

Ptpn5 Cas9-CKO Strategy

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Project Overview



Project Name

Ptpn5

Project type

Cas9-CKO

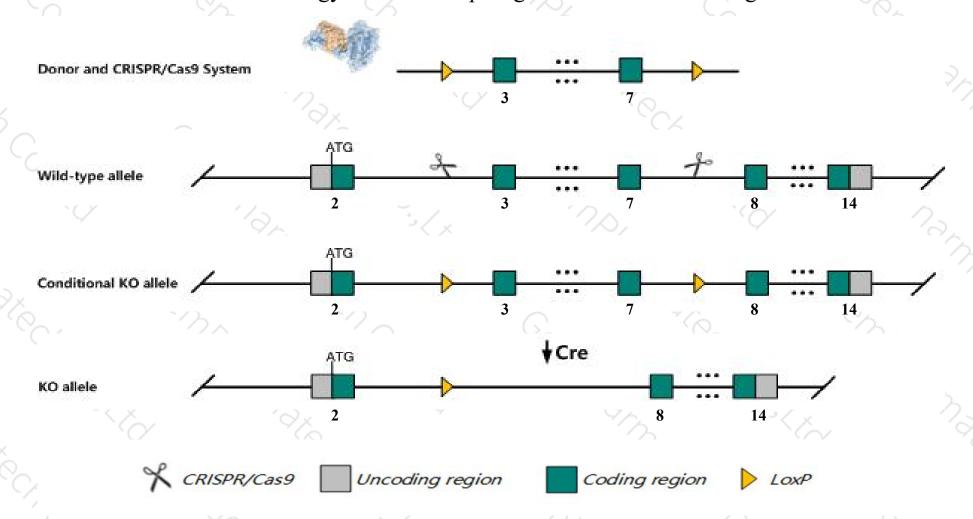
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Ptpn5* gene. The schematic diagram is as follows:



Technical routes



- The *Ptpn5* gene has 11 transcripts. According to the structure of *Ptpn5* gene, exon3-exon7 of *Ptpn5-201* (ENSMUST00000033142.12) transcript is recommended as the knockout region. The region contains 818bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Ptpn5* 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 normal brain development.
- > The *Ptpn5* gene is located on the Chr7. 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)



Ptpn5 protein tyrosine phosphatase, non-receptor type 5 [Mus musculus (house mouse)]

Gene ID: 19259, updated on 17-Sep-2019

Summary



Official Symbol Ptpn5 provided by MGI

Official Full Name protein tyrosine phosphatase, non-receptor type 5 provided by MGI

Primary source MGI:MGI:97807

> Ensembl: ENSMUSG00000030854 See related

Gene type protein coding RefSeg status VALIDATED Organism Mus musculus

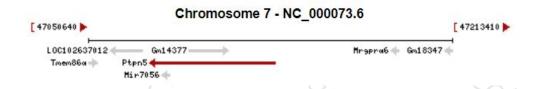
> Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Lineage

Muroidea; Muridae; Murinae; Mus; Mus

Also known as

Biased expression in CNS E18 (RPKM 44.4), whole brain E14.5 (RPKM 40.6) and 9 other tissues See more

Orthologs human all



Transcript information (Ensembl)



The gene has 11 transcripts, all transcripts are shown below:

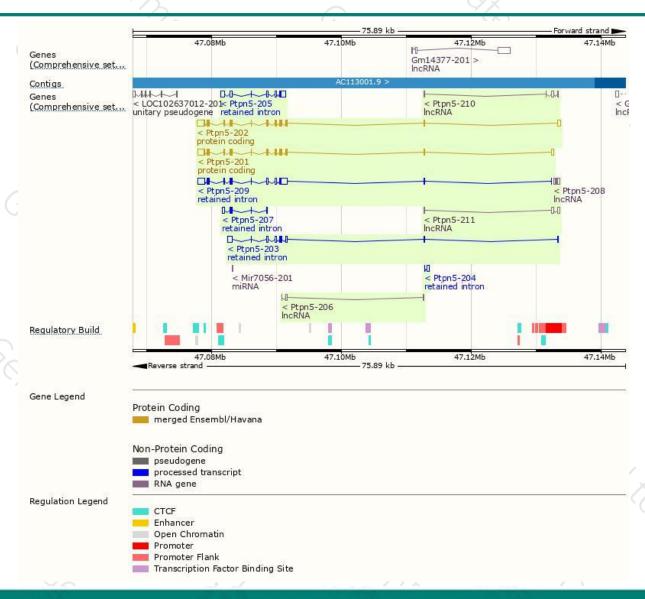
Name 🍦	Transcript ID #	bp 🛊	Protein	Translation ID	Biotype	CCDS	UniProt	Flags		
Ptpn5-201	ENSMUST00000033142.12	3145	<u>541aa</u>	ENSMUSP00000033142.5	Protein coding	CCDS21294©	P54830₽	TSL:1	GENCODE basic	APPRIS P1
Ptpn5-202	ENSMUST00000102626.9	3137	541aa	ENSMUSP00000099686.1	Protein coding	CCDS21294@	P54830@	TSL:1	GENCODE basic	APPRIS P1
Ptpn5-209	ENSMUST00000209161.1	3296	No protein	2	Retained intron	-	ূ	TSL:2		
Ptpn5-205	ENSMUST00000208324.1	2165	No protein	-	Retained intron	(F 4 5)	*	TSL:1		
tpn5-203	ENSMUST00000207172.1	1874	No protein	-	Retained intron	2.52	-	TSL:2		
tpn5-207	ENSMUST00000208531.1	694	No protein	2.	Retained intron	123	2	TSL:3		
tpn5-204	ENSMUST00000207344.1	393	No protein		Retained intron	(+ 2)		TSL:3		
otpn5-211	ENSMUST00000209184.1	765	No protein	1.53	IncRNA	15 <u>T</u> 16		TSL:3		
tpn5-210	ENSMUST00000209179.1	673	No protein	2	IncRNA	848	-	TSL:3		
tpn5-208	ENSMUST00000209057.1	589	No protein	. 50	IncRNA	S 1 89	*	TSL:3		
Ptpn5-206	ENSMUST00000208437.1	348	No protein	- 2	IncRNA	-	2	TSL:3		

The strategy is based on the design of *Ptpn5-201* transcript, The transcription is shown below



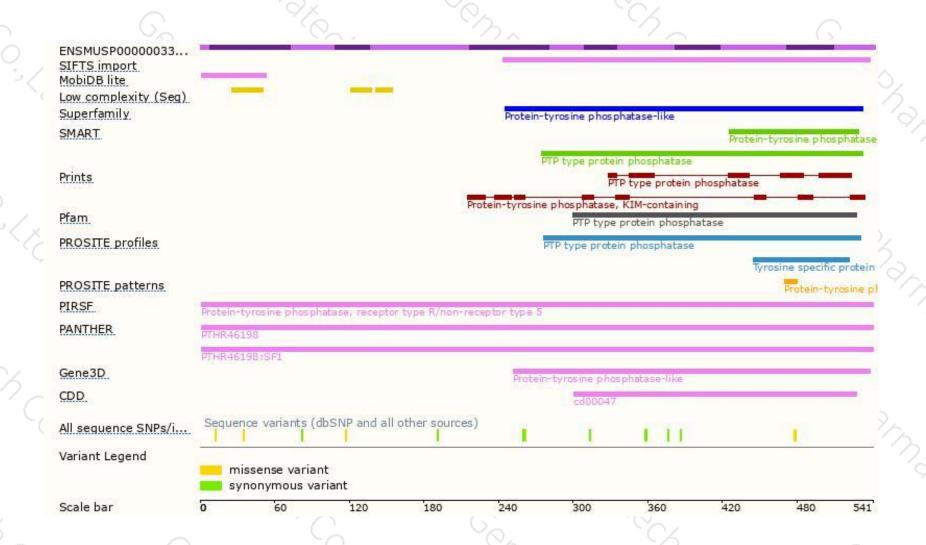
Genomic location distribution





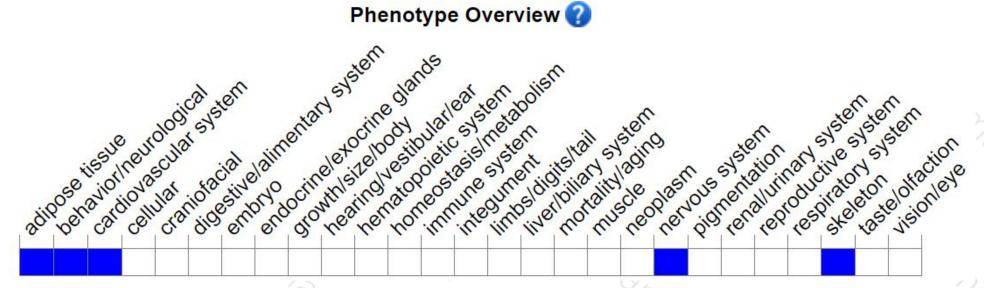
Protein domain





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 normal brain development.



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





