

Slc11a1 Cas9-KO Strategy

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Project Overview



Project Name

Slc11a1

Project type

Cas9-KO

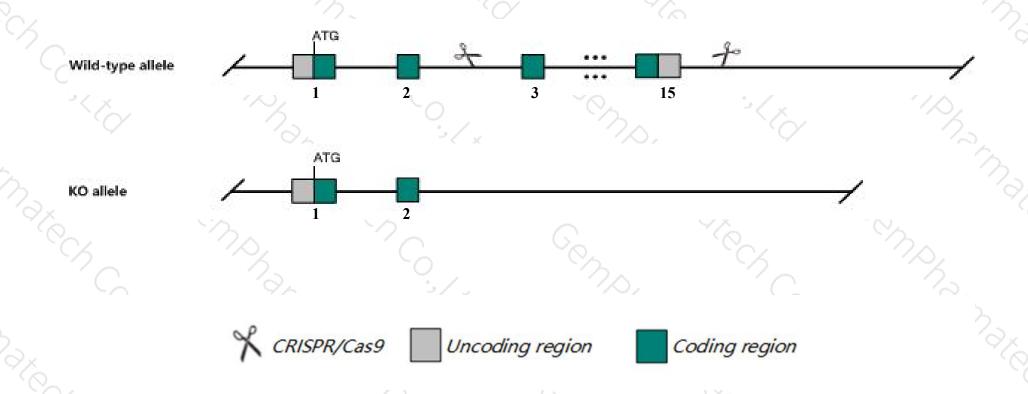
Strain background

C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Slc11a1 gene. The schematic diagram is as follows:



Technical routes



- ➤ The Slc11a1 gene has 11 transcripts. According to the structure of Slc11a1 gene, exon3-exon15 of Slc11a1-201 (ENSMUST00000027368.5) 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 Slc11a1 gene. The brief process is as follows: CRISPR/Cas9 system

Notice



- ➤ According to the existing MGI data, Mutations in this gene are associated with susceptibility to infection with pathogens, including Myobacterium, Salmonella and Leishmania. Depending on the mutation, mutants may exhibit either increased or decreased susceptibility to infection.
- > The Slc11a1 gene is located on the Chr1. 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)



SIc11a1 solute carrier family 11 (proton-coupled divalent metal ion transporters), member 1 [Mus musculus (house mouse)]

Gene ID: 18173, updated on 12-Aug-2019

Summary

☆ ?

Official Symbol Slc11a1 provided by MGI

Official Full Name solute carrier family 11 (proton-coupled divalent metal ion transporters), member 1 provided by MGI

Primary source MGI:MGI:1345275

See related Ensembl: ENSMUSG00000026177

Gene type protein coding
RefSeq status VALIDATED
Organism <u>Mus musculus</u>

Lineage Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha;

Muroidea; Muridae; Murinae; Mus; Mus

Also known as Bcg; Ity; Lsh; ity; Ity1; Nramp; Nramp1

Expression Broad expression in spleen adult (RPKM 37.5), mammary gland adult (RPKM 13.7) and 19 other tissues See more

Orthologs <u>human</u> all

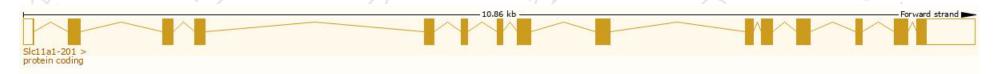
Transcript information (Ensembl)



The gene has 11 transcripts, all transcripts are shown below:

