

Slc6a8 Cas9-KO Strategy

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Project Overview



Project Name

Slc6a8

Project type

Cas9-KO

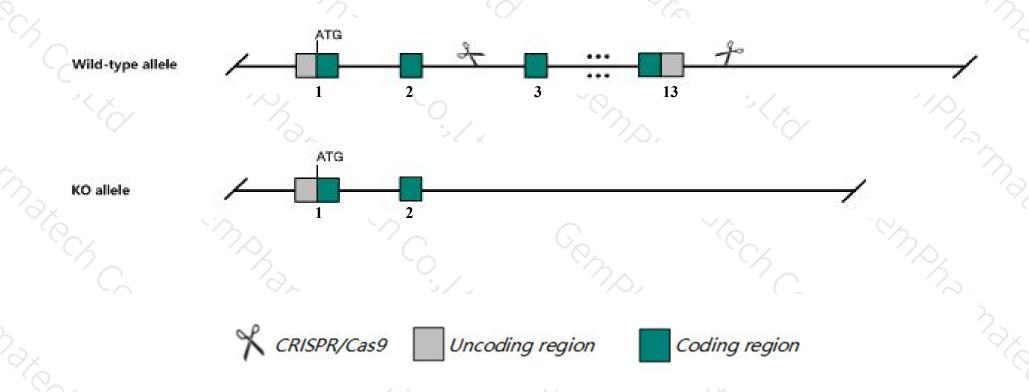
Strain background

C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Slc6a8 gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Slc6a8* gene has 13 transcripts. According to the structure of *Slc6a8* gene, exon3-exon13 of *Slc6a8-201* (ENSMUST00000033752.13) 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 Slc6a8 gene. The brief process is as follows: CRISPR/Cas9 system were

Notice



- ➤ According to the existing MGI data, Male mice hemizygous for a targeted allele exhibit decreased body weight, decreased creatine concentrations, impaired short term object recognition, impaired contextual conditioning, altered locomotor activity, and increased serotonine levels in the brain.
- The N-terminal of Slc6a8 gene will remain several amino acids ,it may remain the partial function of Slc6a8 gene.
- > The Slc6a8 gene is located on the ChrX. 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)



SIc6a8 solute carrier family 6 (neurotransmitter transporter, creatine), member 8 [Mus musculus (house mouse)]

Gene ID: 102857, updated on 10-Oct-2019

Summary

△ ?

Official Symbol Slc6a8 provided by MGI

Official Full Name solute carrier family 6 (neurotransmitter transporter, creatine), member 8 provided by MGI

Primary source MGI:MGI:2147834

See related Ensembl:ENSMUSG00000019558

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 CRT; CT1; CRTR; CTR5; Creat; AA589632

Expression Broad expression in duodenum adult (RPKM 226.8), small intestine adult (RPKM 214.6) and 18 other tissues See more

Orthologs <u>human</u> all

Genomic context



Location: X A7.3; X 37.38 cM

See Slc6a8 in Genome Data Viewer

Exon count: 13

Annotation release	Status	Assembly	Chr	Location	
108	current	GRCm38.p6 (GCF_000001635.26)	X	NC_000086.7 (7367313373682502)	
Build 37.2	previous assembly	MGSCv37 (GCF_000001635.18)	X	NC_000086.6 (7091847270927841)	

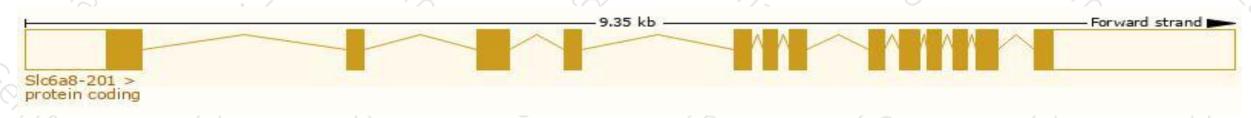
Transcript information (Ensembl)



The gene has 13 transcripts, all transcripts are shown below:

