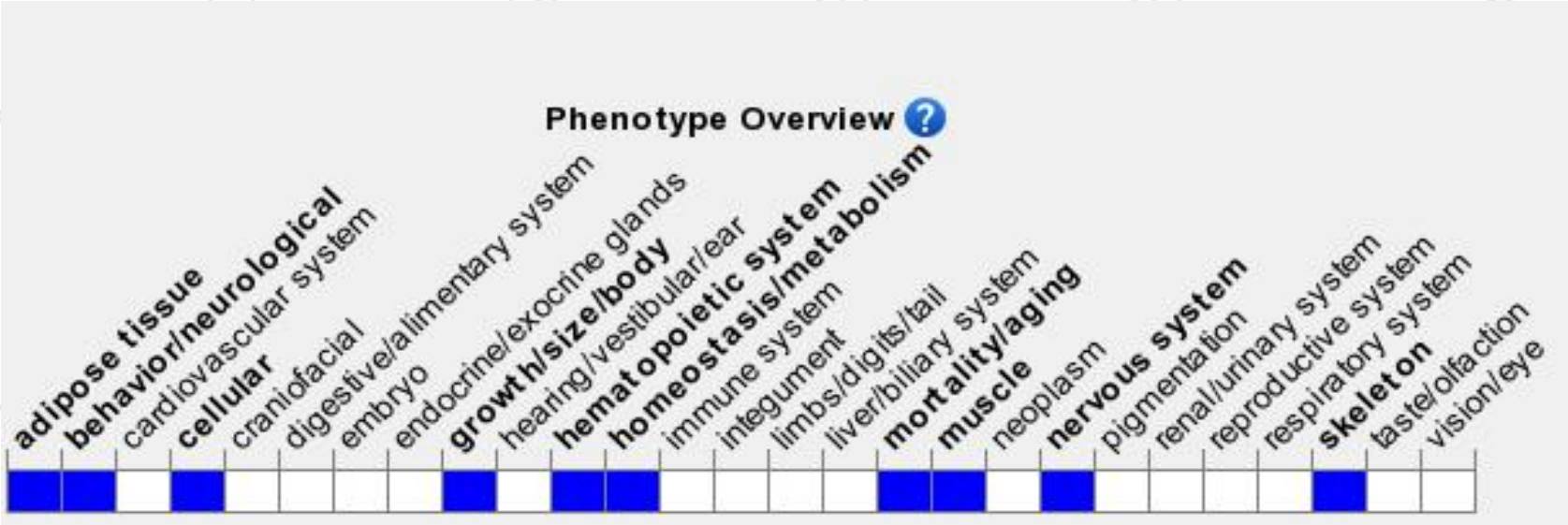
# Genomic location distribution



# 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 homologous for a null allele exhibit transient postnatal size reduction, abnormal gait and abnormal innervation of the spinal cord. Part of null homozygous show preweaning lethality.

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

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