

# Slc22a17 Cas9-CKO Strategy

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#### Overview

#### Target Gene Name

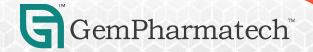
• Slc22a17

#### Project Type

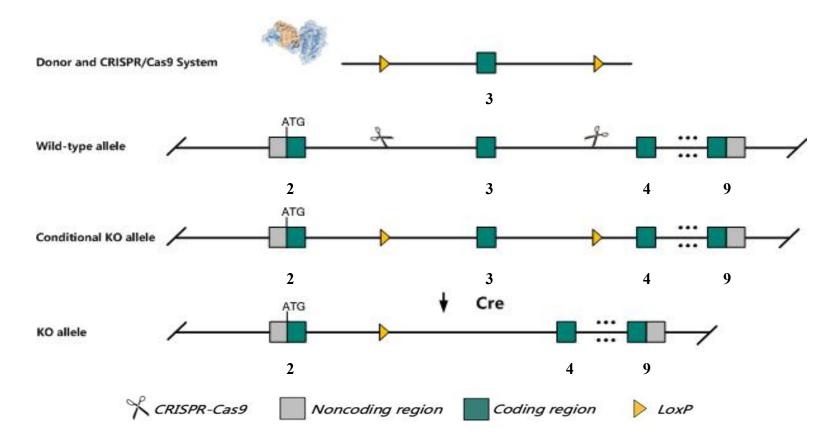
• Cas9-CKO

#### Genetic Background

• C57BL/6JGpt



# Strain Strategy



Schematic representation of CRISPR-Cas9 engineering used to edit the Slc22a17 gene.



#### Technical Information

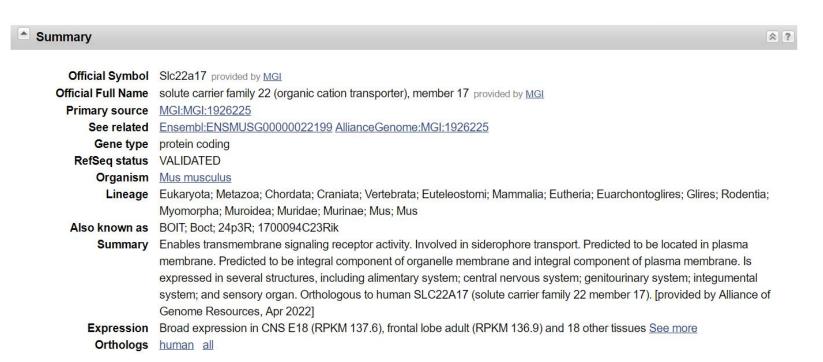
- The *Slc22a17* gene has 12 transcripts. According to the structure of *Slc22a17* gene, exon3 of *Slc22a17-201*(ENSMUST00000050772.10) transcript is recommended as the knockout region. The region contains 155bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR-Cas9 technology to modify *Slc22a17* 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.



#### Gene Information

Slc22a17 solute carrier family 22 (organic cation transporter), member 17 [ Mus musculus (house mouse) ]

Gene ID: 59049, updated on 4-Apr-2023



Source: https://www.ncbi.nlm.nih.gov/

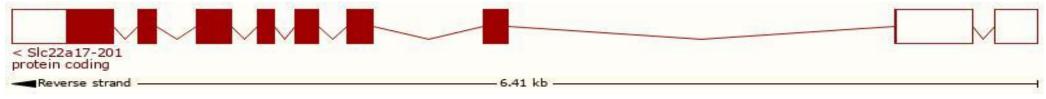


# Transcript Information

The gene has 12 transcripts, all transcripts are shown below:

