

# Shprh Cas9-CKO Strategy

Designer: Daohua Xu

Reviewer: Huimin Su

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## **Project Overview**



**Project Name** 

Shprh

**Project type** 

Cas9-CKO

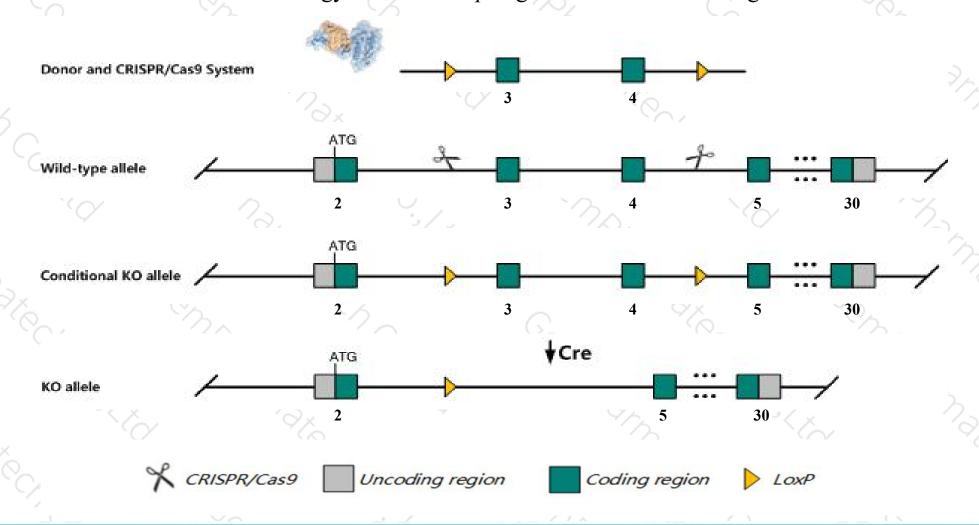
Strain background

C57BL/6JGpt

## Conditional Knockout strategy



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



### Technical routes



- The *Shprh* gene has 8 transcripts. According to the structure of *Shprh* gene, exon3-exon4 of *Shprh-201* (ENSMUST00000044053.12) transcript is recommended as the knockout region. The region contains 349bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Shprh* 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, the gene product is an e3 ligase involved in poly-ubiquitination of
- > The *Shprh* gene is located on the Chr10. 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)



#### Shprh SNF2 histone linker PHD RING helicase [Mus musculus (house mouse)]

Gene ID: 268281, updated on 13-Mar-2020

#### Summary

☆ ?

Official Symbol Shprh provided by MGI

Official Full Name SNF2 histone linker PHD RING helicase provided by MGI

Primary source MGI:MGI:1917581

See related Ensembl: ENSMUSG00000090112

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 2610103K11Rik, AA450458, AU024614, BC006883, D230017O13Rik, E130018M05

Expression Broad expression in cerebellum adult (RPKM 3.8), CNS E18 (RPKM 3.8) and 25 other tissuesSee more

Orthologs <u>human all</u>

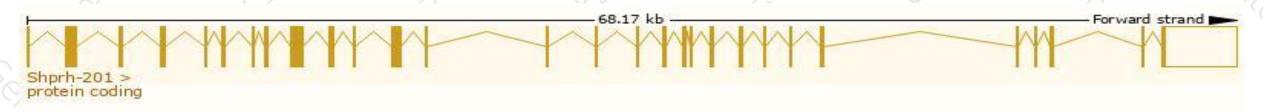
## Transcript information (Ensembl)



The gene has 8 transcripts, all transcripts are shown below:

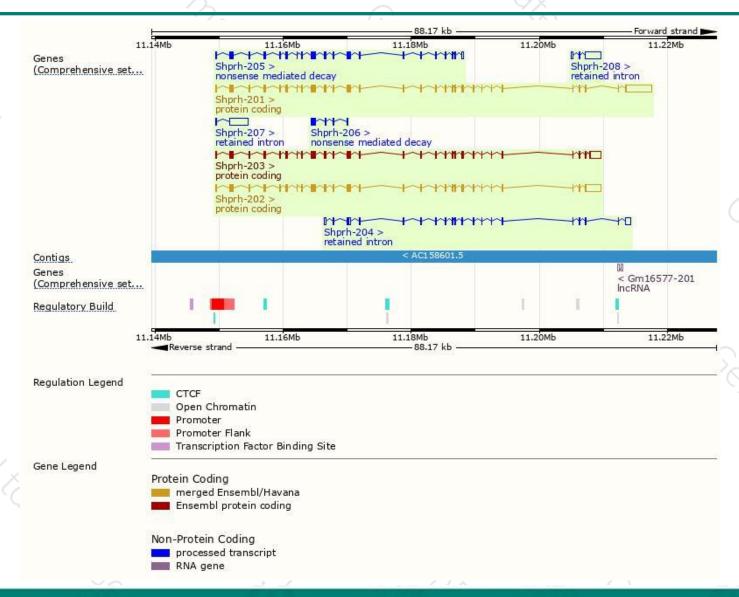
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Shprh-201	ENSMUST00000044053.12	9271	1674aa	Protein coding	CCDS35838	Q7TPQ3	TSL:1 GENCODE basic APPRIS P4
Shprh-202	ENSMUST00000054814.13	7225	1616aa	Protein coding	CCDS35839	Q7TPQ3	TSL:1 GENCODE basic APPRIS ALT
Shprh-203	ENSMUST00000159541.7	6679	1646aa	Protein coding	CCDS70003	Q7TPQ3	TSL:1 GENCODE basic APPRIS ALT
Shprh-205	ENSMUST00000159810.7	4171	<u>1031aa</u>	Nonsense mediated decay	720	Q7TPQ3	TSL:1
Shprh-206	ENSMUST00000160461.7	945	218aa	Nonsense mediated decay	151	F7D4T4	CDS 5' incomplete TSL:3
Shprh-204	ENSMUST00000159553.7	3467	No protein	Retained intron	94.3	19-	TSL:1
Shprh-207	ENSMUST00000161858.1	3028	No protein	Retained intron	12	32	TSL:1
Shprh-208	ENSMUST00000161910.1	2857	No protein	Retained intron	728	02	TSL:1

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



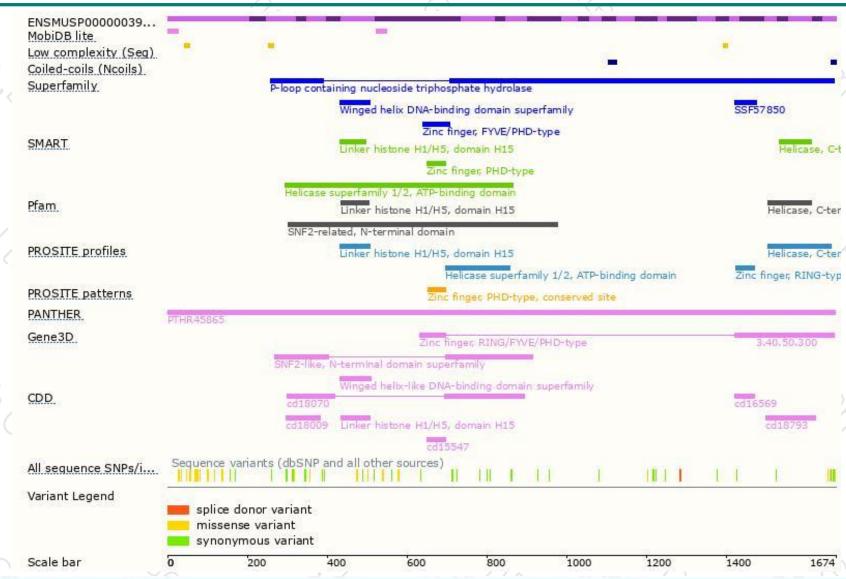
### Genomic location distribution





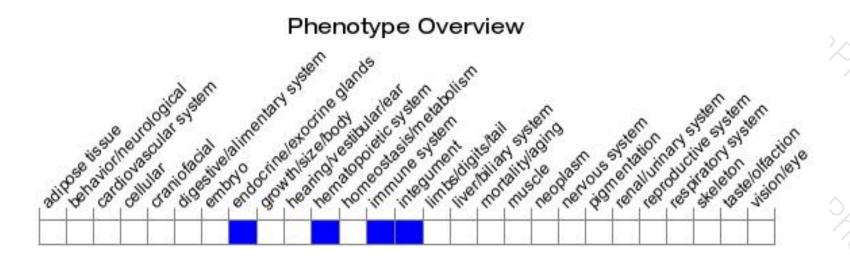
### Protein domain





## 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, the gene product is an E3 ligase involved in poly-ubiquitination of



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





