

Ppp2r5a Cas9-CKO Strategy

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Reviewer: Xiaojing Li

Design Date: 2019-8-30

Project Overview



Project Name

Ppp2r5a

Project type

Cas9-CKO

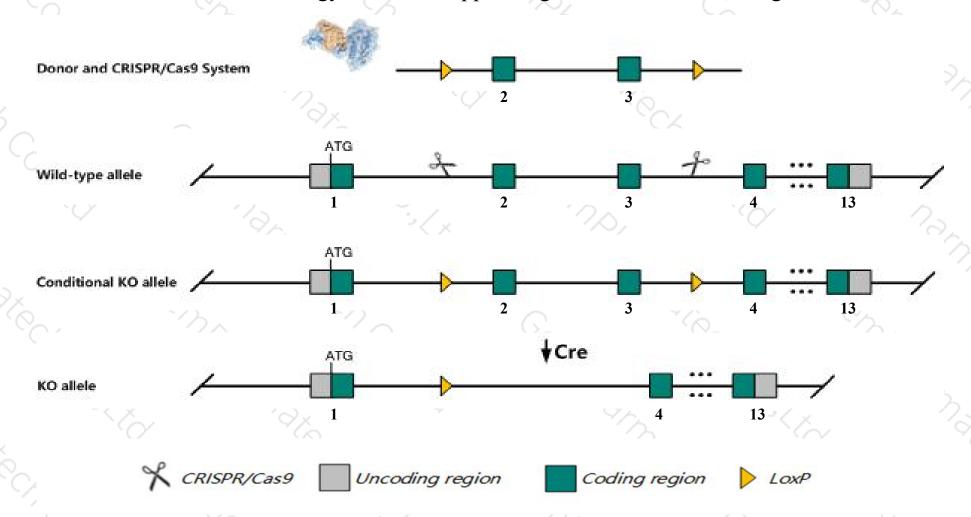
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Ppp2r5a* gene. The schematic diagram is as follows:



Technical routes



- The *Ppp2r5a* gene has 8 transcripts. According to the structure of *Ppp2r5a* gene, exon2-exon3 of *Ppp2r5a-201* (ENSMUST00000067976.8) transcript is recommended as the knockout region. The region contains 299bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Ppp2r5a* 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 gene trap allele are hypomorphic, with the severely reduced expression causing development of spontaneous skin lesions combined with hair loss and enlarged liver, spleen and lymph nodes. They are also more susceptible to chemically induced skin tumors.
- The *Ppp2r5a* gene is located on the Chr1. 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)



Ppp2r5a protein phosphatase 2, regulatory subunit B', alpha [Mus musculus (house mouse)]

Gene ID: 226849, updated on 12-Aug-2019

Summary

△ ?

Official Symbol Ppp2r5a provided by MGI

Official Full Name protein phosphatase 2, regulatory subunit B', alpha provided by MGI

Primary source MGI:MGI:2388479

See related Ensembl: ENSMUSG00000026626

Gene type protein coding
RefSeq status PROVISIONAL
Organism Mus musculus

Lineage Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;

Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus

Also known as PR61alpha

Expression Ubiquitous expression in mammary gland adult (RPKM 92.3), subcutaneous fat pad adult (RPKM 91.4) and 27 other tissues See more

Orthologs human all

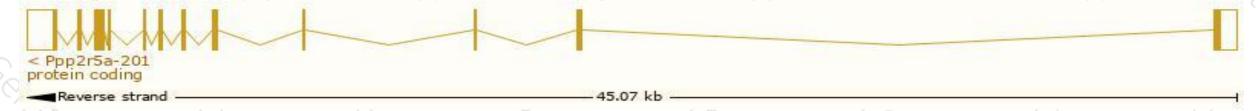
Transcript information (Ensembl)



