

Slc39a5 Cas9-CKO Strategy

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Design Date: 2020-2-19

Project Overview



Project Name

Slc39a5

Project type

Cas9-CKO

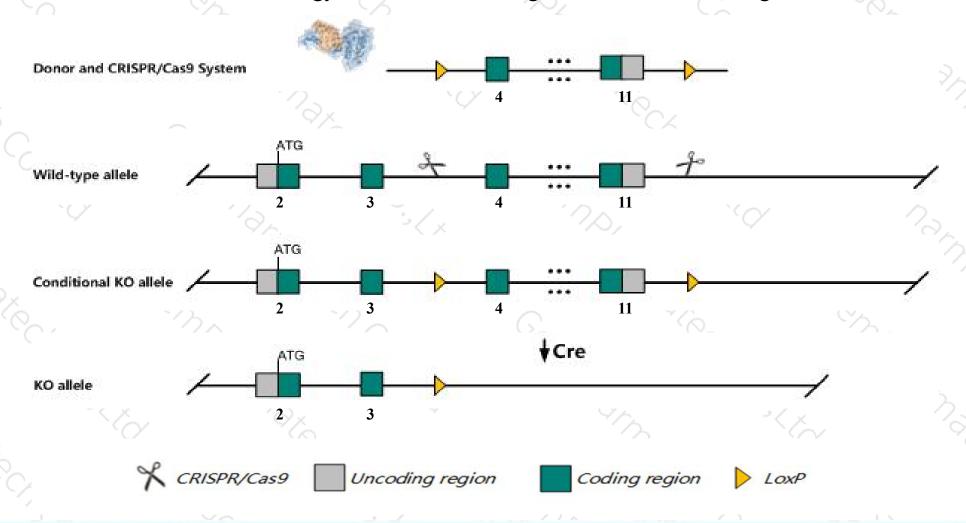
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Slc39a5 gene. The schematic diagram is as follows:



Technical routes



- The *Slc39a5* gene has 5 transcripts. According to the structure of *Slc39a5* gene, exon4-exon11 of *Slc39a5-202* (ENSMUST00000167859.1) 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 *Slc39a5* 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 altered zinc homeostasis and increased susceptibility to zinc-induced pancretitis.
- > The function of Gm17201 may be affected.
- The *Slc39a5* gene is located on the Chr10. 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)



△ ?

Slc39a5 solute carrier family 39 (metal ion transporter), member 5 [Mus musculus (house mouse)]

Gene ID: 72002, updated on 12-Aug-2019

Summary

Official Full Name solute carrier family 39 (metal ion transporter), member 5 provided by MGI

Primary source MGI:MGI:1919336

Official Symbol Slc39a5 provided by MGI

See related Ensembl: ENSMUSG00000039878

Gene type protein coding RefSeg 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 Zip5; 1810013D05Rik; 2010205A06Rik

Expression Biased expression in duodenum adult (RPKM 158.7), large intestine adult (RPKM 83.5) and 7 other tissues See more

Orthologs human all

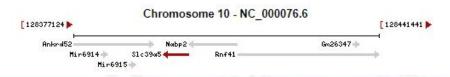
Location: 10: 10 D3

Exon count: 12

	☆ ?
nomic context	

See Slc39a5 in Genome Data Viewer

Annotation release	Status	Assembly	Chr	Location	
108	current	GRCm38.p6 (GCF_000001635.26)	10	NC_000076.6 (128395927128401276, complement)	
Build 37.2	previous assembly	MGSCv37 (GCF_000001635.18)	10	NC_000076.5 (127832987127838280, complement))



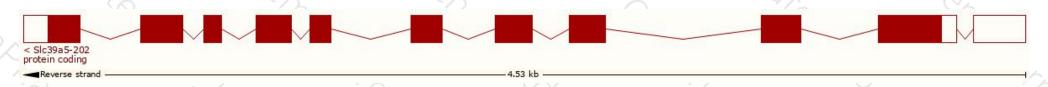
Transcript information (Ensembl)



The gene has 5 transcripts, all transcripts are shown below:

