

# Slc4a5 Cas9-CKO Strategy

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# **Project Overview**



**Project Name** 

Slc4a5

**Project type** 

Cas9-CKO

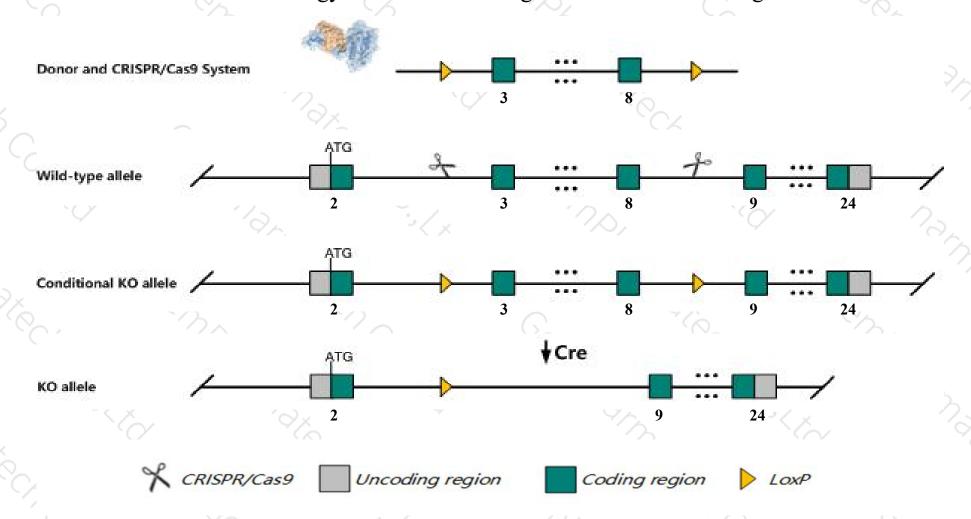
Strain background

C57BL/6JGpt

## Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Slc4a5 gene. The schematic diagram is as follows:



### Technical routes



- ➤ The *Slc4a5* gene has 6 transcripts. According to the structure of *Slc4a5* gene, exon3-exon8 of *Slc4a5-201* (ENSMUST00000039212.7) transcript is recommended as the knockout region. The region contains 712bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Slc4a5* 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 arterial hypertension and renal metabolic acidosis.
- > Transcript *Slc4a5-*205&206 may not be affected.
- The *Slc4a5* gene is located on the Chr6. 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)



#### Slc4a5 solute carrier family 4, sodium bicarbonate cotransporter, member 5 [ Mus musculus (house mouse) ]

Gene ID: 232156, updated on 5-Jan-2020

#### Summary

△ ?

Official Symbol Slc4a5 provided by MGI

Official Full Name solute carrier family 4, sodium bicarbonate cotransporter, member 5 provided by MGI

Primary source MGI:MGI:2443220

See related Ensembl: ENSMUSG00000068323

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 C330006M16; C330016K18Rik

**Expression** Biased expression in lung adult (RPKM 8.8), duodenum adult (RPKM 3.6) and 10 other tissues See more

Orthologs human all

#### Genomic context



Location: 6; 6 C3

See Slc4a5 in Genome Data Viewer

Exon count: 31

Annotation release	Status	Assembly	Chr	Location
108	current	GRCm38.p6 (GCF_000001635.26)	6	NC_000072.6 (8320758083304945)
Build 37.2	previous assembly	MGSCv37 (GCF_000001635.18)	6	NC_000072.5 (8318736983254939)

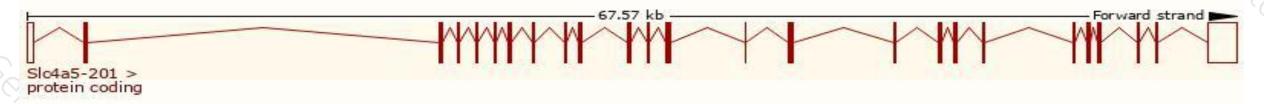
# Transcript information (Ensembl)



