

Slitrk5 Cas9-CKO Strategy

Designer: Daohua Xu

Reviewer: Huimin Su

Design Date: 2020-4-15

Project Overview



Project Name

Slitrk5

Project type

Cas9-CKO

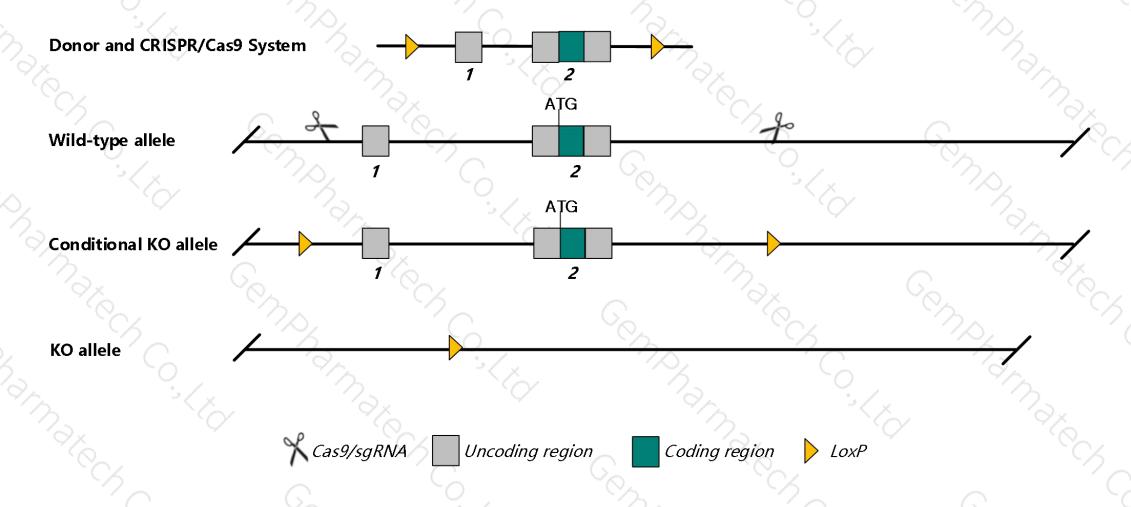
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Slitrk5 gene. The schematic diagram is as follows:



Technical routes



- The *Slitrk5* gene has 2 transcripts. According to the structure of *Slitrk5* gene, exon1-exon2 of *Slitrk5-201* (ENSMUST00000042767.8) transcript is recommended as the knockout region. The region contains all 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 *Slitrk5* 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 null allele have abnormal medium spiny neuron morphology and exhibit behavioral abnormalities.
- The *Slitrk5* 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.
- > 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)



Slitrk5 SLIT and NTRK-like family, member 5 [Mus musculus (house mouse)]

Gene ID: 75409, updated on 13-Mar-2020

Summary

☆ ?

Official Symbol Slitrk5 provided by MGI

Official Full Name SLIT and NTRK-like family, member 5 provided by MGI

Primary source MGI:MGI:2679448

See related Ensembl: ENSMUSG00000033214

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 2610019D03Rik

Expression Biased expression in frontal lobe adult (RPKM 6.2), CNS E18 (RPKM 6.1) and 6 other tissuesSee more

Orthologs human all

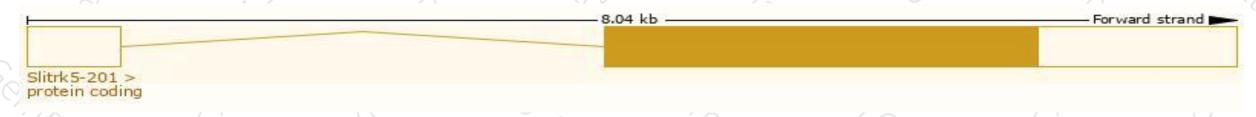
Transcript information (Ensembl)



The gene has 2 transcripts, all transcripts are shown below:

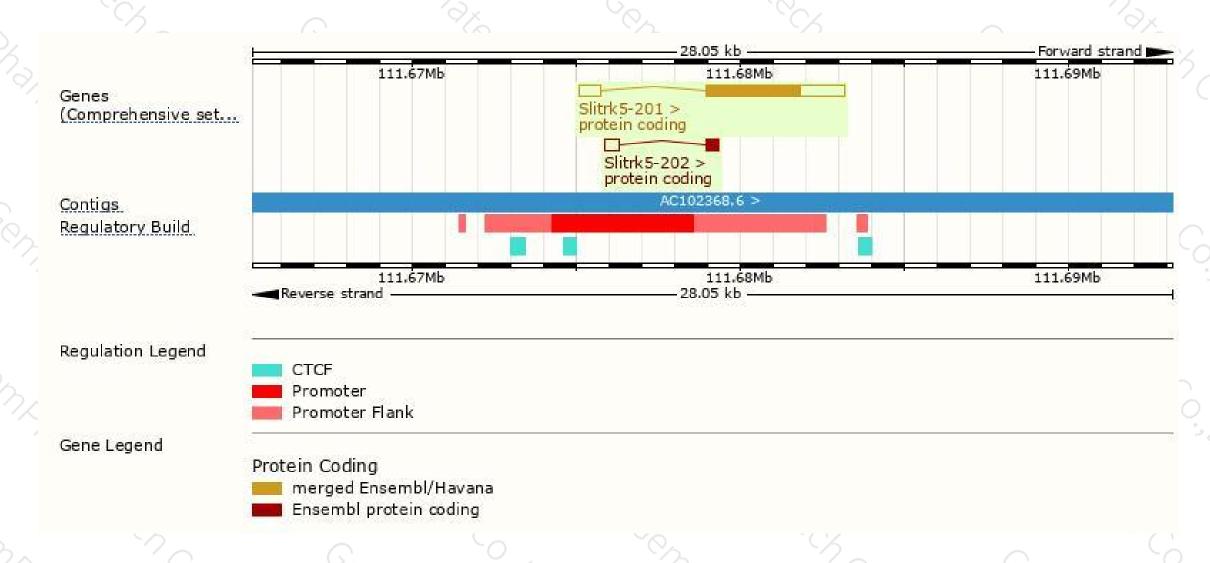
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Slitrk5-201	ENSMUST00000042767.8	4831	957aa	Protein coding		Q810B7	TSL:2 GENCODE basic APPRIS is a system to annotate alternatively spliced transcripts based on a range of computational methods to identify the most functionally important transcript(s) of a gene. APPRIS P1
Slitrk5-202	ENSMUST00000227891.1	837	132aa	Protein coding		A0A2I3BQY9	CDS 3' incomplete

The strategy is based on the design of *Slitrk5-201* transcript, the transcription is shown below:



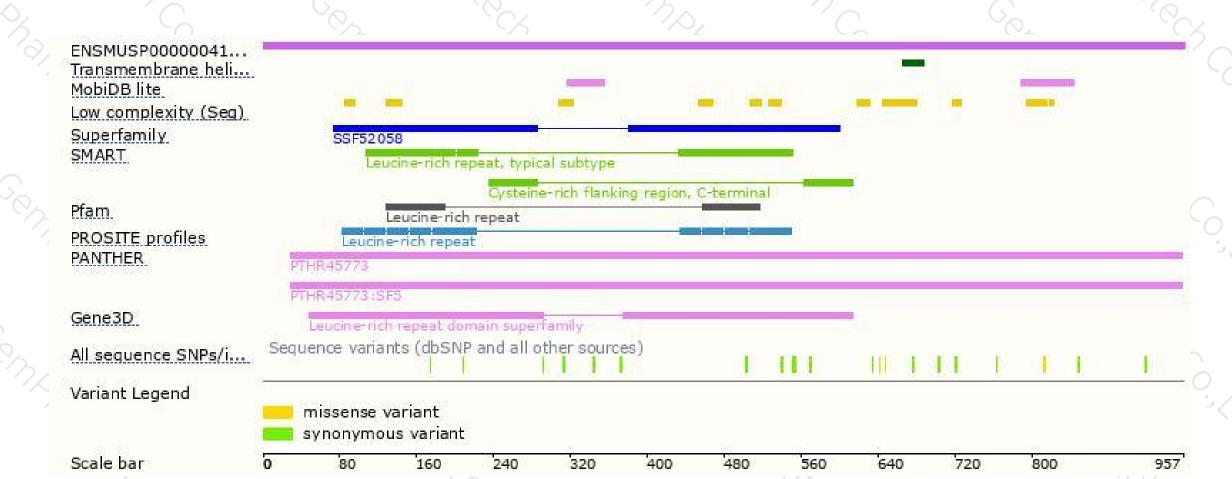
Genomic location distribution





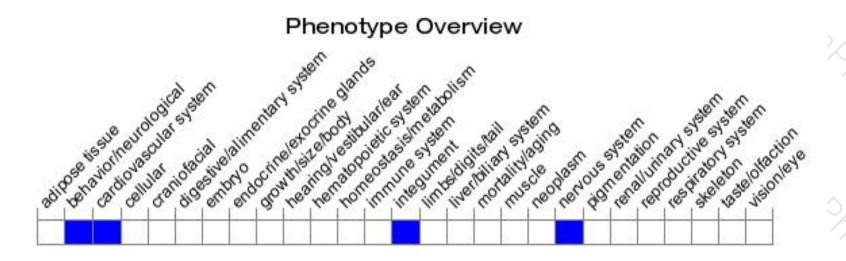
Protein domain





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 null allele have abnormal medium spiny neuron morphology and exhibit behavioral abnormalities.



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





