

# Slc15a2 Cas9-CKO Strategy

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## **Project Overview**



**Project Name** 

Slc15a2

**Project type** 

Cas9-CKO

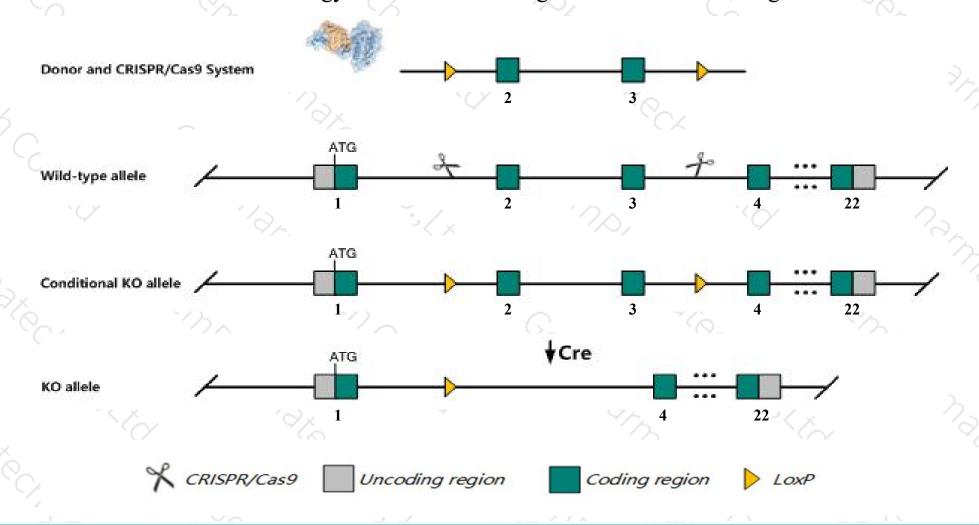
Strain background

C57BL/6JGpt

## Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Slc15a2 gene. The schematic diagram is as follows:



### Technical routes



- ➤ The *Slc15a2* gene has 14 transcripts. According to the structure of *Slc15a2* gene, exon2-exon3 of *Slc15a2-201* (ENSMUST00000023616.9) transcript is recommended as the knockout region. The region contains 230bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Slc15a2* 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, Homozygous mutant mice have impairments of dipeptide transportion, however, show no gross defects.
- ➤ Transcript *Slc15a2*-202&203&204&206&209&210&211&214 may not be affected.
- The *Slc15a2* gene is located on the Chr16. 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)



#### SIc15a2 solute carrier family 15 (H+/peptide transporter), member 2 [ Mus musculus (house mouse) ]

Gene ID: 57738, updated on 12-Aug-2019

#### Summary

☆ ?

Official Symbol Slc15a2 provided by MGI

Official Full Name solute carrier family 15 (H+/peptide transporter), member 2 provided by MGI

Primary source MGI:MGI:1890457

See related Ensembl: ENSMUSG00000022899

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 Pept2; C78862; 8430408C16Rik

Expression Broad expression in kidney adult (RPKM 7.8), frontal lobe adult (RPKM 3.5) and 17 other tissues See more

Orthologs human all

#### Genomic context



**Location:** 16; 16 B3

See Slc15a2 in Genome Data Viewer

Exon count: 24

Annotation release	Status	Assembly	Chr	Location
108	current	GRCm38.p6 (GCF_000001635.26)	16	NC_000082.6 (3675016136785158, complement)
Build 37.2	previous assembly	MGSCv37 (GCF_000001635.18)	16	NC_000082.5 (3675025036785048, complement)

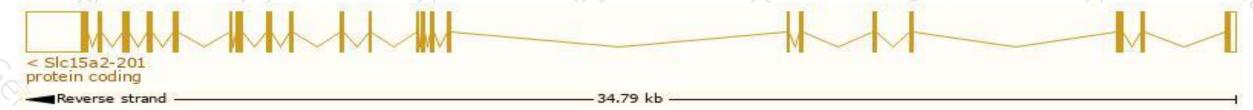
## Transcript information (Ensembl)



