

Sod1 Cas9-CKO Strategy

Designer:

Daohua Xu

Reviewer:

Huimin Su

Design Date:

2019-9-17

Project Overview



Project Name

Sod1

Project type

Cas9-CKO

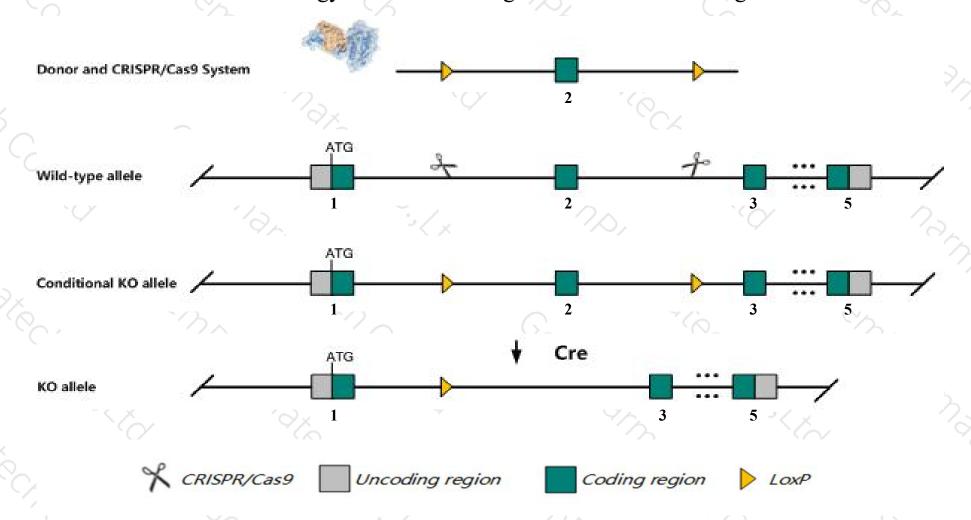
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Sod1 gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Sod1* gene has 2 transcripts. According to the structure of *Sod1* gene, exon2 of *Sod1-201*(ENSMUST00000023707.10) transcript is recommended as the knockout region. The region contains 97bp coding sequence.

 Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Sod1* 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, Homozygous mutants exhibit increased motor neuron loss after axonal injury and enhanced susceptibility to ischemic reperfusion injury. Homozygous females have irregular and small litters, and for some alleles exhibit immature ovarian follicles with few corpora lutea.
- > The *Sod1* gene is located on the Chr16. 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)



Sod1 superoxide dismutase 1, soluble [Mus musculus (house mouse)]

Gene ID: 20655, updated on 31-Jan-2019

Summary

☆ ?

Official Symbol Sod1 provided by MGI

Official Full Name superoxide dismutase 1, soluble provided by MGI

Primary source MGI:MGI:98351

See related Ensembl:ENSMUSG00000022982

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 B430204E11Rik, Cu/Zn-SOD, CuZnSOD, Ipo-1, Ipo1, SODC, Sod-1

Expression Ubiquitous expression in liver adult (RPKM 521.8), liver E14 (RPKM 341.6) and 28 other tissues See more

Orthologs human all

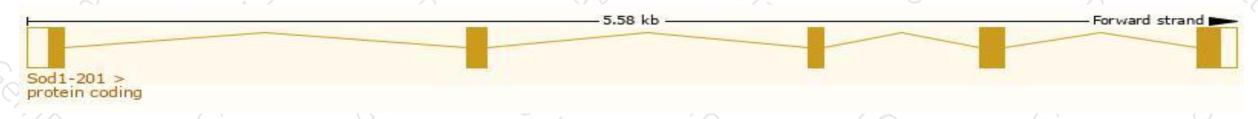
Transcript information (Ensembl)