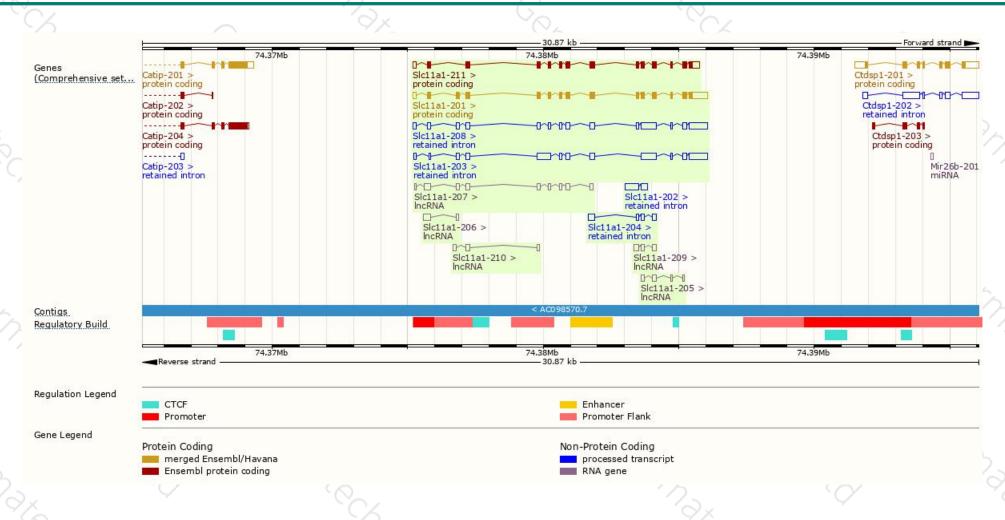
Name 🎂	Transcript ID .	bp 🌲	Protein 🌢	Biotype	CCDS 🍦	UniProt 👙	Flags
Slc11a1-201	ENSMUST00000027368.5	2315	<u>548aa</u>	Protein coding	CCDS15047 ₽	P41251₽	TSL:1 GENCODE basic APPRIS P2
Slc11a1-211	ENSMUST00000187516.6	1893	<u>507aa</u>	Protein coding	2	A0A087WNR0 ₽	TSL:5 GENCODE basic APPRIS ALT2
Slc11a1-203	ENSMUST00000131511.7	2726	No protein	Retained intron	2	2	TSL:2
Slc11a1-208	ENSMUST00000147233.7	2584	No protein	Retained intron	<u>-</u>	2	TSL:2
Slc11a1-202	ENSMUST00000127487.1	704	No protein	Retained intron	2	2	TSL:3
Slc11a1-204	ENSMUST00000133896.7	604	No protein	Retained intron	2	_	TSL:5
Slc11a1-207	ENSMUST00000136786.7	1055	No protein	IncRNA	2	_	TSL:5
Slc11a1-209	ENSMUST00000149166.7	457	No protein	I IncRNA	2	_	TSL:2
Slc11a1-205	ENSMUST00000134189.1	426	No protein	IncRNA	2	2	TSL:5
Slc11a1-206	ENSMUST00000135137.1	378	No protein	IncRNA	2	2	TSL:2
Slc11a1-210	ENSMUST00000155865.1	331	No protein	IncRNA	2	2	TSL:3

The strategy is based on the design of Slc11a1-201 transcript, The transcription is shown below



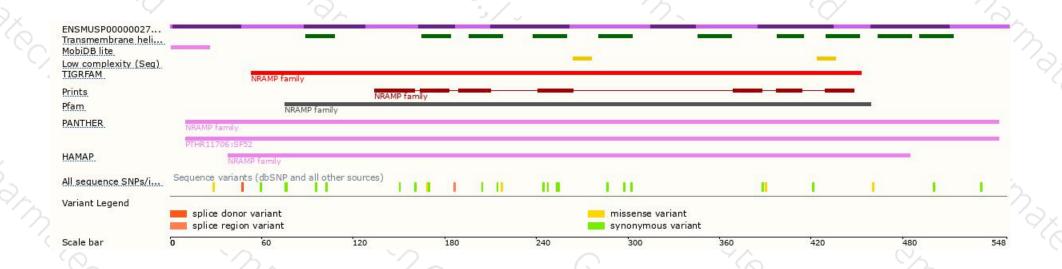
Genomic location distribution





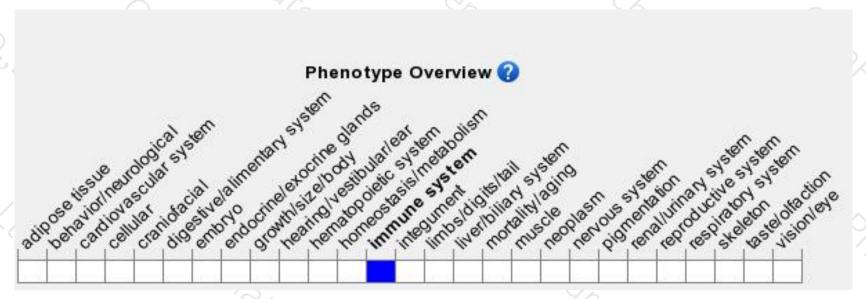
Protein domain





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, Mutations in this gene are associated with susceptibility to infection with pathogens, including Myobacterium, Salmonella and Leishmania. Depending on the mutation, mutants may exhibit either incordecreased susceptibility to infection.



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





