

Slc39a7 Cas9-KO Strategy

Designer:Xueting Zhang

Project Overview



Project Name

Slc39a7

Project type

Cas9-KO

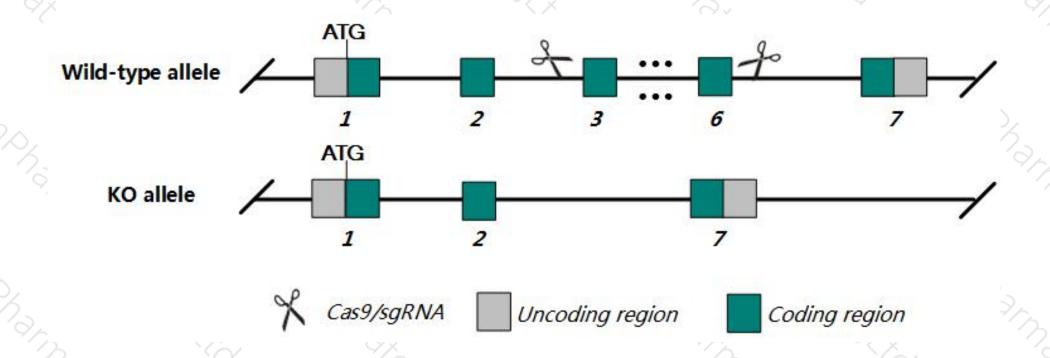
Strain background

C57BL/6J

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Slc39a7 gene. The schematic diagram is as follows:



Technical routes



- ➤ The Slc39a7 gene has 6 transcripts. According to the structure of Slc39a7 gene, exon3-exon6 of Slc39a7-201 (ENSMUST00000025186.15) transcript is recommended as the knockout region. The region contains 617bp coding sequence Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Slc39a7* gene. The brief process is as follows: sgRNA was transcribed in vitro.Cas9 and sgRNA were microinjected into the fertilized eggs of C57BL/6J 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/6J mice.

Notice



- > According to the existing MGI data, Mice homozygous for a conditional allele activated in intestinal epithelial cells exhibit premature death, loss of epithelial integrity, reduced enterocyte proliferation and increased enterocyte apoptosis.
- The knpckout region is near to the N-terminal of $Rxrb \Col11a2 \H2-ke6 \Mir219a \Gm26940 \Ring1$ gene, this strategy may influence the regulatory function of the N-terminal of these gene.
- ightharpoonup The Gm20427 gene will be deleted together in this strategy.
- ➤ The *Slc39a7* gene is located on the Chr11. 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 gene transcription and translation processes, all risks cannot be predicted under existing information.

Gene information (NCBI)



SIc39a7 solute carrier family 39 (zinc transporter), member 7 [Mus musculus (house mouse)]

Gene ID: 14977, updated on 15-Apr-2019

Summary

☆ ?

Official Symbol Slc39a7 provided by MGI

Official Full Name solute carrier family 39 (zinc transporter), member 7 provided by MGI

Primary source MGI:MGI:95909

See related Ensembl: ENSMUSG00000024327

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 Ke4; Ke-4; Zip7; Ring5; H-2Ke4; H2-Ke4; AA408174; Al117660; AL024048

Expression Ubiquitous expression in ovary adult (RPKM 99.5), adrenal adult (RPKM 66.7) and 28 other tissues See more

Orthologs human all

Genomic context



Location: 17 B1; 17 17.98 cM

See Slc39a7 in Genome Data Viewer

Exon count: 7

Annotation release	Status	Assembly	Chr	Location	
106	current	GRCm38.p4 (GCF_000001635.24)	17	NC_000083.6 (3402826634031690, complement)	
Build 37.2	previous assembly	MGSCv37 (GCF_000001635.18)	17	NC_000083.5 (3416521134168635, complement)	

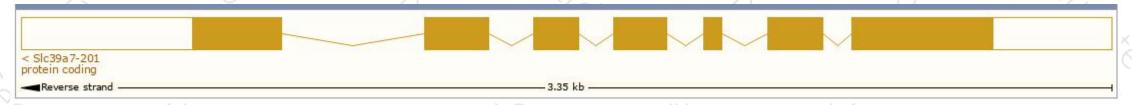
Transcript information (Ensembl)



The gene has 6 transcripts, all transcripts are shown below:

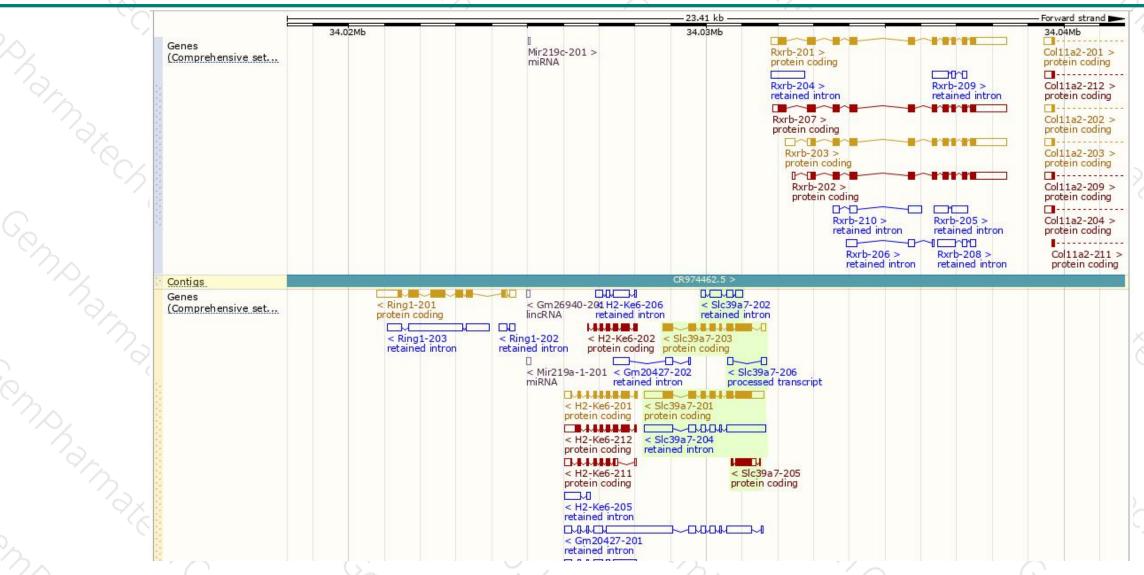
Name	Transcript ID A	bp 🌲	Protein	Biotype	CCDS 🍦	UniProt	Flags
Slc39a7-201	ENSMUST00000025186.15	2320	476aa	Protein coding	CCDS37576@	A0A068BIT0 @ Q31125 @	TSL:1 GENCODE basic APPRIS P1
Slc39a7-202	ENSMUST00000167145.1	844	No protein	Retained intron	-	-	TSL:5
Slc39a7-203	ENSMUST00000169397.8	1550	476aa	Protein coding	CCDS37576@	A0A068BIT0@ Q31125@	TSL:5 GENCODE basic APPRIS P1
Slc39a7-204	ENSMUST00000170491.7	2433	No protein	Retained intron	-		TSL:1
Slc39a7-205	ENSMUST00000171872.2	620	<u>161aa</u>	Protein coding	-	<u>E9PV10</u> ₽	CDS 3' incomplete TSL:5
Slc39a7-206	ENSMUST00000173810.1	277	No protein	Processed transcript	-	14	TSL:5

The strategy is based on the design of Slc39a7-201 transcript, The transcription is shown below



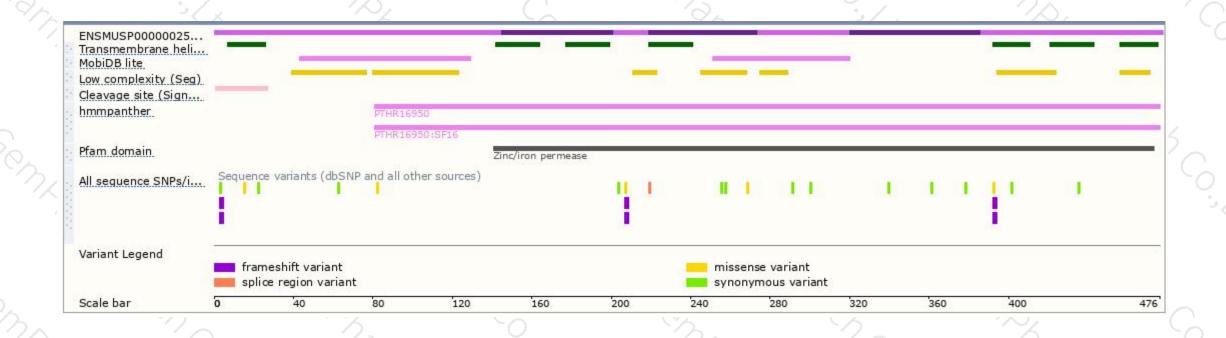
Genomic location distribution





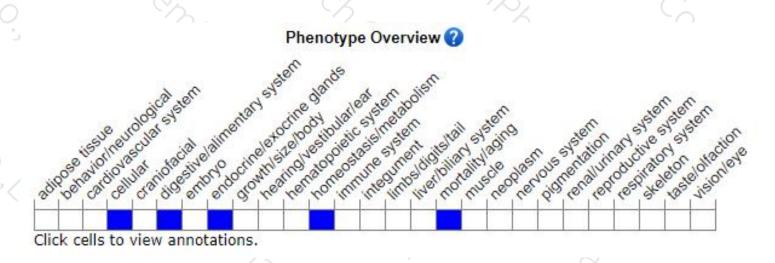
Protein domain





Mouse phenotype description(MGI)





Phenotypes affected by the gene are marked in blue.Data quoted from MGI database(http://www.informatics.jax.org/).

Mice homozygous for a conditional allele activated in intestinal epithelial cells exhibit premature death, loss of epithelial integrity, reduced enterocyte proliferation and increased enterocyte apoptosis.



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

Tel: 025-5864 1534