#### The gene has 14 transcripts, all transcripts are shown below:

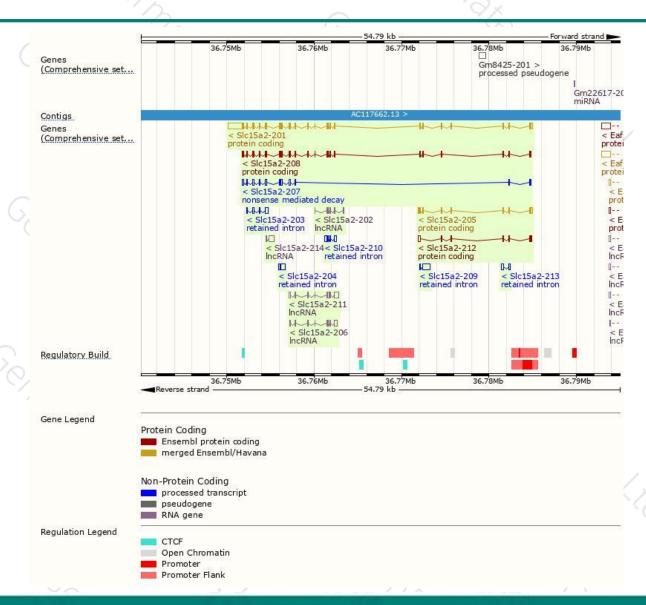
Name	Transcript ID	bp	Protein	Biotype	ccds	UniProt	Flags
SIc15a2-201	ENSMUST00000023616.9	3994	740aa	Protein coding	CCDS28156	E9QMN8	TSL:1 GENCODE basic APPRIS P2
SIc15a2-205	ENSMUST00000164579.7	1068	259aa	Protein coding	CCDS49842	E9Q329	TSL:1 GENCODE basic
SIc15a2-208	ENSMUST00000165531.7	2158	709aa	Protein coding	-	E9PYQ9	TSL:5 GENCODE basic APPRIS ALT
SIc15a2-212	ENSMUST00000168279.1	958	189aa	Protein coding	2	G3XA51	TSL:3 GENCODE basic
SIc15a2-207	ENSMUST00000165380.7	1292	84aa	Nonsense mediated decay		E9Q0L2	TSL:5
SIc15a2-209	ENSMUST00000166399.1	928	No protein	Retained intron	-	P#3	TSL:2
SIc15a2-210	ENSMUST00000167909.1	729	No protein	Retained intron	-	120	TSL:2 TSL:3
SIc15a2-204	ENSMUST00000163964.1	679	No protein	Retained intron	-		
SIc15a2-203	ENSMUST00000163471.1	622	No protein	Retained intron		1271	TSL:3
SIc15a2-213	ENSMUST00000169644.1	516	No protein	Retained intron	-	1943	TSL:3
SIc15a2-211	ENSMUST00000167941.7	1170	No protein	IncRNA	2	020	TSL:5
SIc15a2-206	ENSMUST00000164770.7	874	No protein	IncRNA	-		TSL:5 TSL:3
SIc15a2-214	ENSMUST00000172382.1	694	No protein	IncRNA			
SIc15a2-202	ENSMUST00000100308.9	430	No protein	IncRNA			TSL:3

The strategy is based on the design of Slc15a2-201 transcript, The transcription is shown below



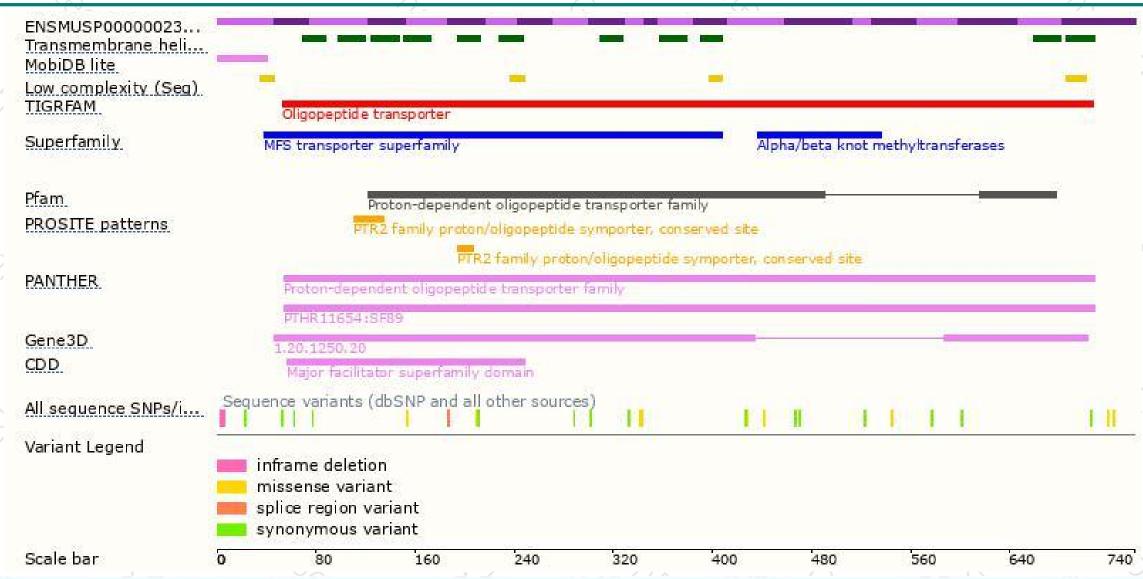
### Genomic location distribution





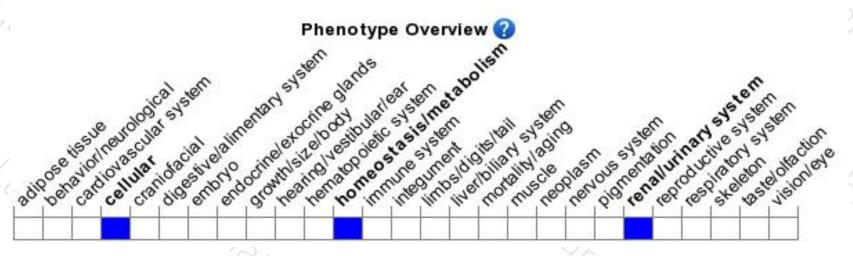
### Protein domain





## 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, Homozygous mutant mice have impairments of dipeptide transportion, however, show no gross defects.



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





