

Ptpn2 Cas9-KO Strategy To hall alto color color

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Project Overview



Project Name

Ptpn2

Project type

Cas9-KO

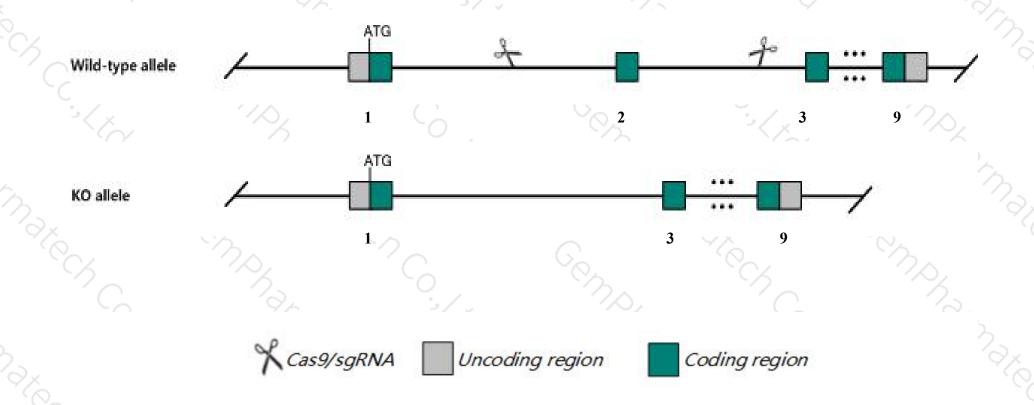
Strain background

C57BL/6J

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Ptpn2* gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Ptpn2* gene has 4 transcripts. According to the structure of *Ptpn2* gene, exon2 of *Ptpn2-203*(ENSMUST00000122412.1) transcript is recommended as the knockout region. The region contains 91bp coding sequence.

 Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Ptpn2* gene. The brief process is as follows: sgRNA was transcribed in vitro.Cas9 and sgRNA were microinjected into the fertilized eggs of C57BL/6J 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/6J mice.

Notice



- ➤ According to the existing MGI data, Mice homozygous for disruptions in this gene have a reduced life span, abnormalities of the hematopoietic system and an increased succeptibility to inflammatory disease.
- ➤ The *Ptpn2* gene is located on the Chr18. 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 gene transcription and translation processes, all risks cannot be predicted under existing information.

Gene information (NCBI)



Ptpn2 protein tyrosine phosphatase, non-receptor type 2 [Mus musculus (house mouse)]

Gene ID: 19255, updated on 12-Mar-2019

Summary

☆ ?

Official Symbol Ptpn2 provided by MGI

Official Full Name protein tyrosine phosphatase, non-receptor type 2 provided by MGI

Primary source MGI:MGI:97806

See related Ensembl: ENSMUSG00000024539

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 Al325124, Ptpt, TC-PTP

Expression Ubiquitous expression in CNS E11.5 (RPKM 3.3), CNS E14 (RPKM 3.0) and 27 other tissuesSee more

Orthologs <u>human</u> all

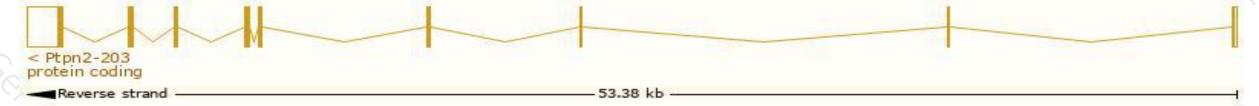
Transcript information (Ensembl)