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

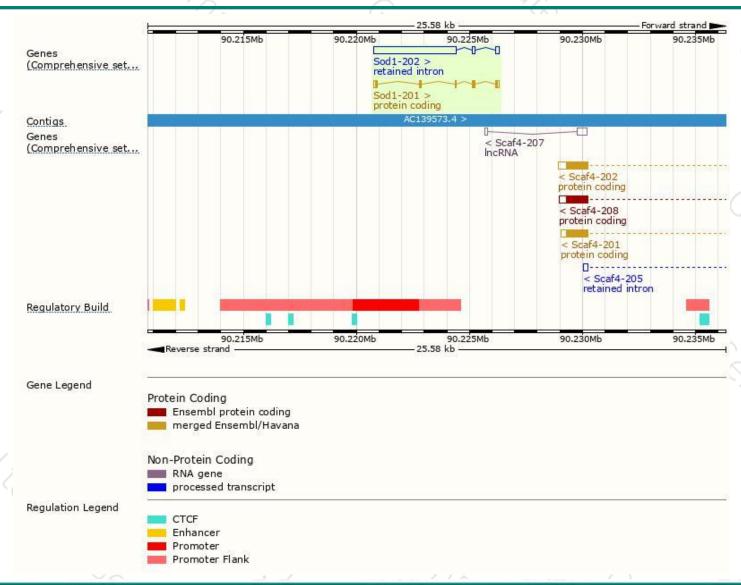
Name 🍦	Transcript ID 👙	bp #	Protein #	Biotype	CCDS	UniProt 🌲	Flags		
Sod1-201	ENSMUST00000023707.10	641	<u>154aa</u>	Protein coding	CCDS37395 ₽	P08228@	TSL:1	GENCODE basic	APPRIS P1
Sod1-202	ENSMUST00000232505.1	3970	No protein	Retained intron			=		

The strategy is based on the design of *Sod1-201* transcript, The transcription is shown below



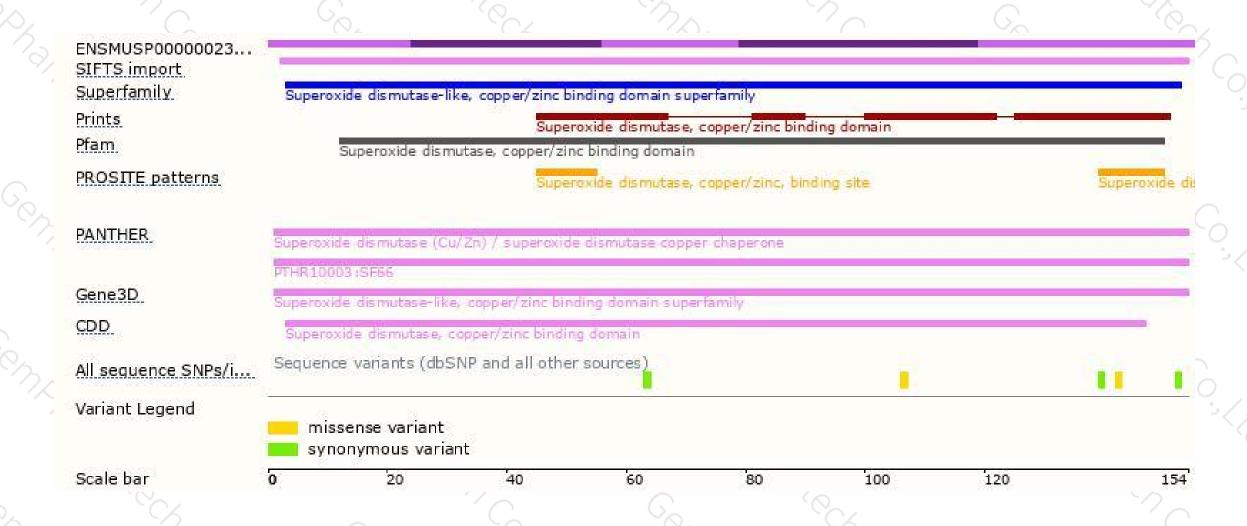
Genomic location distribution





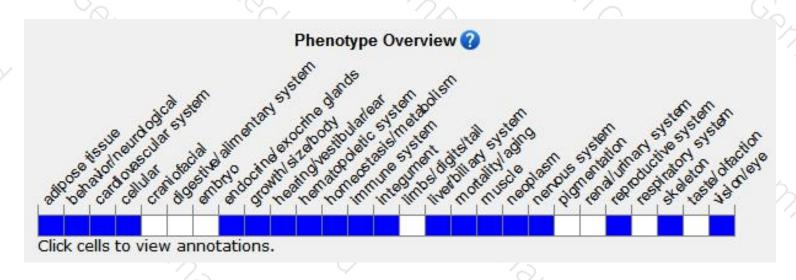
Protein domain





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, Homozygous mutants exhibit increased motor neuron loss after axonal injury and enhanced susceptibility to ischemic reperfusion injury. Homozygous females have irregular and small litters, and for some alleles exhibit immature ovarian follicles with few corpora lutea.



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





