

Pdzd7 Cas9-CKO Strategy

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Project Overview



Project Name

Pdzd7

Project type

Cas9-CKO

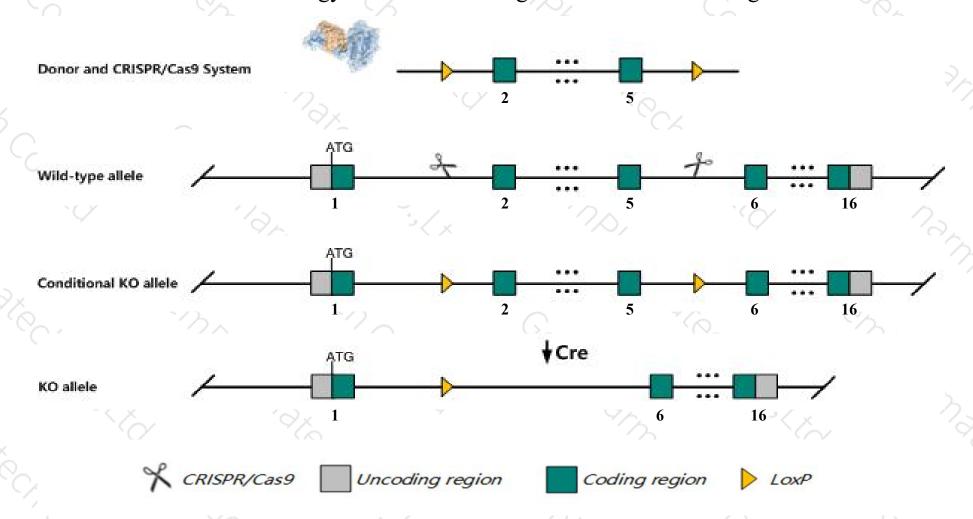
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Pdzd7* gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Pdzd7* gene has 8 transcripts. According to the structure of *Pdzd7* gene, exon2-exon5 of *Pdzd7-203*(ENSMUST00000169459.3) transcript is recommended as the knockout region. The region contains 641bp coding sequence.

 Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Pdzd7* 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 knock-out allele exhibit profound deafness due to abnormal outer cochlear hair cell morphology and function.
- The floxed region is near to the N-terminal of *Sfxn3* gene, this strategy may influence the regulatory function of the N-terminal of *Sfxn3* gene.
- ightharpoonup Transcript Pdzd7-201 may not be affected.
- The *Pdzd7* gene is located on the Chr19. 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)



Pdzd7 PDZ domain containing 7 [Mus musculus (house mouse)]

Gene ID: 100503041, updated on 10-Oct-2019

Summary

△ ?

Official Symbol Pdzd7 provided by MGI

Official Full Name PDZ domain containing 7 provided by MGI

Primary source MGI:MGI:3608325

See related Ensembl:ENSMUSG00000074818

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 Pdzk7; EG435601; 9130207N01

Expression Biased expression in adrenal adult (RPKM 18.9), colon adult (RPKM 4.4) and 9 other tissues See more

Orthologs human all

Genomic context



Location: 19; 19 C3

See Pdzd7 in Genome Data Viewer

Exon count: 21

Annotation release	Status	Assembly	Chr	Location	
108	current	GRCm38.p6 (GCF_000001635.26)	19	NC_000085.6 (4502690745058633, complement)	
Build 37.2	previous assembly	MGSCv37 (GCF_000001635.18)	19	NC_000085.5 (4510210545120133, complement)	

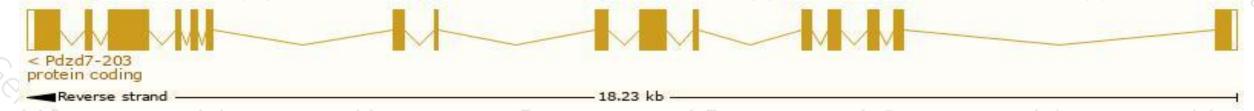
Transcript information (Ensembl)