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

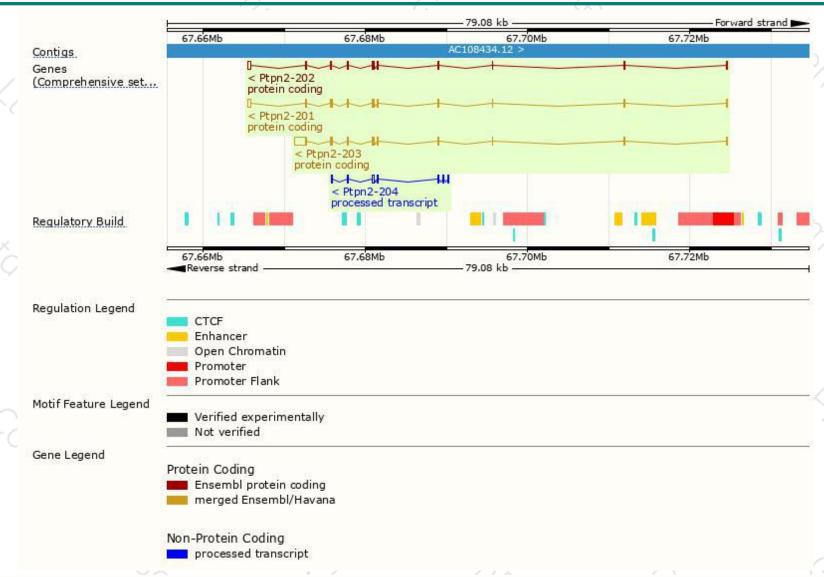
Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
ENSMUST00000122412.1	2667	<u>406aa</u>	Protein coding	CCDS50311	Q06180	TSL:1 GENCODE basic APPRIS ALT2
ENSMUST00000025420.13	1568	382aa	Protein coding	CCDS37851	Q06180	TSL:1 GENCODE basic APPRIS P3
ENSMUST00000120934.7	1497	363aa	Protein coding	49	D3Z6W2	TSL:5 GENCODE basic APPRIS ALT2
ENSMUST00000128169.1	760	No protein	Processed transcript	29	20	TSL:3
	ENSMUST00000122412.1 ENSMUST00000025420.13 ENSMUST00000120934.7	ENSMUST00000122412.1 2667 ENSMUST00000025420.13 1568 ENSMUST00000120934.7 1497	ENSMUST00000122412.1 2667 406aa ENSMUST00000025420.13 1568 382aa ENSMUST00000120934.7 1497 363aa	ENSMUST00000122412.1 2667 406aa Protein coding ENSMUST00000025420.13 1568 382aa Protein coding ENSMUST00000120934.7 1497 363aa Protein coding	ENSMUST00000122412.1 2667 406aa Protein coding CCDS50311 ENSMUST00000025420.13 1568 382aa Protein coding CCDS37851 ENSMUST00000120934.7 1497 363aa Protein coding -	ENSMUST00000122412.1 2667 406aa Protein coding CCDS50311 Q06180 ENSMUST00000025420.13 1568 382aa Protein coding CCDS37851 Q06180 ENSMUST00000120934.7 1497 363aa Protein coding - D3Z6W2

The strategy is based on the design of *Ptpn2-203* transcript, The transcription is shown below



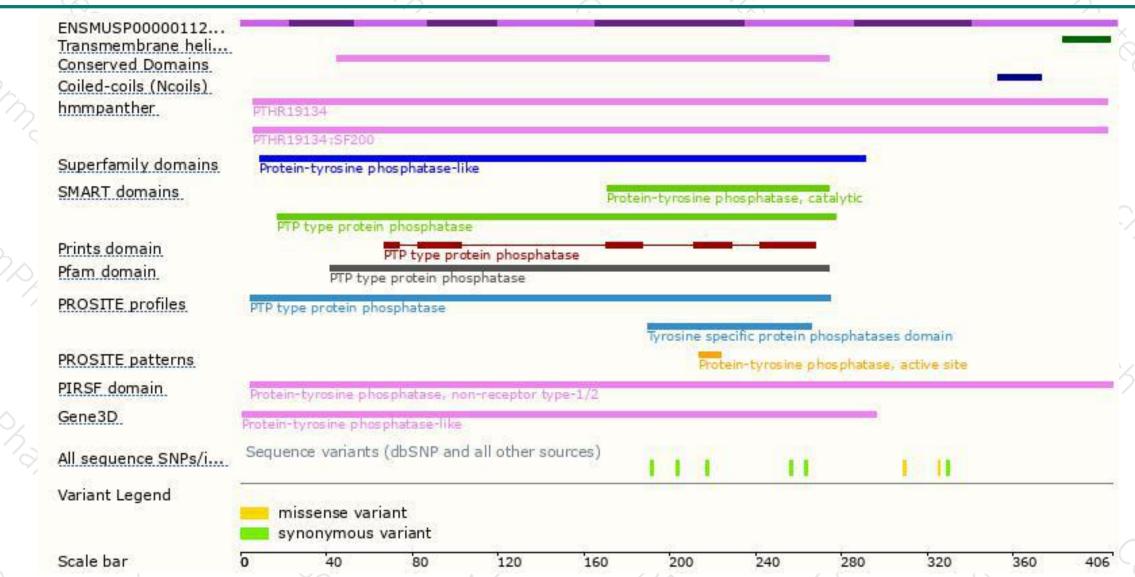
Genomic location distribution





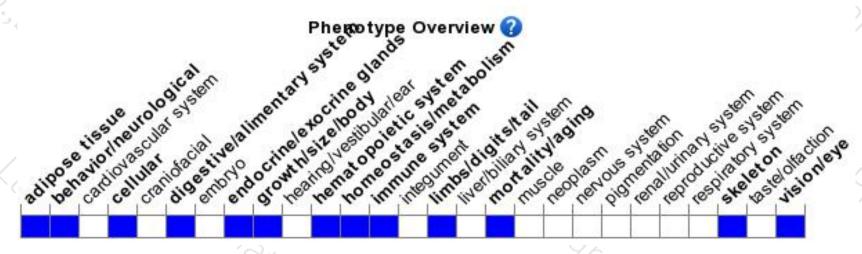
Protein domain





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 disruptions in this gene have a reduced life span, abnormalities of the hematopoietic system and an increased succeptibility to inflammatory disease.



If you have any questions, you are welcome to inquire.

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