

Snap29 Cas9-KO Strategy

Designer:Xueting Zhang

Reviewer: Yanhua Shen

Date:2019-12-02

Project Overview



Project Name

Snap29

Project type

Cas9-KO

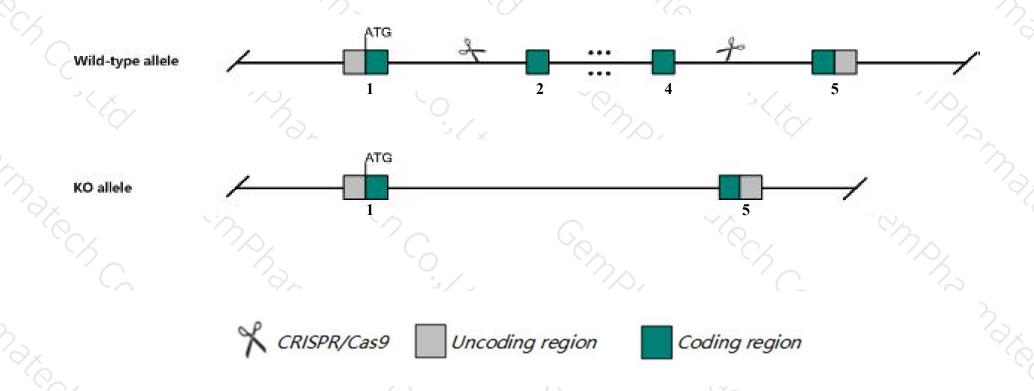
Strain background

C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Snap29 gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Snap29* gene has 1 transcript. According to the structure of *Snap29* gene, exon2-exon4 of *Snap29-201* (ENSMUST00000023449.10) transcript is recommended as the knockout region. The region contains 388bp coding sequence Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Snap29* gene. The brief process is as follows: gRNA was transcribed in vitro.Cas9 and gRNA 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.

Notice



- > According to the existing MGI data, Mice homozygous for a knock-out allele exhibit slightly reduced birth body size and a congenital ichtyotic phenotype associated with scaly and tight skin, hyperkeratosis, acanthosis, abnormalities in epidermal differentiation and autophagy, and increased endoplasmic reticulum stress.
- > The *Snap29* gene is located on the Chr16. 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)



Snap29 synaptosomal-associated protein 29 [Mus musculus (house mouse)]

Gene ID: 67474, updated on 12-Aug-2019

Summary

△ ?

Official Symbol Snap29 provided by MGI

Official Full Name synaptosomal-associated protein 29 provided by MGI

Primary source MGI:MGI:1914724

See related Ensembl: ENSMUSG00000022765

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 Gs32; Al891940; AU020222; BB131856; 1300018G05Rik

Expression Ubiquitous expression in placenta adult (RPKM 6.2), CNS E18 (RPKM 4.5) and 28 other tissues See more

Orthologs human all

Genomic context

☆ ?

Location: 16; 16 A3

See Snap29 in Genome Data Viewer

Exon count: 5

| Annotation release | Status | Assembly | Chr | Location | |
|--------------------|-------------------|------------------------------|-----|--------------------------------|--|
| 108 | current | GRCm38.p6 (GCF_000001635.26) | 16 | NC_000082.6 (1740600017430826) | |
| Build 37.2 | previous assembly | MGSCv37 (GCF_000001635.18) | 16 | NC_000082.5 (1740609317430919) | |

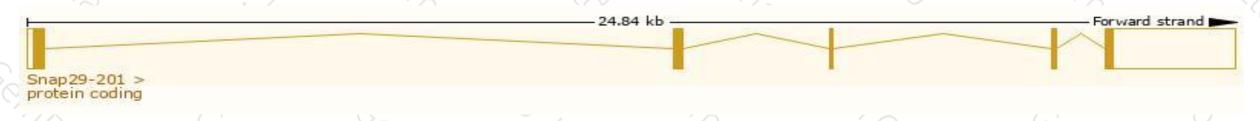
Transcript information (Ensembl)



The gene has 1 transcript, and the transcript is shown below:

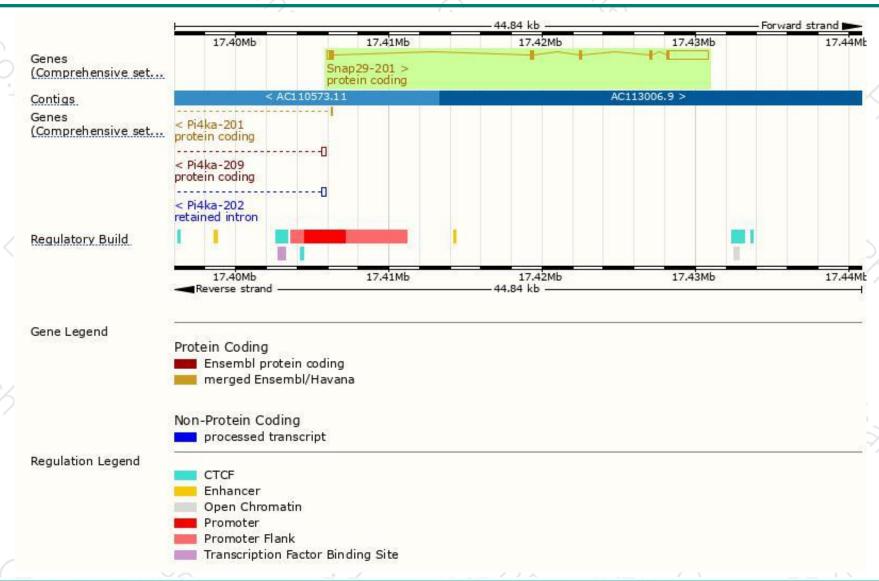
| Name | Transcript ID | bp | Protein | Biotype | CCDS | UniProt | Flags |
|------------|-----------------------|------|--------------|----------------|-----------|---------|-------------------------------|
| Snap29-201 | ENSMUST00000023449.10 | 3447 | <u>260aa</u> | Protein coding | CCDS28001 | Q9ERB0 | TSL:1 GENCODE basic APPRIS P1 |

The strategy is based on the design of Snap29-201 transcript, The transcription is shown below



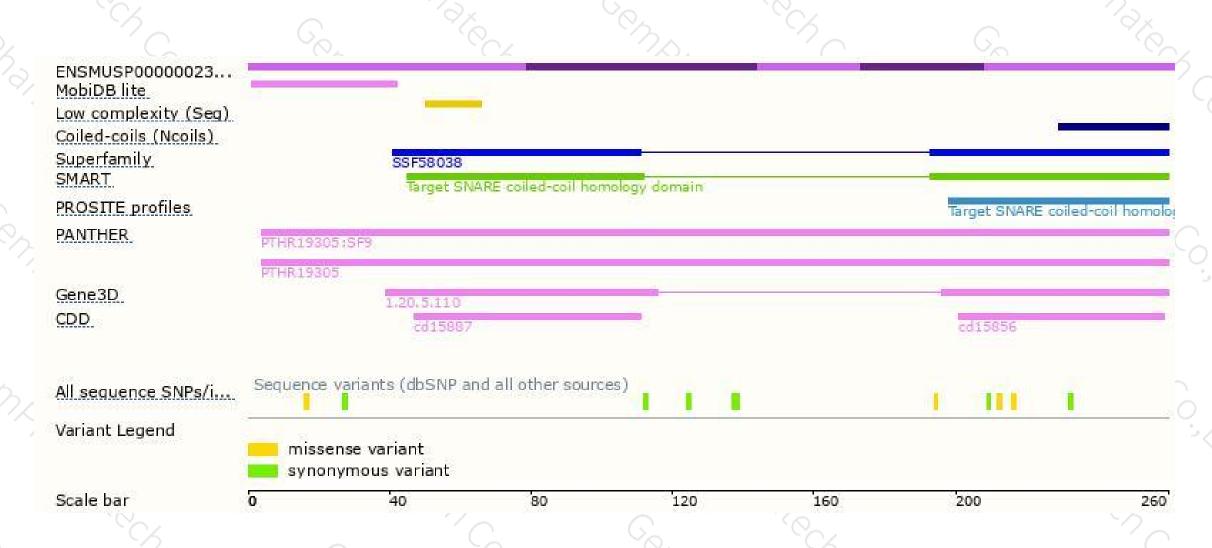
Genomic location distribution





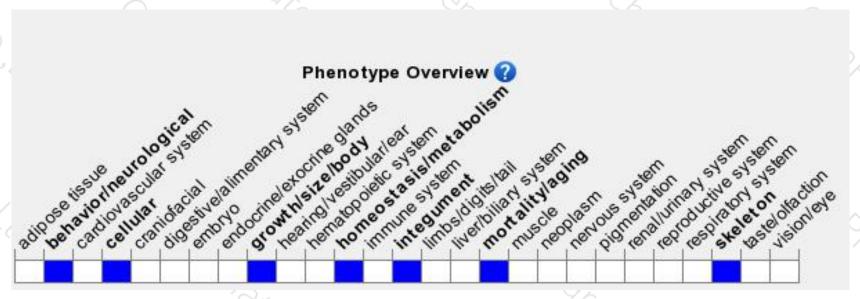
Protein domain





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, Mice homozygous for a knock-out allele exhibit slightly reduced birth body size and a congenital ichtyotic phenotype associated with scaly and tight skin, hyperkeratosis, acanthosis, abnormalities in epidermal differentiation and autophagy, and increased endoplasmic reticulum stress.



If you have any questions, you are welcome to inquire. Tel: 400-9660890





