

# Ptk2 Cas9-KO Strategy

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**Design Date:** 2019-8-3

## **Project Overview**



Project Name Ptk2

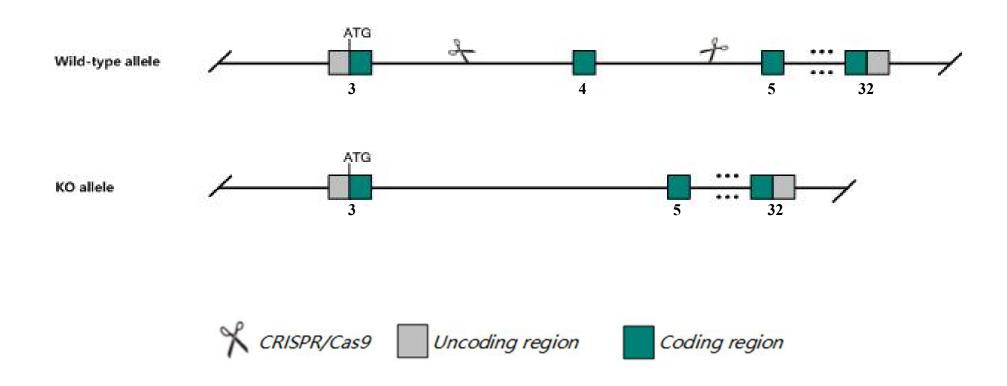
Project type Cas9-KO

Strain background C57BL/6JGpt

## **Knockout strategy**



This model will use CRISPR/Cas9 technology to edit the *Ptk2* gene. The schematic diagram is as follows:



### **Technical routes**



The Ptk2 gene has 15 transcripts. According to the structure of Ptk2 gene, exon4 of Ptk2-201

(ENSMUST00000110036.10) transcript is recommended as the knockout region. The region contains 167bp coding sequence.

Knock out the region will result in disruption of protein function.

In this project we use CRISPR/Cas9 technology to modify Ptk2 gene. The brief process is as follows: CRISPR/Cas9 system v

### **Notice**



According to the existing MGI data, Mice homozygous for a null allele die before or during organogenesis with growth retardation, abnormal embryonic and extra embryonic tissue development, and abnormal vascular development.

The *Ptk2* gene is located on the Chr15. 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 the gene knockout on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.

## Gene information NCBI



#### Ptk2 PTK2 protein tyrosine kinase 2 [Mus musculus (house mouse)]

Gene ID: 14083, updated on 2-Apr-2019

#### Summary

☆ ?

Official Symbol Ptk2 provided by MGI

Official Full Name PTK2 protein tyrosine kinase 2 provided by MGI

Primary source MGI:MGI:95481

See related Ensembl:ENSMUSG00000022607

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 FADK 1, FAK, FRNK, Fadk, p125FAK

Expression Ubiquitous expression in cortex adult (RPKM 12.3), CNS E11.5 (RPKM 11.8) and 28 other tissuesSee more

