

Slc44a2 Cas9-CKO Strategy

Designer:Xueting Zhang

Reviwer: Yanhua Shen

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Project Overview



Project Name

Slc44a2

Project type

Cas9-CKO

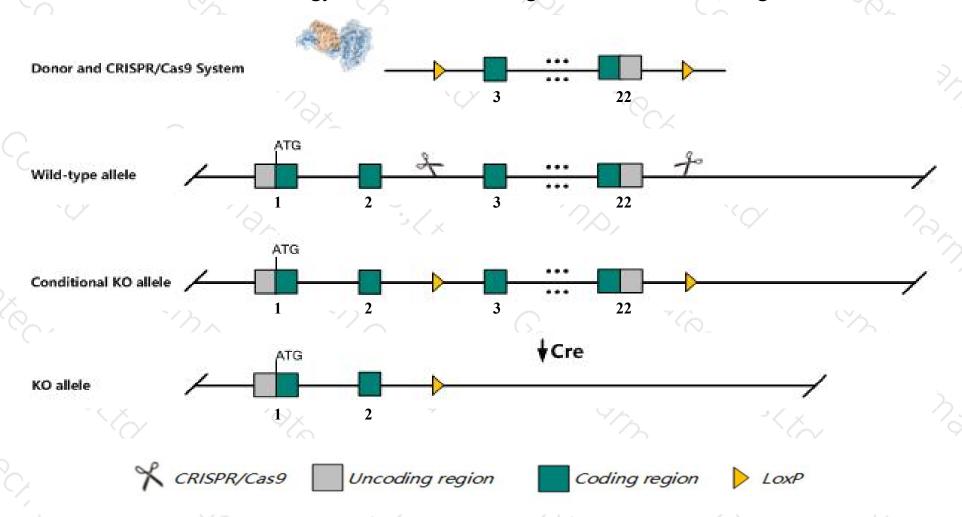
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Slc44a2 gene. The schematic diagram is as follows:



Technical routes



- The *Slc44a2* gene has 10 transcripts. According to the structure of *Slc44a2* gene, exon3-exon22 of *Slc44a2-201* (ENSMUST00000034697.7) transcript is recommended as the knockout region. The region contains most of the coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Slc44a2* 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 homozygous for a knock-out allele exhibit cochlear hair cell loss, spiral ganglion degeneration, and progressive sensorineural hearing loss.
- The *Slc44a2* gene is located on the Chr9. 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)



Slc44a2 solute carrier family 44, member 2 [Mus musculus (house mouse)]

Gene ID: 68682, updated on 18-Feb-2020

Summary

☆ ?

Official Symbol Slc44a2 provided by MGI

Official Full Name solute carrier family 44, member 2 provided by MGI

Primary source MGI:MGI:1915932

See related Ensembl: ENSMUSG00000057193

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 CTL2; 1110028E10Rik

Expression Broad expression in thymus adult (RPKM 112.2), lung adult (RPKM 80.3) and 23 other tissues See more

Orthologs <u>human</u> all

Genomic context



Location: 9; 9 A3

See Slc44a2 in Genome Data Viewer

Exon count: 25

| Annotation release | Status | Assembly | Chr | Location | |
|--------------------|-------------------|------------------------------|-----|--------------------------------|--|
| 108 | current | GRCm38.p6 (GCF_000001635.26) | 9 | NC_000075.6 (2132071921355028) | |
| Build 37.2 | previous assembly | MGSCv37 (GCF_000001635.18) | 9 | NC_000075.5 (2114229621159472) | |

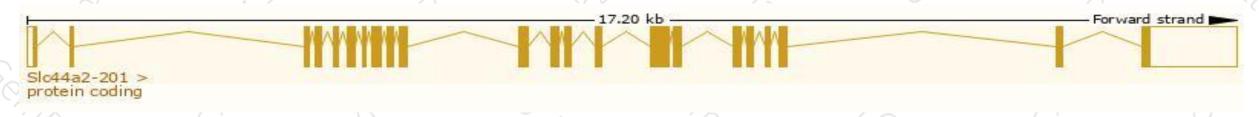
Transcript information (Ensembl)



