

# ***Pdzd7*** **Cas9-KO Strategy**

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Date: 2020-02-20

# Project Overview

**Project Name**

***Pdzd7***

**Project type**

**Cas9-KO**

**Strain background**

**C57BL/6JGpt**

# Knockout strategy

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



- 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

- 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 knockout region is near to the N-terminal of *Sfxn3* gene, this strategy may influence the regulatory function of the N-terminal of *Sfxn3* gene.
- 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 the gene knockout on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology 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

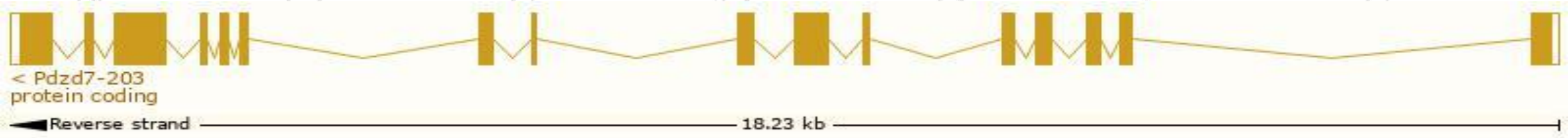
Annotation release	Status	Assembly	Chr	Location
<a href="#">108</a>	current	GRCm38.p6 ( <a href="#">GCF_000001635.26</a> )	19	NC_000085.6 (45026907..45058633, complement)
Build 37.2	previous assembly	MGSCv37 ( <a href="#">GCF_000001635.18</a> )	19	NC_000085.5 (45102105..45120133, complement)

# Transcript information (Ensembl)

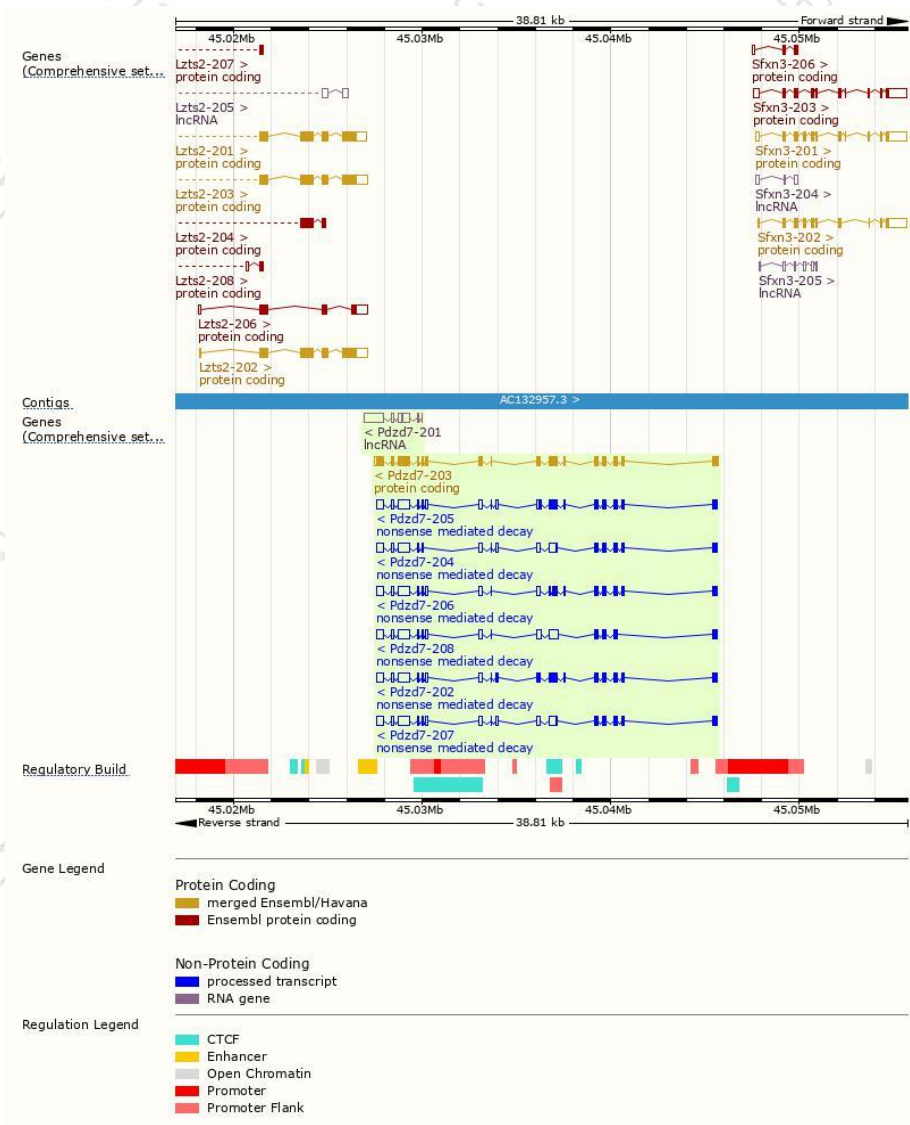
The gene has 8 transcripts,all transcripts are shown below:

