

Rsad2 Cas9-CKO Strategy

Designer: Daohua Xu

Reviewer: Huimin Su

Design Date: 2020-2-11

Project Overview



Project Name Rsad2

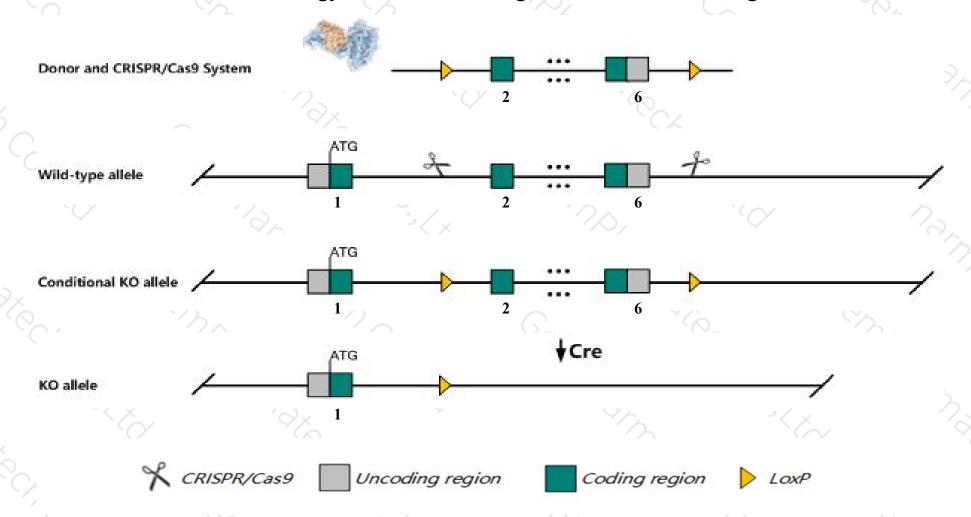
Project type Cas9-CKO

Strain background C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Rsad2 gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Rsad2* gene has 3 transcripts. According to the structure of *Rsad2* gene, exon2-exon6 of *Rsad2-201* (ENSMUST00000020970.9) transcript is recommended as the knockout region. The region contains 740bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Rsad2* 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 null allele exhibit impaired T-helper 2 differentitation.
- > The *Rsad2* gene is located on the Chr12. 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)



Rsad2 radical S-adenosyl methionine domain containing 2 [Mus musculus (house mouse)]

Gene ID: 58185, updated on 31-Jan-2019

Summary

↑ ?

Official Symbol Rsad2 provided by MGI

Official Full Name radical S-adenosyl methionine domain containing 2 provided by MGI

Primary source MGI:MGI:1929628

See related Ensembl: ENSMUSG00000020641

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 2510004L01Rik, Vig1, cig5

Expression Biased expression in liver E18 (RPKM 29.0), spleen adult (RPKM 9.1) and 7 other tissuesSee more

Orthologs <u>human all</u>

Transcript information (Ensembl)



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

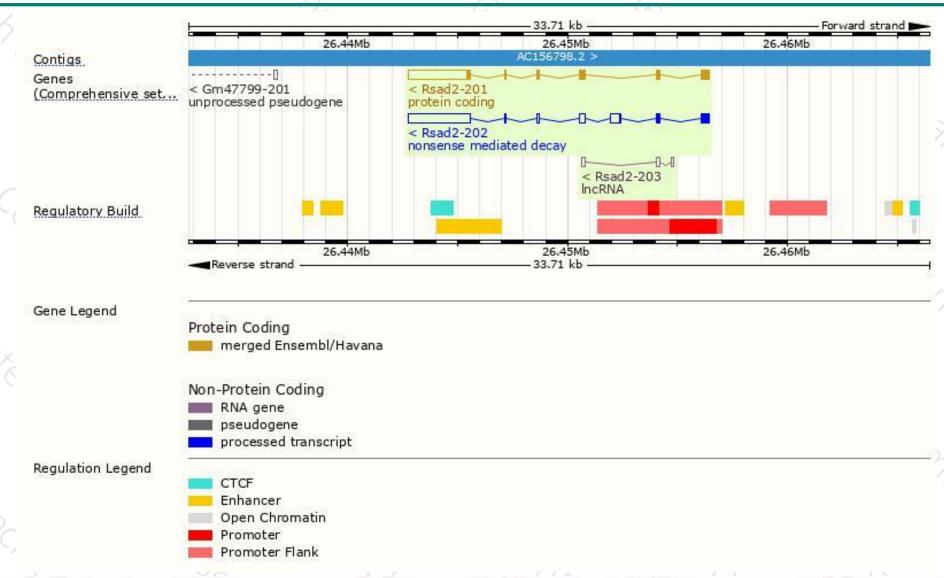
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Rsad2-201	ENSMUST00000020970.9	3782	362aa	Protein coding	CCDS25848	Q8CBB9	TSL:1 GENCODE basic APPRIS P1
Rsad2-202	ENSMUST00000137792.6	4189	<u>187aa</u>	Nonsense mediated decay	5	D6RJ49	TSL:1
Rsad2-203	ENSMUST00000142732.1	430	No protein	IncRNA	ų.	-	TSL:2

