

Snx27 Cas9-CKO Strategy

Designer:

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Design Date:

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Project Overview



Project Name

Snx27

Project type

Cas9-CKO

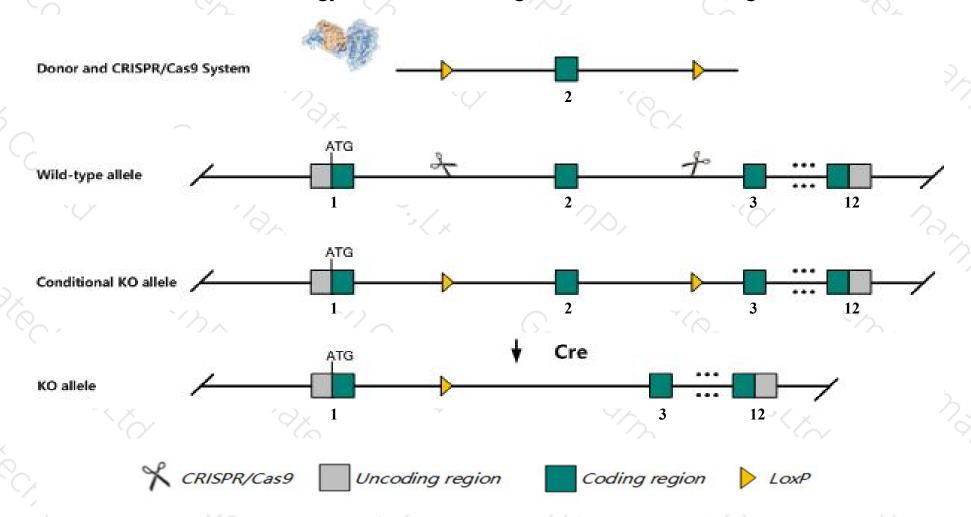
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Snx27* gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Snx27* gene has 8 transcripts. According to the structure of *Snx27* gene, exon2 of *Snx27-201*(ENSMUST00000029783.15) transcript is recommended as the knockout region. The region contains 232bp coding sequence.

 Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Snx27* 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 knock-out allele exhibit prenatal and postnatal lethality, decreased organ size, slow postnatal weight gain, and decreased endocytosis of Grin2c.
- The *Snx27* gene is located on the Chr3. 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)



Snx27 sorting nexin family member 27 [Mus musculus (house mouse)]

Gene ID: 76742, updated on 19-Mar-2019

Summary

☆ ?

Official Symbol Snx27 provided by MGI

Official Full Name sorting nexin family member 27 provided by MGI

Primary source MGI:MGI:1923992

See related Ensembl:ENSMUSG00000028136

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 5730552M22Rik, ESTM45, ESTM47, R75405

Expression Ubiquitous expression in CNS E18 (RPKM 6.3), cerebellum adult (RPKM 5.7) and 28 other tissuesSee more

Orthologs <u>human</u> all

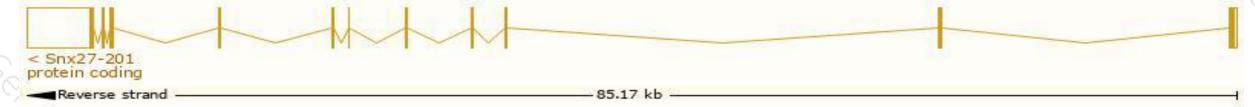
Transcript information (Ensembl)