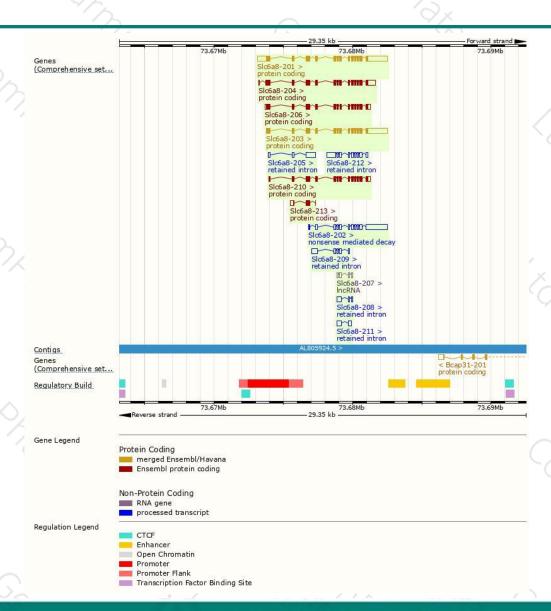
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
SIc6a8-201	ENSMUST00000033752.13	3964	<u>640aa</u>	Protein coding	CCDS30208	Q8VBW1	TSL:1 GENCODE basic
SIc6a8-203	ENSMUST00000114465.8	3314	<u>635aa</u>	Protein coding	CCDS53098	A2ALM6 Q8VBW1	TSL:1 GENCODE basic APPRIS P1
SIc6a8-204	ENSMUST00000114467.8	2509	<u>635aa</u>	Protein coding	CCDS53098	A2ALM6 Q8VBW1	TSL:1 GENCODE basic APPRIS P1
SIc6a8-206	ENSMUST00000164449.7	2151	<u>625aa</u>	Protein coding	-	E9Q151	TSL:5 GENCODE basic
SIc6a8-210	ENSMUST00000168831.7	1919	<u>576aa</u>	Protein coding		F6UKB4	CDS 5' incomplete TSL:1
SIc6a8-213	ENSMUST00000171398.1	537	<u>107aa</u>	Protein coding	-	E9Q6T2	CDS 3' incomplete TSL:3
SIc6a8-202	ENSMUST00000114464.8	2729	<u>19aa</u>	Nonsense mediated decay	-	F7C668	CDS 5' incomplete TSL:1
SIc6a8-212	ENSMUST00000170614.7	1613	No protein	Retained intron	÷	(<u>4</u>	TSL:2
SIc6a8-205	ENSMUST00000146796.2	940	No protein	Retained intron		15	TSL:2
SIc6a8-209	ENSMUST00000168764.7	843	No protein	Retained intron	-	H=	TSL:3
SIc6a8-211	ENSMUST00000170574.7	527	No protein	Retained intron	0	N=	TSL:3
SIc6a8-208	ENSMUST00000167828.1	493	No protein	Retained intron	-	-	TSL:1
SIc6a8-207	ENSMUST00000166200.7	404	No protein	IncRNA		-	TSL:3
4							·

The strategy is based on the design of Slc6a8-201 transcript, The transcription is shown below



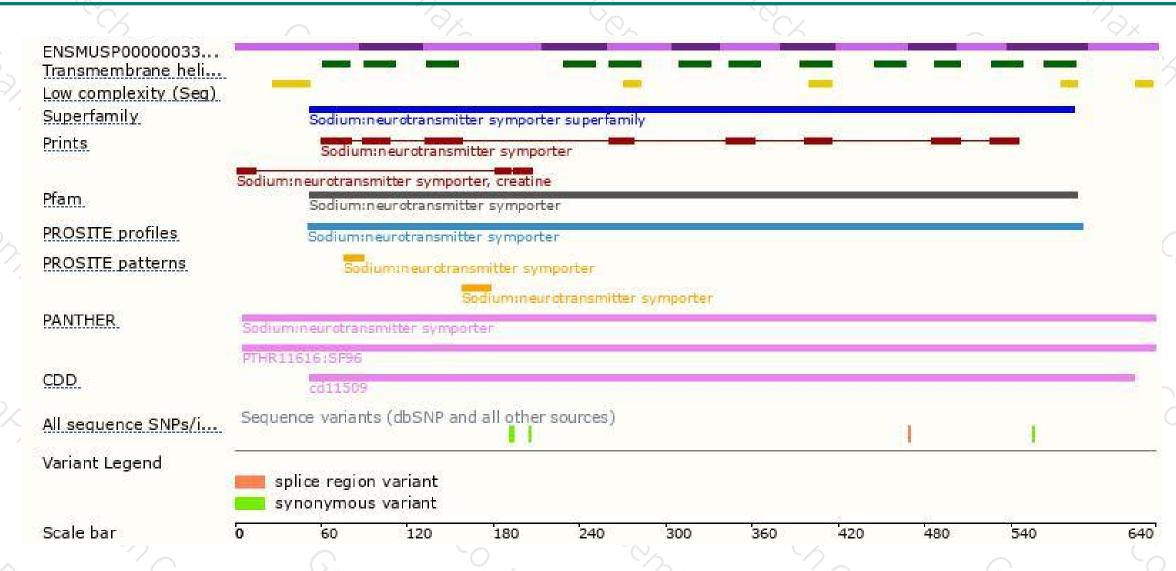
Genomic location distribution





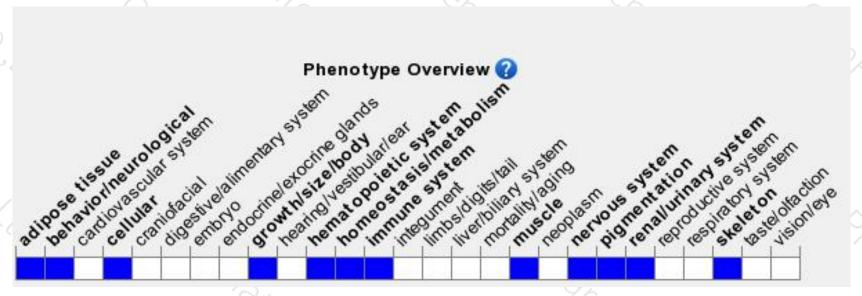
Protein domain





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, Male mice hemizygous for a targeted allele exhibit decreased body weight, decreased creatine concentrations, impaired short term object recognition, impaired contextual conditioning, altered locomoto activity, and increased serotonine levels in the brain.



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





