

Shroom4 Cas9-CKO Strategy

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Overview

Target Gene Name

• Shroom4

Project Type

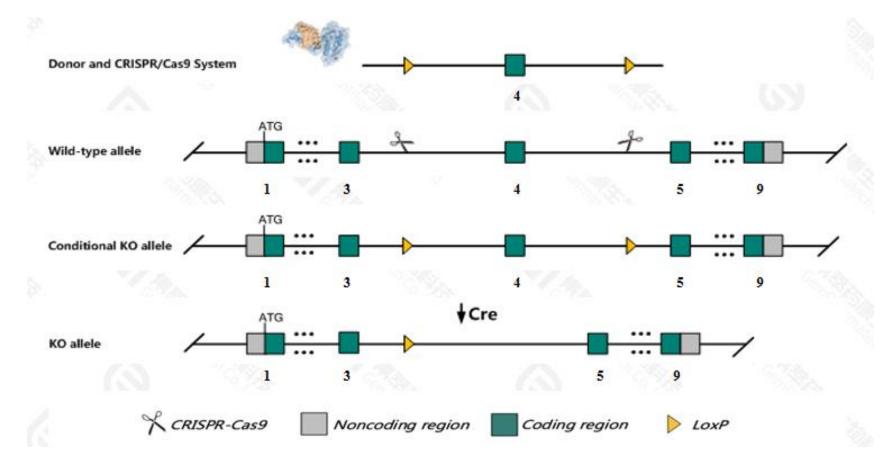
• Cas9-CKO

Genetic Background

• C57BL/6JGpt



Strain Strategy



Schematic representation of CRISPR-Cas9 engineering used to edit the *Shroom4* gene.



Technical Information

- The *Shroom4* gene has 3 transcripts. According to the structure of *Shroom4* gene, exon4 of 202(ENSMUST00000103005.10) transcript is recommended as the knockout region. The region contains 2464bp of coding sequences. Knocking out the region will result in disruption of protein function.
- In this project we use CRISPR-Cas9 technology to modify *Shroom4* 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 on-target amplicon sequencing. A stable F1-generation mouse strain was obtained by mating positive F0-generation mice with C57BL/6JGpt mice and confirmation of the desired mutant allele was carried out by PCR and on-target amplicon sequencing.
- 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.



Gene Information

Shroom4 shroom family member 4 [Mus musculus (house mouse)]

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Gene ID: 208431, updated on 26-Sep-2022

Summary

☆ ?

Official Symbol Shroom4 provided by MGI

Official Full Name shroom family member 4 provided by MGI

Primary source MGI:MGI:2685570

See related Ensembl: ENSMUSG00000068270 AllianceGenome: MGI: 2685570

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 Gm724; Shrm4; D430043L16Rik

Summary Enables actin filament binding activity and myosin II binding activity. Acts upstream of or within actin filament organization. Located in several cellular components,

including apical plasma membrane; basal plasma membrane; and stress fiber. Colocalizes with cortical actin cytoskeleton. Is expressed in several structures, including brain; genitourinary system; heart; liver; and lung. Human ortholog(s) of this gene implicated in Stocco Dos Santos type X-linked intellectual disability. Orthologous to

human SHROOM4 (shroom family member 4). [provided by Alliance of Genome Resources, Apr 2022]

Expression Broad expression in kidney adult (RPKM 3.7), lung adult (RPKM 2.9) and 16 other tissues See more

Orthologs human all

Try the new Gene table

Try the new Transcript table

Genomic context

☆ ?

Location: X; X A1.1

See Shroom4 in Genome Data Viewer

Exon count: 11

Source: https://www.ncbi.nlm.nih.gov/

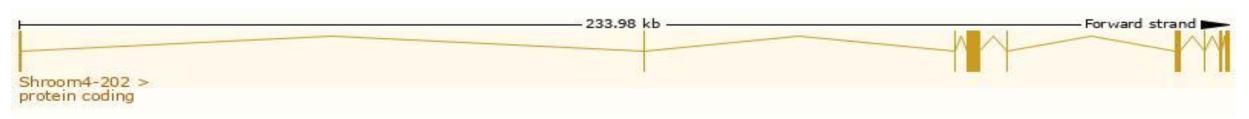


Transcript Information

The gene has 3 transcripts, all transcripts are shown below:

Transcript ID	Name 🍦	bp 🍦	Protein 🍦	Biotype	CCDS .	UniProt Match	Flags
ENSMUST00000103005.10	Shroom4-202	4839	<u>1475aa</u>	Protein coding	CCDS29959 ₽	Q1W617-1&	Ensembl Canonical GENCODE basic APPRIS P1 TSL:1
ENSMUST00000089520.3	Shroom4-201	8201	1359aa	Protein coding	CCDS85744 ₺	Q1W617-2₽	GENCODE basic TSL:1
ENSMUST00000143641.4	Shroom4-203	755	200aa	Protein coding		E9PUX3₺	TSL:3 CDS 3' incomplete

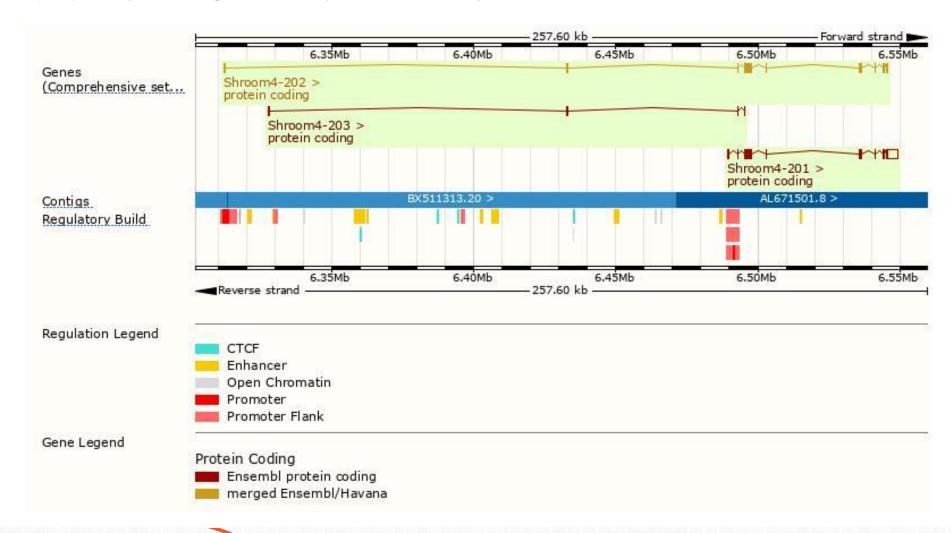
The strategy is based on the design of *Shroom4-202* transcript, the transcription is shown below:

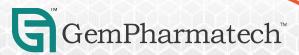


Source: https://www.ensembl.org



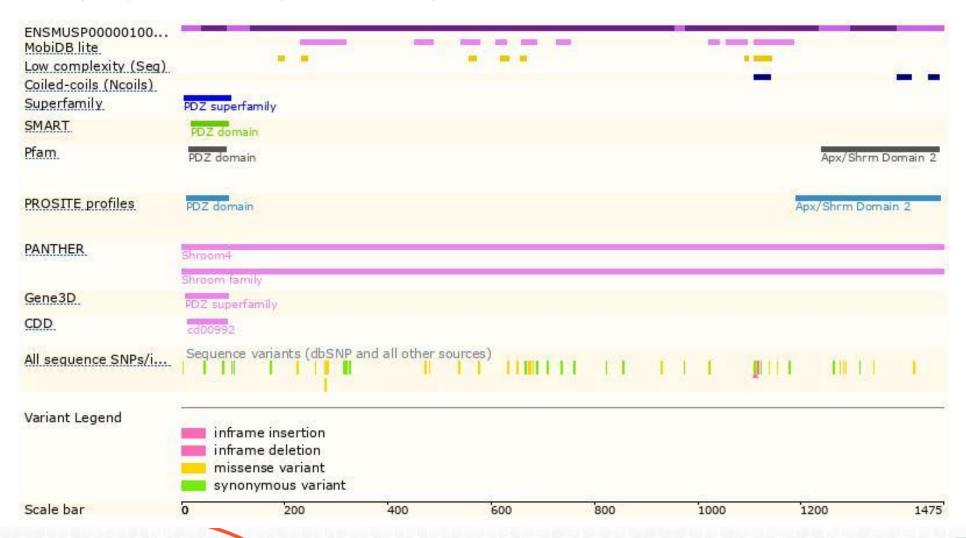
Genomic Information





Source: : https://www.ensembl.org

Protein Information



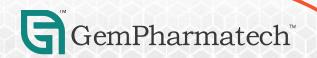


Source: : https://www.ensembl.org

Mouse Phenotype Information (MGI)



• Phenotypes affected by the mutations of Shroom4 gene are marked in blue. Data quoted from MGI database(http://www.informatics.jax.org/).



Important Information

- The *Shroom4* 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 loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at existing technological level.