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

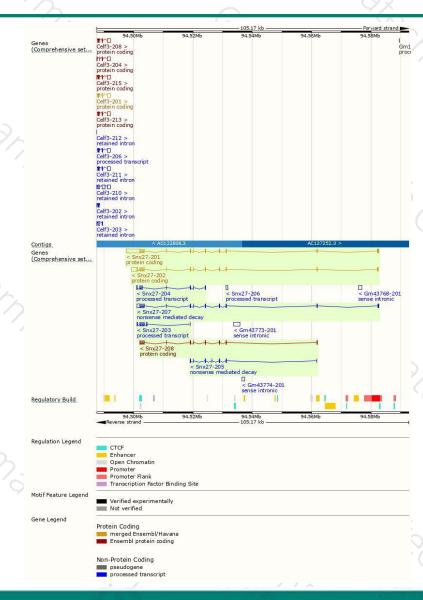
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Snx27-201	ENSMUST00000029783.15	6261	<u>526aa</u>	Protein coding	CCDS38538	Q3UHD6	TSL:1 GENCODE basic
Snx27-202	ENSMUST00000107283.7	3531	<u>539aa</u>	Protein coding	CCDS38537	Q3UHD6	TSL:1 GENCODE basic APPRIS P1
Snx27-208	ENSMUST00000200642.4	1308	<u>435aa</u>	Protein coding	120	A0A0G2JF85	TSL:5 GENCODE basic
Snx27-207	ENSMUST00000199462.4	2510	<u>63aa</u>	Nonsense mediated decay	757	A0A0G2JG07	TSL:5
Snx27-205	ENSMUST00000198426.1	599	<u>110aa</u>	Nonsense mediated decay	1871	A0A0G2JGD5	CDS 5' incomplete TSL:3
Snx27-204	ENSMUST00000196675.4	734	No protein	Processed transcript	N#3	ě	TSL:5
Snx27-206	ENSMUST00000198902.1	649	No protein	Processed transcript	020	ū.	TSL:NA
Snx27-203	ENSMUST00000196251.1	599	No protein	Processed transcript	797	22	TSL:5

The strategy is based on the design of *Snx27-201* transcript, The transcription is shown below



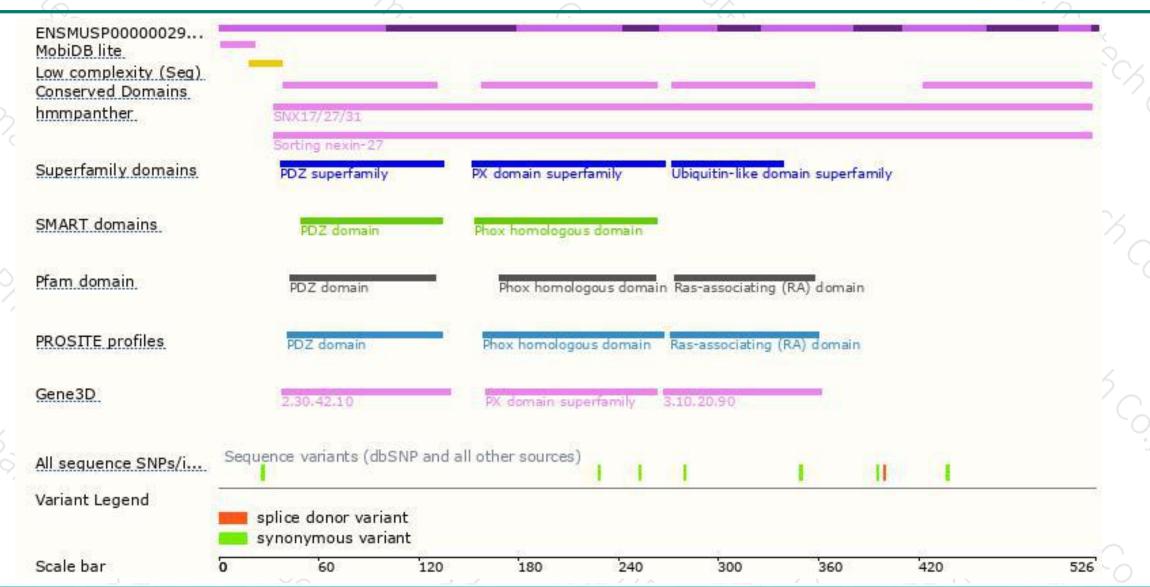
Genomic location distribution





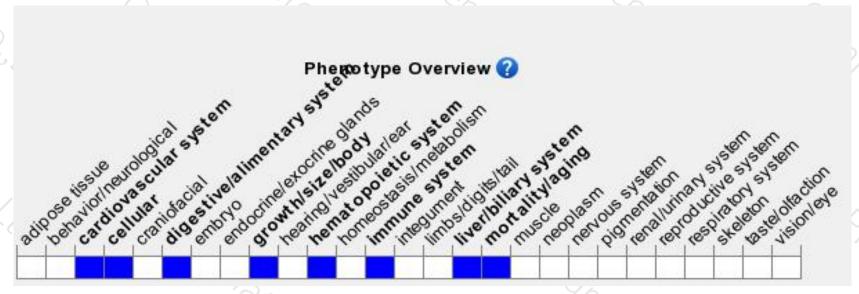
Protein domain





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 knock-out allele exhibit prenatal and postnatal lethality, decreased organ size, slow postnatal weight gain, and decreased endocytosis of Grin2c.



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