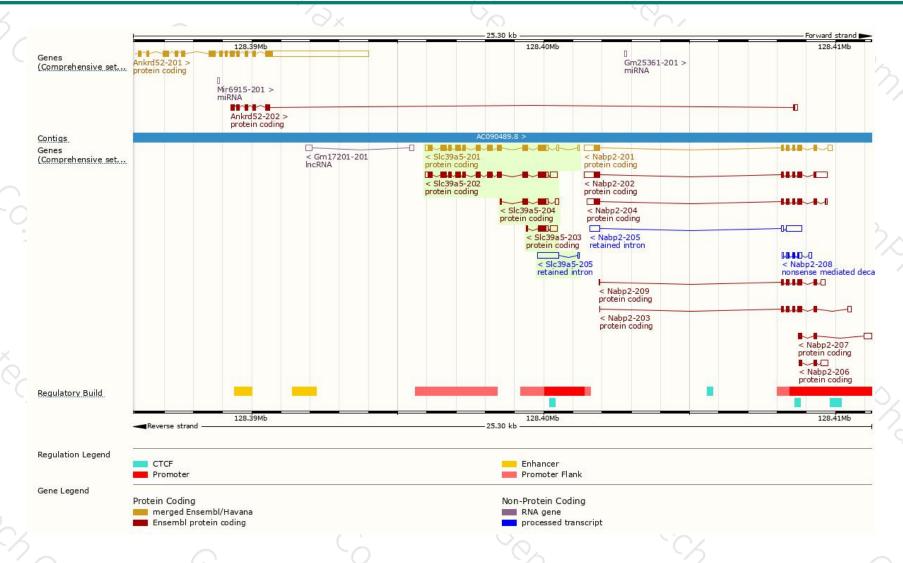
Name 🍦	Transcript ID	bp 👙	Protein #	Biotype	CCDS 🍦	UniProt #	Flags		
Slc39a5-202	ENSMUST00000167859.1	2025	<u>535aa</u>	Protein coding	CCDS24275₽	Q9D856₽	TSL:1	GENCODE basic	APPRIS P1
Slc39a5-201	ENSMUST00000042666.12	1947	<u>535aa</u>	Protein coding	CCDS24275₽	Q9D856┏	TSL:1	GENCODE basic	APPRIS P1
Slc39a5-204	ENSMUST00000219131.1	709	<u>176aa</u>	Protein coding	7.0	A0A1W2P7R0₽		CDS 3' incomplete	TSL:5
Slc39a5-203	ENSMUST00000218858.1	648	<u>114aa</u>	Protein coding	7.	A0A1W2P7Q2₽	1	CDS 3' incomplete	TSL:5
Slc39a5-205	ENSMUST00000219222.1	766	No protein	Retained intron		17.		TSL:3	

The strategy is based on the design of Slc39a5-202 transcript, The transcription is shown below



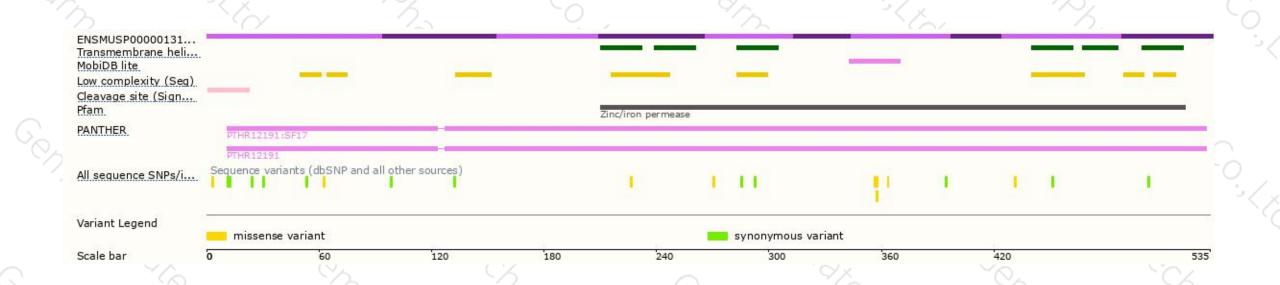
Genomic location distribution





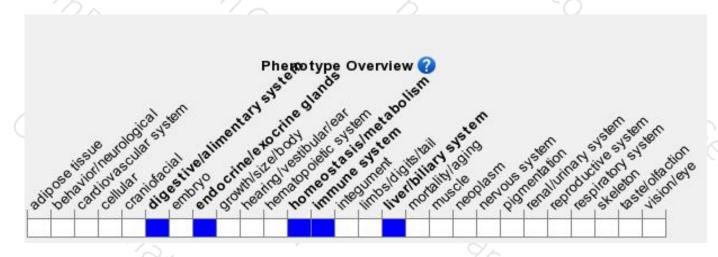
Protein domain





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 altered zinc homeostasis and increased susceptibility to zinc-induced pancretitis.



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