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

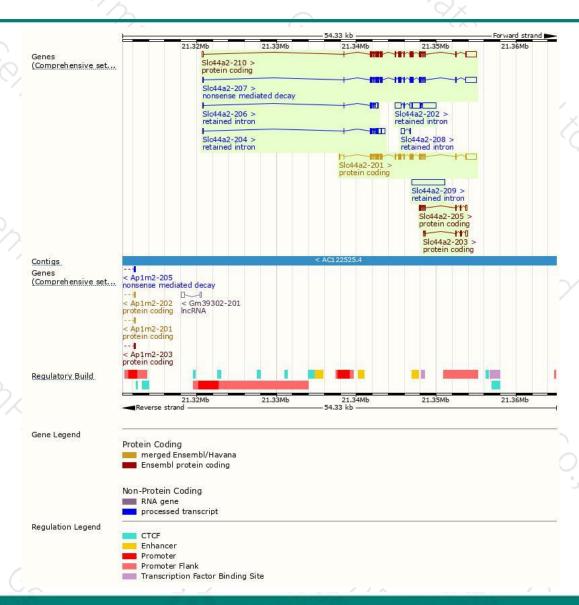
| Name | Transcript ID | bp | Protein | Biotype | CCDS | UniProt | Flags |
|-------------|----------------------|------|--------------|-------------------------|-----------------|------------|---------------------------------|
| SIc44a2-201 | ENSMUST00000034697.7 | 3456 | 706aa | Protein coding | CCDS22903 | Q8BY89 | TSL:1 GENCODE basic APPRIS P2 |
| SIc44a2-210 | ENSMUST00000217461.1 | 3478 | 704aa | Protein coding | 19 5 | Q8BY89 | TSL:1 GENCODE basic APPRIS ALT1 |
| SIc44a2-205 | ENSMUST00000214268.1 | 684 | <u>170aa</u> | Protein coding | <u> </u> | A0A1L1SU40 | CDS 5' incomplete TSL:5 |
| SIc44a2-203 | ENSMUST00000213535.1 | 679 | <u>111aa</u> | Protein coding | 12 | A0A1L1SS87 | CDS 5' incomplete TSL:3 |
| SIc44a2-207 | ENSMUST00000215574.1 | 3428 | <u>554aa</u> | Nonsense mediated decay | | A0A1L1SVG6 | TSL:1 |
| SIc44a2-209 | ENSMUST00000217453.1 | 4077 | No protein | Retained intron | 19- | * | TSL:NA |
| SIc44a2-202 | ENSMUST00000213499.1 | 3744 | No protein | Retained intron | <u> </u> | 24 | TSL:2 |
| SIc44a2-204 | ENSMUST00000213758.1 | 1315 | No protein | Retained intron | 2 | <u>(2)</u> | TSL:1 |
| SIc44a2-206 | ENSMUST00000215528.1 | 713 | No protein | Retained intron | | | TSL:5 |
| SIc44a2-208 | ENSMUST00000216266.1 | 515 | No protein | Retained intron | 19- | -8 | TSL:1 |
| | | | 7 | | | | |

The strategy is based on the design of Slc44a2-201 transcript, The transcription is shown below



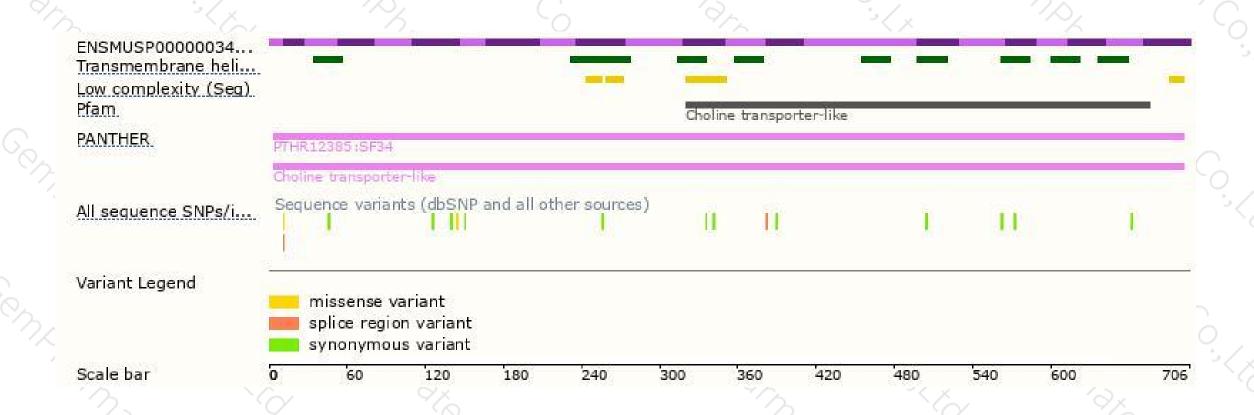
Genomic location distribution





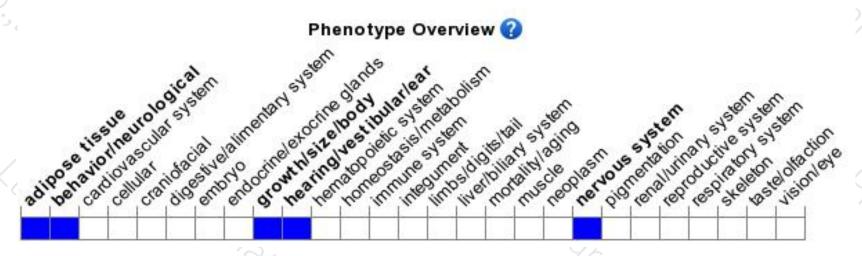
Protein domain





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 cochlear hair cell loss, spiral ganglion degeneration, and progressive sensorineural hearing loss.



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