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

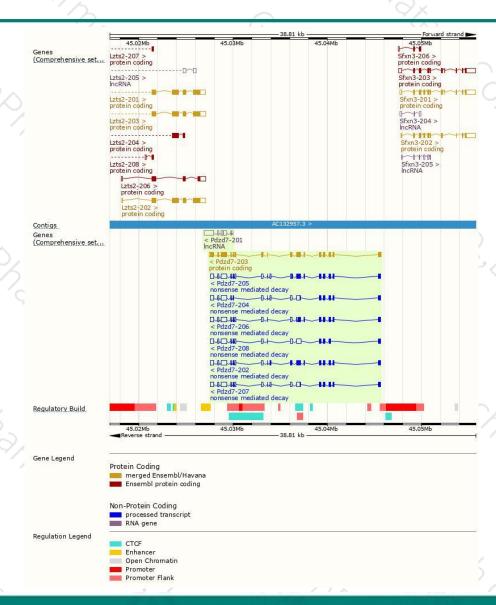
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Pdzd7-203	ENSMUST00000169459.3	3265	<u>1021aa</u>	Protein coding	CCDS57145	E9Q9W7	TSL:5 GENCODE basic APPRIS P1
Pdzd7-205	ENSMUST00000237227.1	3190	<u>476aa</u>	Nonsense mediated decay	ä	-	
Pdzd7-202	ENSMUST00000145391.8	3161	<u>553aa</u>	Nonsense mediated decay	-	E9Q9W7	TSL:5
Pdzd7-207	ENSMUST00000237962.1	3100	289aa	Nonsense mediated decay	2	-	
Pdzd7-204	ENSMUST00000237077.1	3008	289aa	Nonsense mediated decay	ē	-	
Pdzd7-206	ENSMUST00000237833.1	2981	409aa	Nonsense mediated decay	ë	-	
Pdzd7-208	ENSMUST00000238074.1	2975	245aa	Nonsense mediated decay	-	X4ZEG9	
Pdzd7-201	ENSMUST00000038901.5	1799	No protein	IncRNA	<u>_</u>		TSL:1

The strategy is based on the design of Pdzd7-203 transcript, The transcription is shown below



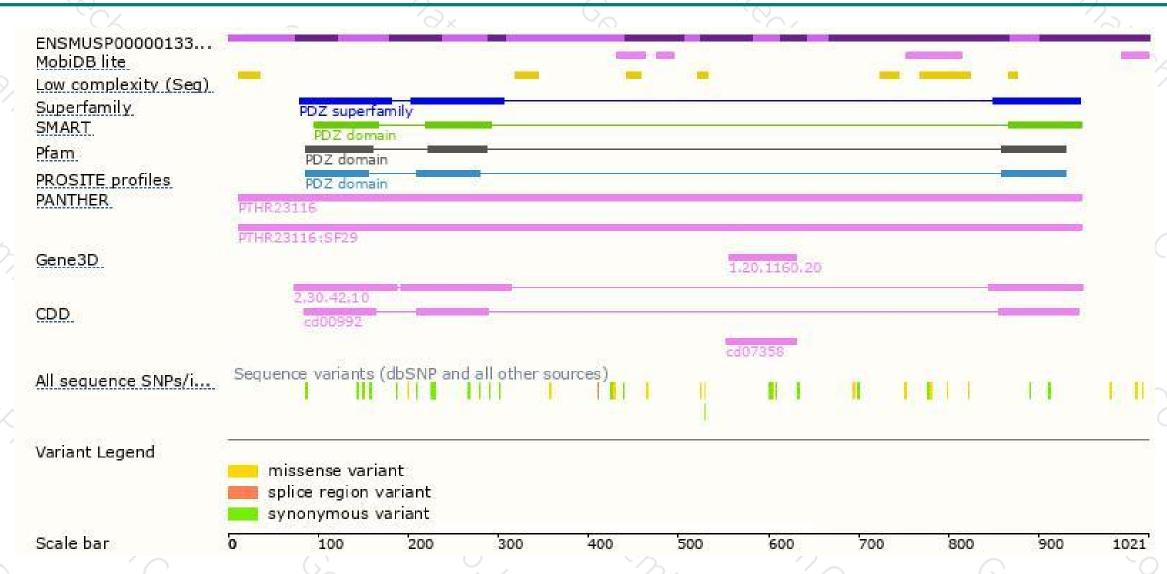
Genomic location distribution





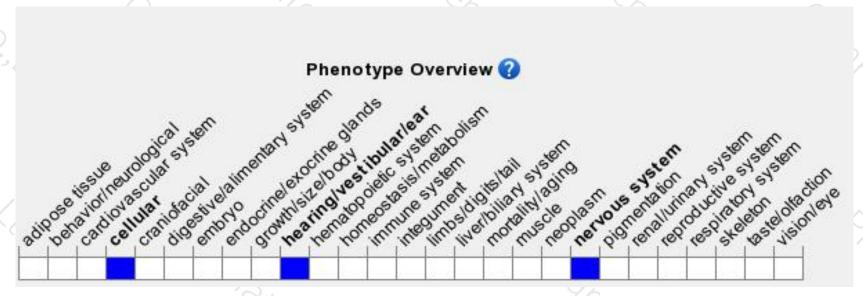
Protein domain





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 a knock-out allele exhibit profound deafness due to abnormal outer cochlear hair cell morphology and function.



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