Orthologs <u>human</u> all

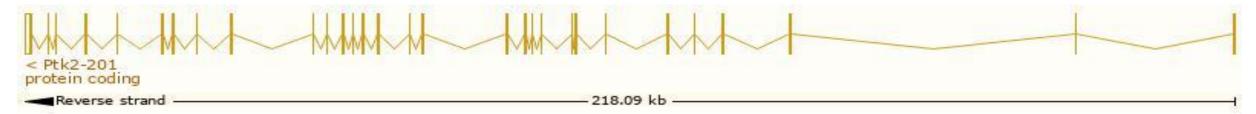
## Transcript information Ensembl



#### The gene has 15 transcripts, all transcripts are shown below:

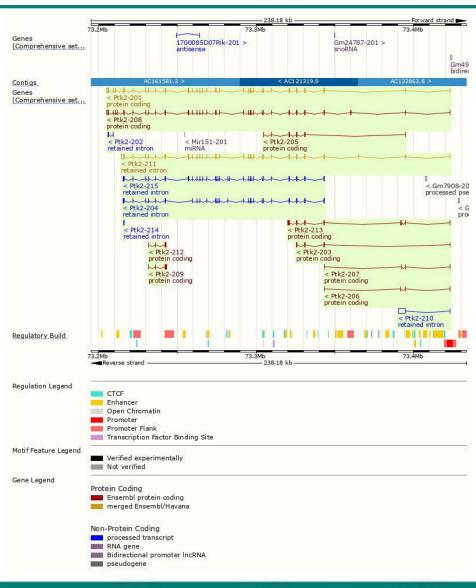
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Ptk2-201	ENSMUST00000110036.10	4417	1052aa	Protein coding	CCDS37099	P34152	TSL:1 GENCODE basic APPRIS P2
Ptk2-208	ENSMUST00000226988.2	4675	1055aa	Protein coding	-	P34152	GENCODE basic APPRIS ALT2
Ptk2-213	ENSMUST00000228180.2	1333	198aa	Protein coding	020	P34152	GENCODE basic
Ptk2-205	ENSMUST00000226791.1	517	<u>119aa</u>	Protein coding	72.5	A0A2I3BRG7	CDS 3' incomplete
Ptk2-209	ENSMUST00000227395.1	498	81aa	Protein coding	151	A0A2I3BR46	CDS 3' incomplete
Ptk2-203	ENSMUST00000226466.1	488	<u>126aa</u>	Protein coding	16.0	A0A2I3BQ85	CDS 3' incomplete
Ptk2-212	ENSMUST00000227686.1	485	114aa	Protein coding	(2)	A0A2I3BPK4	CDS 3' incomplete
Ptk2-206	ENSMUST00000226848.1	364	14aa	Protein coding	725	Q2VQT9	CDS 3' incomplete
Ptk2-207	ENSMUST00000226893.1	318	<u>14aa</u>	Protein coding	1.5	Q2VQT9	CDS 3' incomplete
Ptk2-211	ENSMUST00000227569.1	4678	No protein	Retained intron	-		
Ptk2-210	ENSMUST00000227435.1	4580	No protein	Retained intron	(2)	-	
Ptk2-215	ENSMUST00000228628.1	3254	No protein	Retained intron	72.5	2	
Ptk2-204	ENSMUST00000226742.1	2820	No protein	Retained intron	1.5	-	
Ptk2-202	ENSMUST00000226454.1	542	No protein	Retained intron			
Ptk2-214	ENSMUST00000228457.1	259	No protein	Retained intron	0.20	-	

The strategy is based on the design of *Ptk2-201* transcript, The transcription is shown below



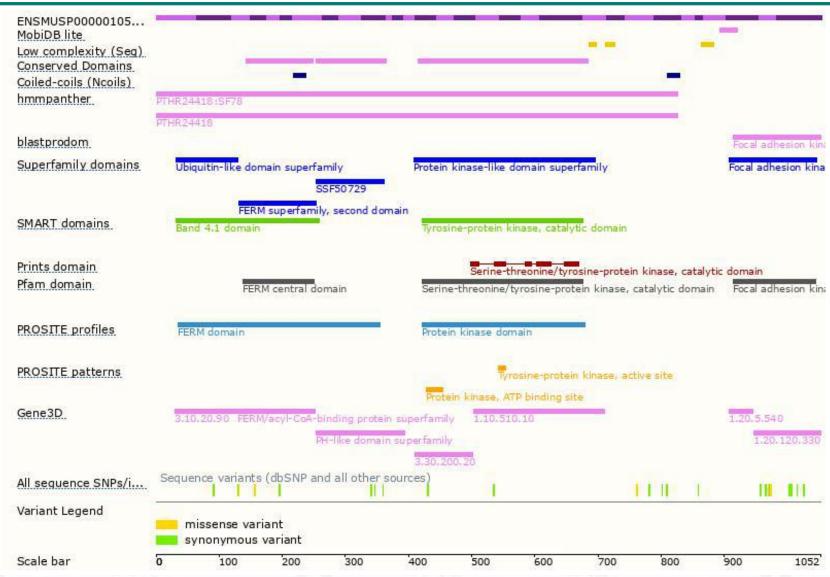
### Genomic location distribution





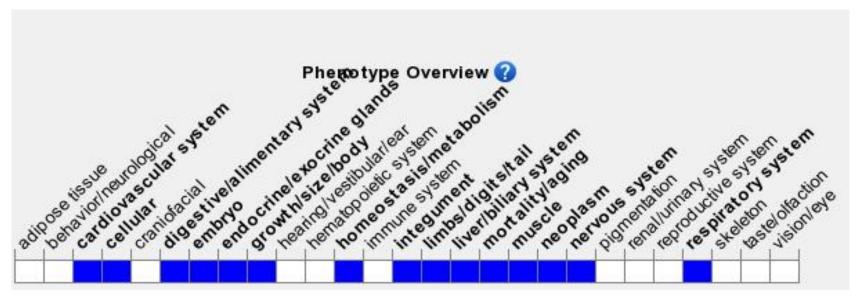
### Protein domain





## 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 die before or during organogenesis with growth retardation, abnormal embryonic and extra embryonic tissue development, and abnormal vascular development.



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