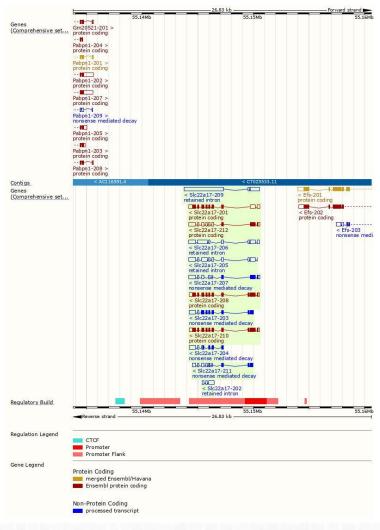
Show/hide columns (1 hidden)							Filter
Transcript ID	Name 🍦	bp 🖕	Protein	Biotype	CCDS	UniProt Match	Flags
ENSMUST00000228495.3	Slc22a17-210	2340	<u>627aa</u>	Protein coding		A0A2I3BQG7&	Ensembl Canonical GENCODE basic APPRIS P4
ENSMUST00000228119.3	Slc22a17-208	2348	<u>626aa</u>	Protein coding		A0A2I3BQL1 &	GENCODE basic   APPRIS ALT2
ENSMUST00000050772.10	Slc22a17-201	2294	401aa	Protein coding	CCDS36924 母	Q9D9E0-1 &	GENCODE basic TSL:1
ENSMUST00000231305.2	Slc22a17-212	2290	240aa	Protein coding		A0A2I3BPI6译	GENCODE basic
ENSMUST00000227880.3	Slc22a17-207	2041	240aa	Nonsense mediated decay		A0A2I3BPI6译	y-
ENSMUST00000226467.3	Slc22a17-203	1999	418aa	Nonsense mediated decay		A0A2I3BPH7&	CDS 5' incomplete
ENSMUST00000228588.2	Slc22a17-211	1446	209aa	Nonsense mediated decay		A0A2I3BR74 &	CDS 5' incomplete
ENSMUST00000226690.2	Slc22a17-204	1406	<u>198aa</u>	Nonsense mediated decay		A0A2I3BRR2₽	CDS 5' incomplete
ENSMUST00000228249.2	Slc22a17-209	4488	No protein	Retained intron		-	-
ENSMUST00000227600.2	Slc22a17-206	2595	No protein	Retained intron		-	Ε.
ENSMUST00000226718.2	Slc22a17-205	2302	No protein	Retained intron		-	7-
ENSMUST00000226456.2	Slc22a17-202	767	No protein	Retained intron		-	-

The strategy is based on the design of *Slc22a17-201* transcript, the transcription is shown below:





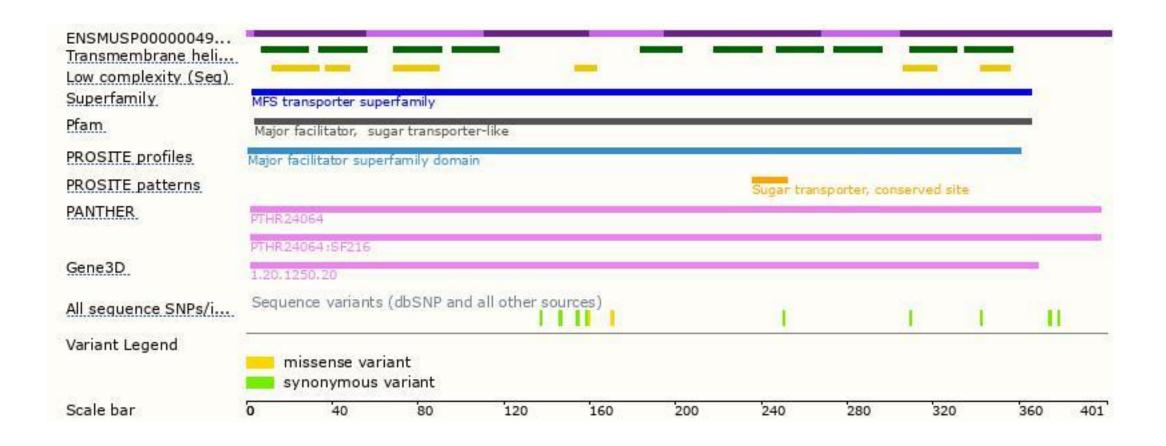
### Genomic Information





Source: : https://www.ensembl.org

#### Protein Information





Source: : https://www.ensembl.org

## Important Information

- The *Slc22a17* gene is located on the Chr14. 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.
- Exon3 of transcript Slc22a17-209 will be destroyed.
- 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.

