

Samd8 Cas9-CKO Strategy

Designer: Daohua Xu

Reviewer: Huimin Su

Design Date: 2019-9-28

Project Overview



Project Name

Samd8

Project type

Cas9-CKO

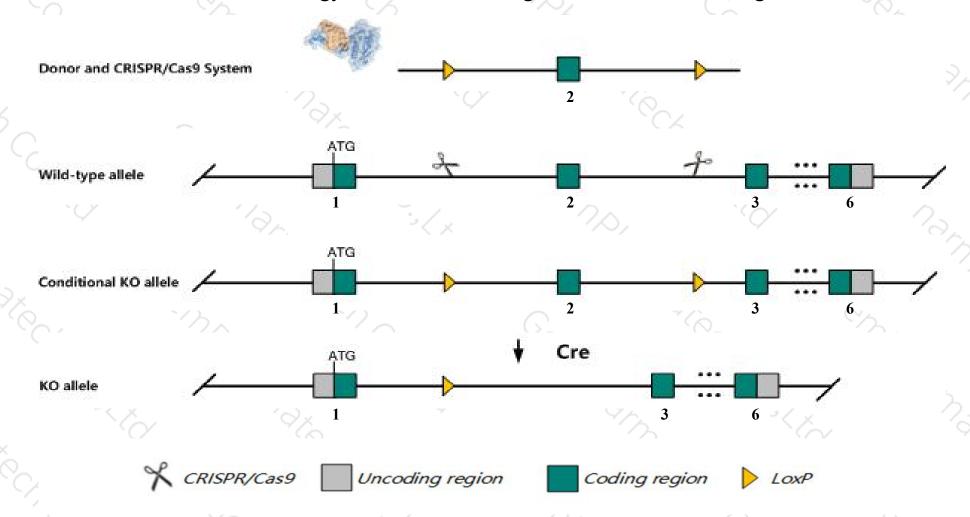
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Samd8 gene. The schematic diagram is as follows:



Technical routes



- ➤ The Samd8 gene has 4 transcripts. According to the structure of Samd8 gene, exon2 of Samd8-201

 (ENSMUST00000022292.9) transcript is recommended as the knockout region. The region contains 593bp coding sequence.

 Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Samd8* 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 decreased ceramide phosphoethanolamine synthase activity but normal liver, kidney and spleen histology.
- The Samd8 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)



Samd8 sterile alpha motif domain containing 8 [Mus musculus (house mouse)]

Gene ID: 67630, updated on 31-Jan-2019

Summary

☆ ?

Official Symbol Samd8 provided by MGI

Official Full Name sterile alpha motif domain containing 8 provided by MGI

Primary source MGI:MGI:1914880

See related Ensembl: ENSMUSG00000021770

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 1110053F04Rik, 1700010P07Rik, SMSr

Expression Ubiquitous expression in testis adult (RPKM 10.0), CNS E18 (RPKM 6.8) and 28 other tissuesSee more

Orthologs <u>human all</u>

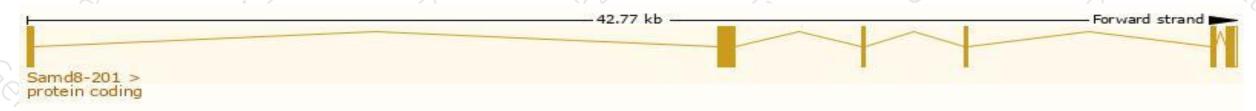
Transcript information (Ensembl)



The gene has 4 transcripts, all transcripts are shown below:

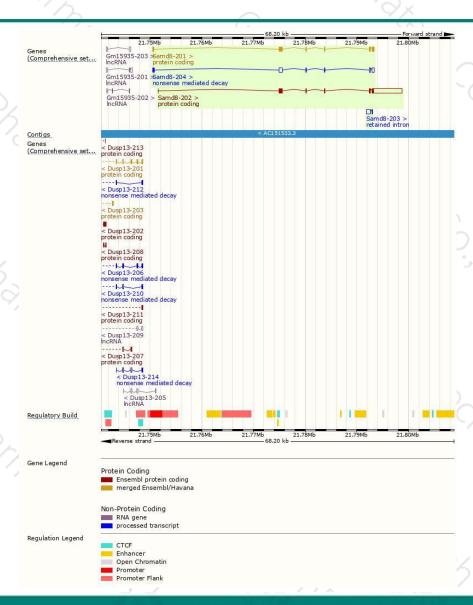
J 10.	Alter.			and the second s			d kus.
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Samd8-201	ENSMUST00000022292.9	1574	478aa	Protein coding	CCDS26866	Q14AQ4 Q9DA37	TSL:1 GENCODE basic
Samd8-202	ENSMUST00000119430.2	6801	<u>415aa</u>	Protein coding		Q3UH82	TSL:1 GENCODE basic APPRIS P1
Samd8-204	ENSMUST00000144061.1	1613	<u>64aa</u>	Nonsense mediated decay	-	D6REI9	TSL:1
Samd8-203	ENSMUST00000142023.1	722	No protein	Retained intron	92	42	TSL:2

The strategy is based on the design of Samd8-201 transcript, The transcription is shown below



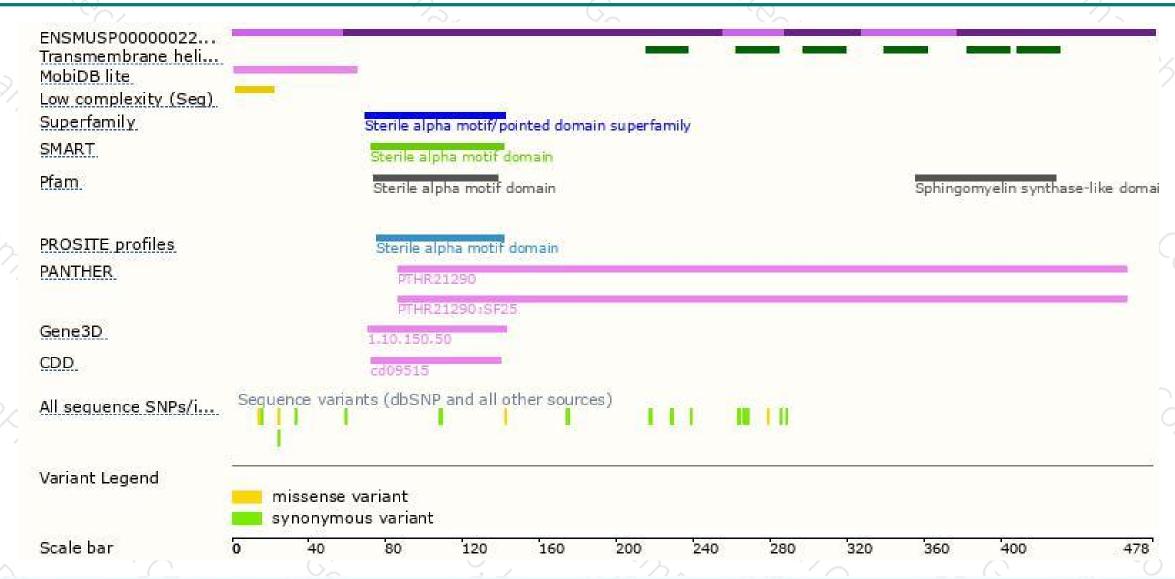
Genomic location distribution





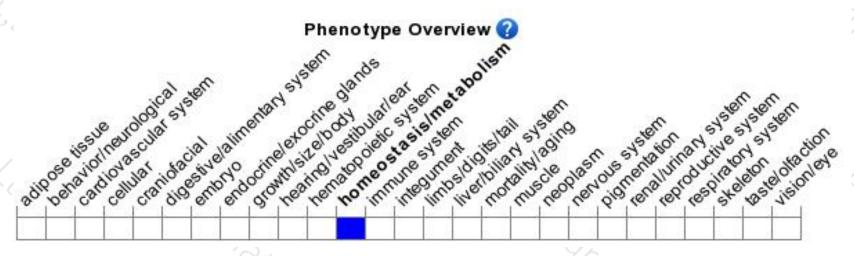
Protein domain





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 decreased ceramide phosphoethanolamine synthase activity but normal liver, kidney and spleen histology.



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