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

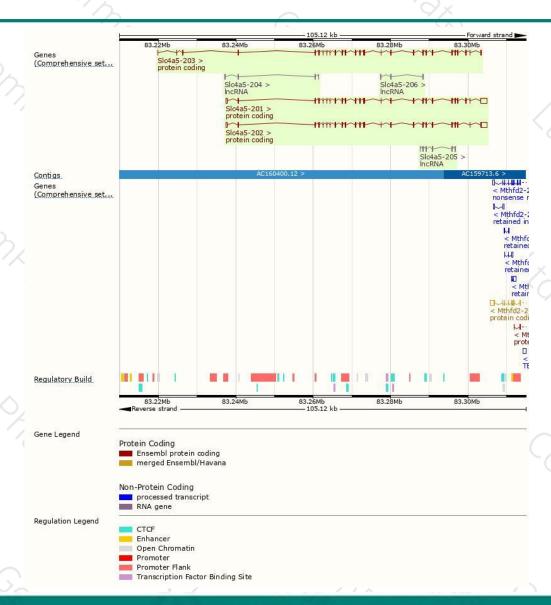
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
SIc4a5-201	ENSMUST00000039212.7	5123	<u>1001aa</u>	Protein coding	CCDS51819	D3YVG0	TSL:5 GENCODE basic APPRIS P2
SIc4a5-202	ENSMUST00000113899.7	5039	<u>1041aa</u>	Protein coding	99.	D3YX36	TSL:5 GENCODE basic APPRIS ALT2
SIc4a5-203	ENSMUST00000113900.7	3630	<u>1116aa</u>	Protein coding	120	E9Q3M5	TSL:5 GENCODE basic APPRIS ALT2
SIc4a5-205	ENSMUST00000131432.1	751	No protein	IncRNA	757	-	TSL:3
SIc4a5-206	ENSMUST00000141863.7	542	No protein	IncRNA	127	-	TSL:3
SIc4a5-204	ENSMUST00000122897.1	437	No protein	IncRNA	363	-	TSL:5

The strategy is based on the design of *Slc4a5-201* transcript, The transcription is shown below



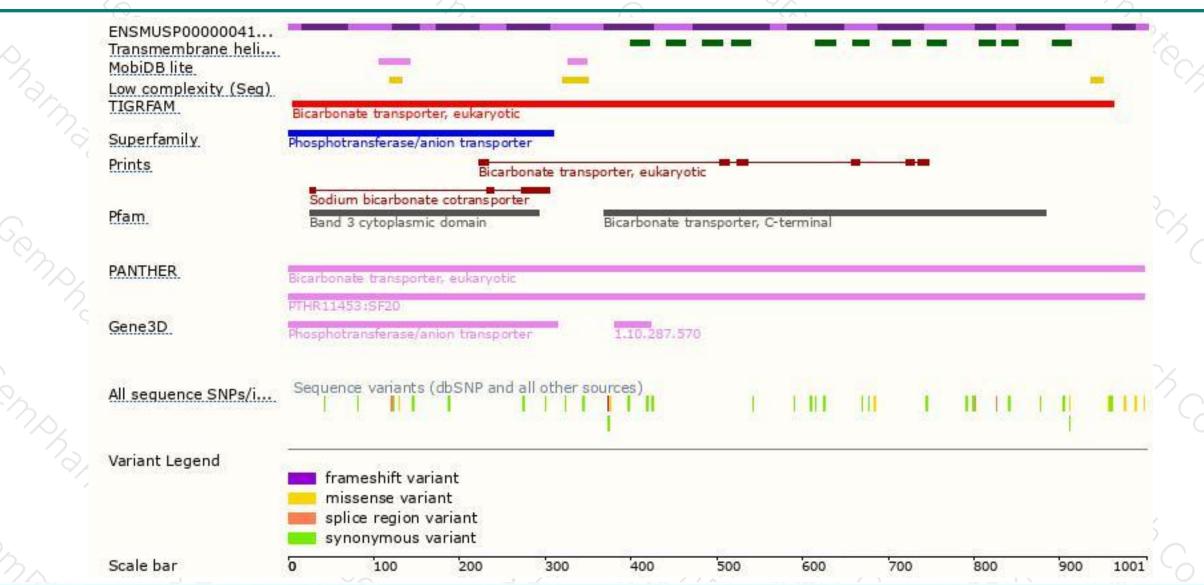
### Genomic location distribution





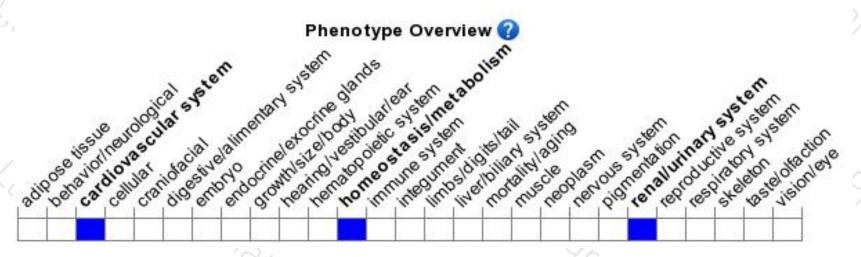
### Protein domain





# 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 arterial hypertension and renal metabolic acidosis.



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