The strategy is based on the design of Rsad2-201 transcript, The transcription is shown below



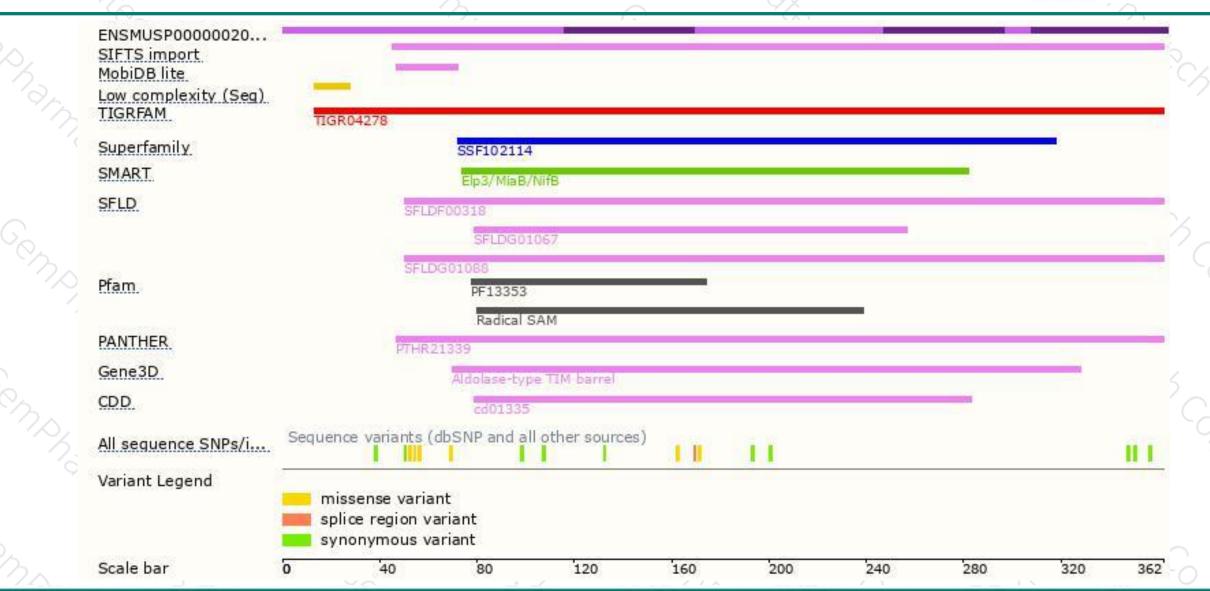
Genomic location distribution





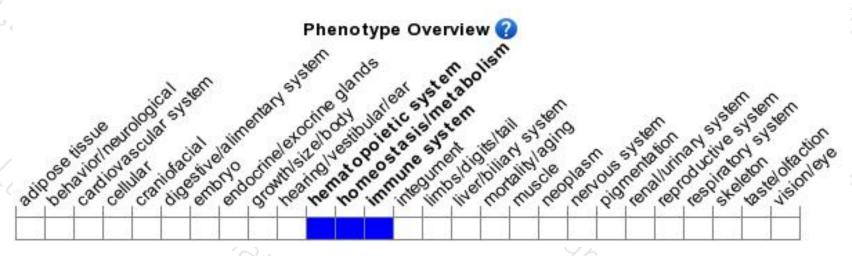
Protein domain





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 null allele exhibit impaired T-helper 2 differentitation.



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