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

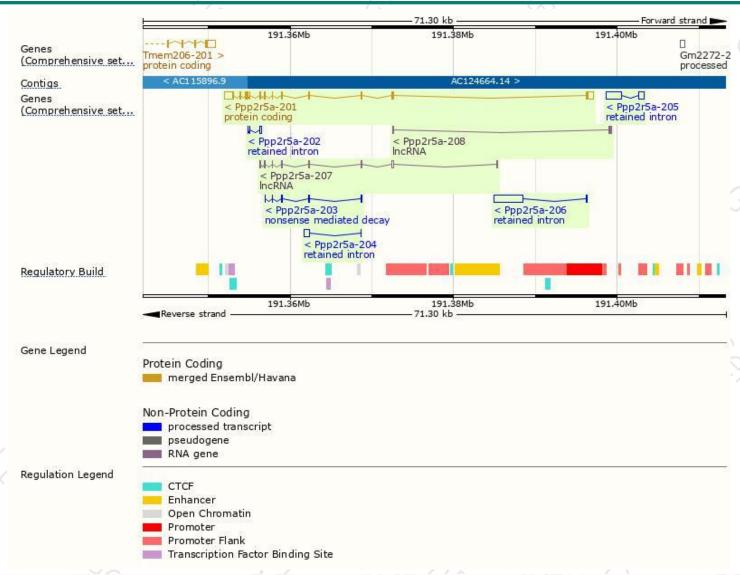
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Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Ppp2r5a-201	ENSMUST00000067976.8	3088	486aa	Protein coding	CCDS15620	Q6PD03	TSL:1 GENCODE basic APPRIS P1
Ppp2r5a-203	ENSMUST00000191925.1	429	<u>60aa</u>	Nonsense mediated decay		A0A0A6YVV9	CDS 5' incomplete TSL:3
Ppp2r5a-206	ENSMUST00000194953.1	3541	No protein	Retained intron	142	2	TSL:1
Ppp2r5a-205	ENSMUST00000194832.1	2421	No protein	Retained intron	82	<u> </u>	TSL:1
Ppp2r5a-204	ENSMUST00000192334.1	706	No protein	Retained intron			TSL:3
Ppp2r5a-202	ENSMUST00000191920.1	240	No protein	Retained intron		*	TSL:3
Ppp2r5a-207	ENSMUST00000195605.5	779	No protein	IncRNA	142	2	TSL:3
Ppp2r5a-208	ENSMUST00000195859.1	297	No protein	IncRNA	14	20	TSL:3

The strategy is based on the design of *Ppp2r5a-201* transcript, The transcription is shown below



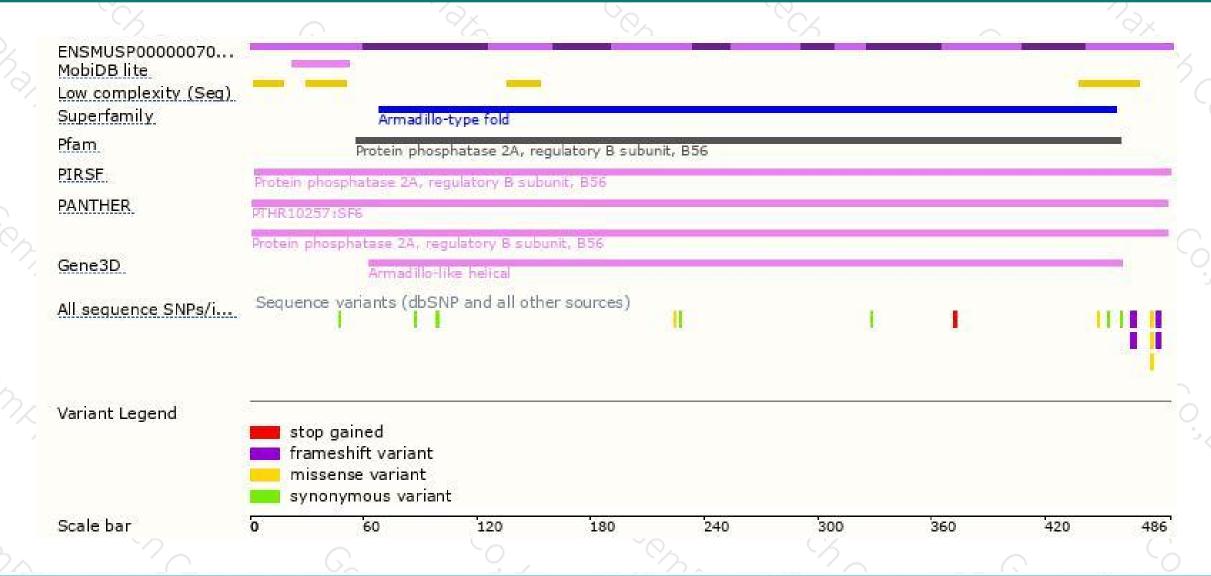
Genomic location distribution





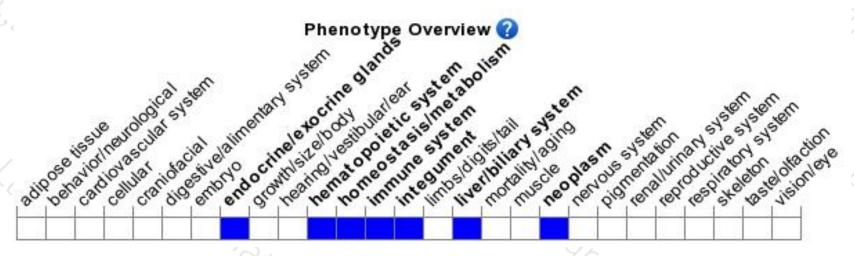
Protein domain





Mouse phenotype description(MGI)





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

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





