

# Slc24a2 Cas9-KO Strategy

Designer: JiaYu

Reviewer: Xiaojing Li

**Design Date:** 2020-2-19

# **Project Overview**



**Project Name** 

Slc24a2

**Project type** 

Cas9-KO

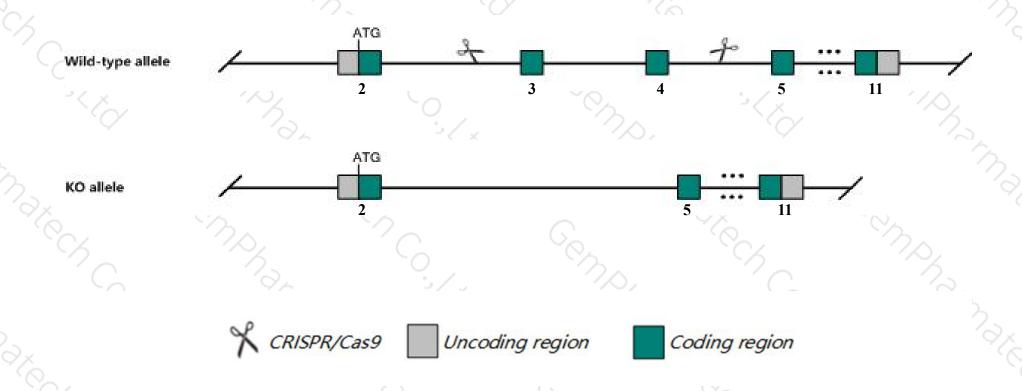
Strain background

C57BL/6JGpt

# **Knockout strategy**



This model will use CRISPR/Cas9 technology to edit the Slc24a2 gene. The schematic diagram is as follows:



### **Technical routes**



- ➤ The Slc24a2 gene has 9 transcripts. According to the structure of Slc24a2 gene, exon3-exon4 of Slc24a2-203 (ENSMUST00000107157.8) transcript is recommended as the knockout region. The region contains 148bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify Slc24a2 gene. The brief process is as follows: CRISPR/Cas9 syste

### **Notice**



- ➤ According to the existing MGI data, Homozygous mutation of this gene results in loss of long term potentiation and an increase in long term depression and deficits in motor learning and spatial working memory.
- > Some amino acids will remain at the N-terminus and some functions may be retained.
- The Slc24a2 gene is located on the Chr4. 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)



#### Slc24a2 solute carrier family 24 (sodium/potassium/calcium exchanger), member 2 [Mus musculus (house mouse)]

Gene ID: 76376, updated on 5-Mar-2019

#### Summary



Official Symbol Slc24a2 provided by MGI

Official Full Name solute carrier family 24 (sodium/potassium/calcium exchanger), member 2 provided by MGI

Primary source MGI:MGI:1923626

See related Ensembl:ENSMUSG00000037996

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 2810021B17Rik, 6330417K15Rik, Al847460, Nckx2

Expression Biased expression in cortex adult (RPKM 27.0), cerebellum adult (RPKM 20.2) and 2 other tissuesSee more

Orthologs <u>human</u> all

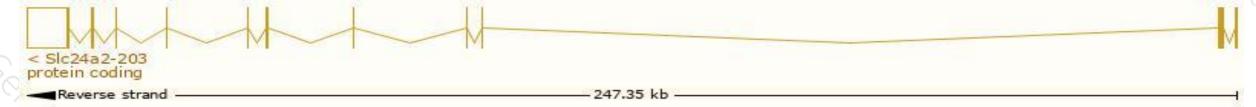
# Transcript information (Ensembl)



The gene has 9 transcripts, all transcripts are shown below:

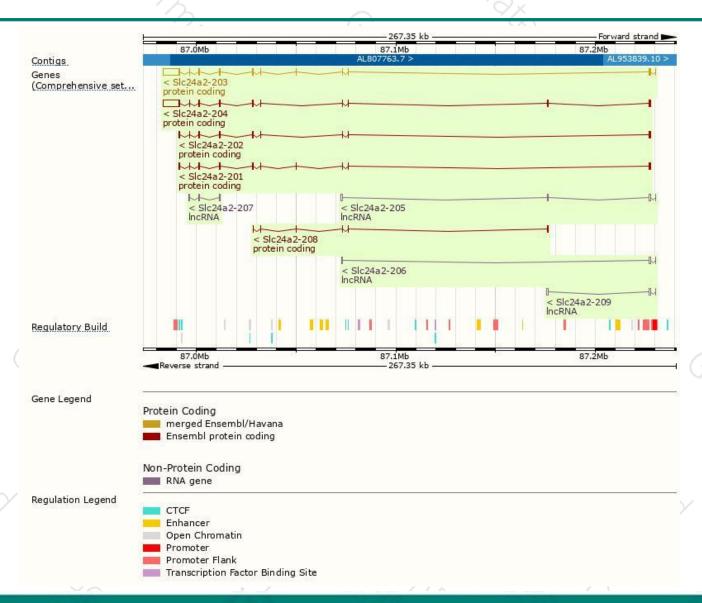
- 1		1			-		
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
SIc24a2-203	ENSMUST00000107157.8	10531	666aa	Protein coding	CCDS18312	Q8BUN9	TSL:1 GENCODE basic APPRIS P2
SIc24a2-204	ENSMUST00000107158.8	10499	<u>711aa</u>	Protein coding	CCDS51220	Q14BI1	TSL:1 GENCODE basic
SIc24a2-201	ENSMUST00000044990.10	2218	<u>662aa</u>	Protein coding	5	B1AXF2	TSL:5 GENCODE basic APPRIS ALT1
SIc24a2-202	ENSMUST00000107155.7	2171	645aa	Protein coding	-	B1AXF3	TSL:5 GENCODE basic APPRIS ALT1
SIc24a2-208	ENSMUST00000146815.1	504	<u>168aa</u>	Protein coding		F6RT95	5' and 3' truncations in transcript evidence prevent annotation of the start and the end of the CDS. CDS 5' and 3' incomplete TSL:
SIc24a2-205	ENSMUST00000134248.7	2230	No protein	IncRNA	į, ė	3.5	TSL:2
SIc24a2-209	ENSMUST00000155361.1	1796	No protein	IncRNA	-	0.20	TSL:2
SIc24a2-206	ENSMUST00000134643.7	1655	No protein	IncRNA	-	1528	TSL:2
SIc24a2-207	ENSMUST00000140780.1	679	No protein	IncRNA			TSL:3

The strategy is based on the design of Slc24a2-203 transcript, The transcription is shown below



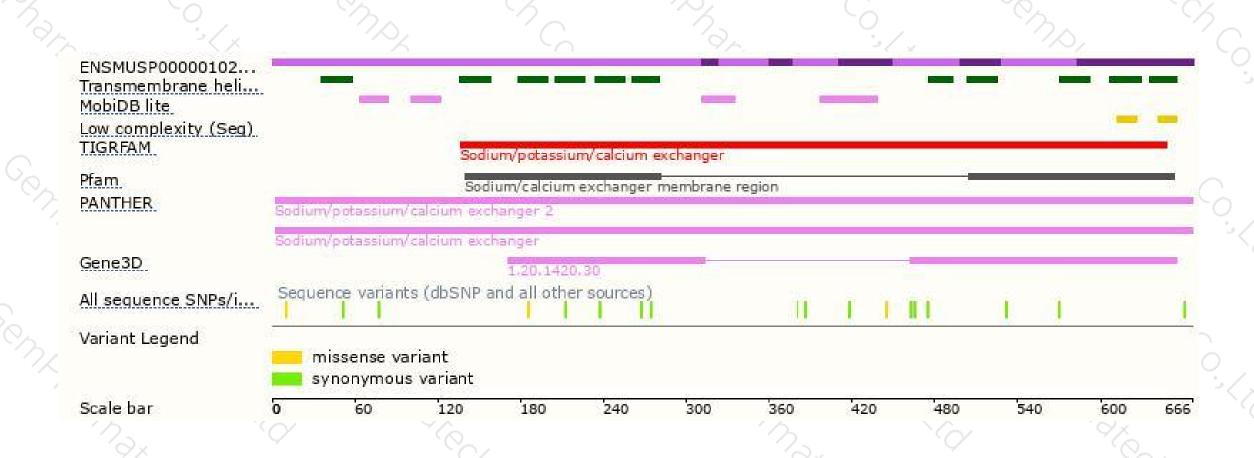
### Genomic location distribution





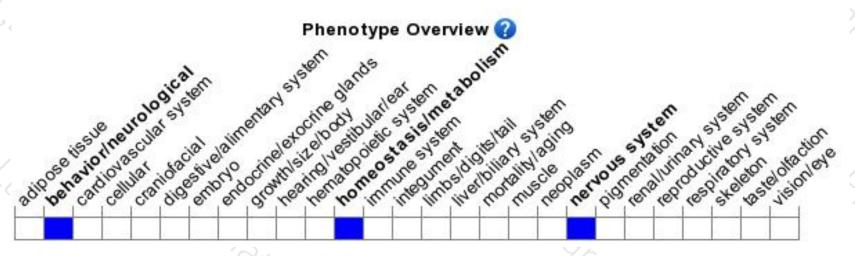
### Protein domain





# 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 mutation of this gene results in loss of long term potentiation and an increase in long term depression and deficits in motor learning and spatial working memory.



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