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

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

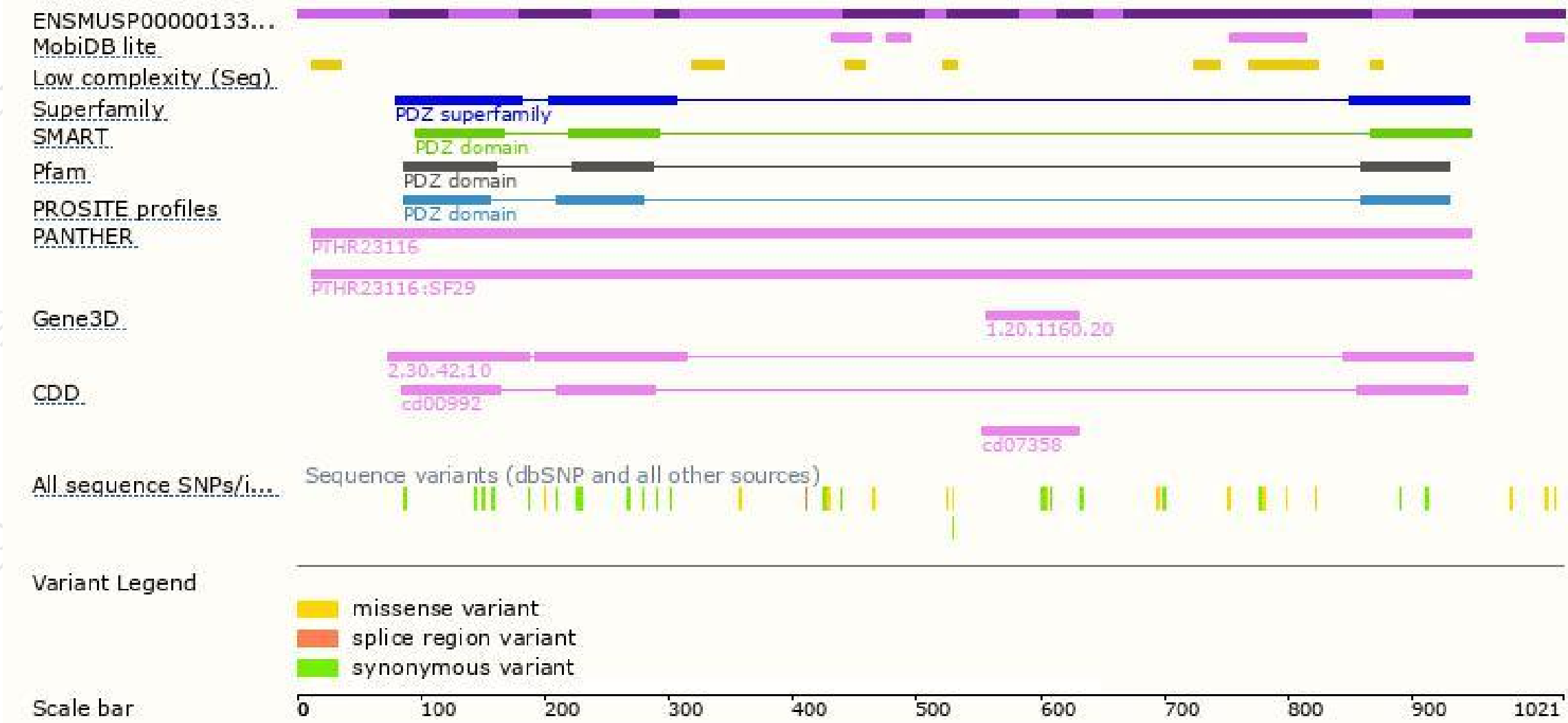


# Genomic location distribution

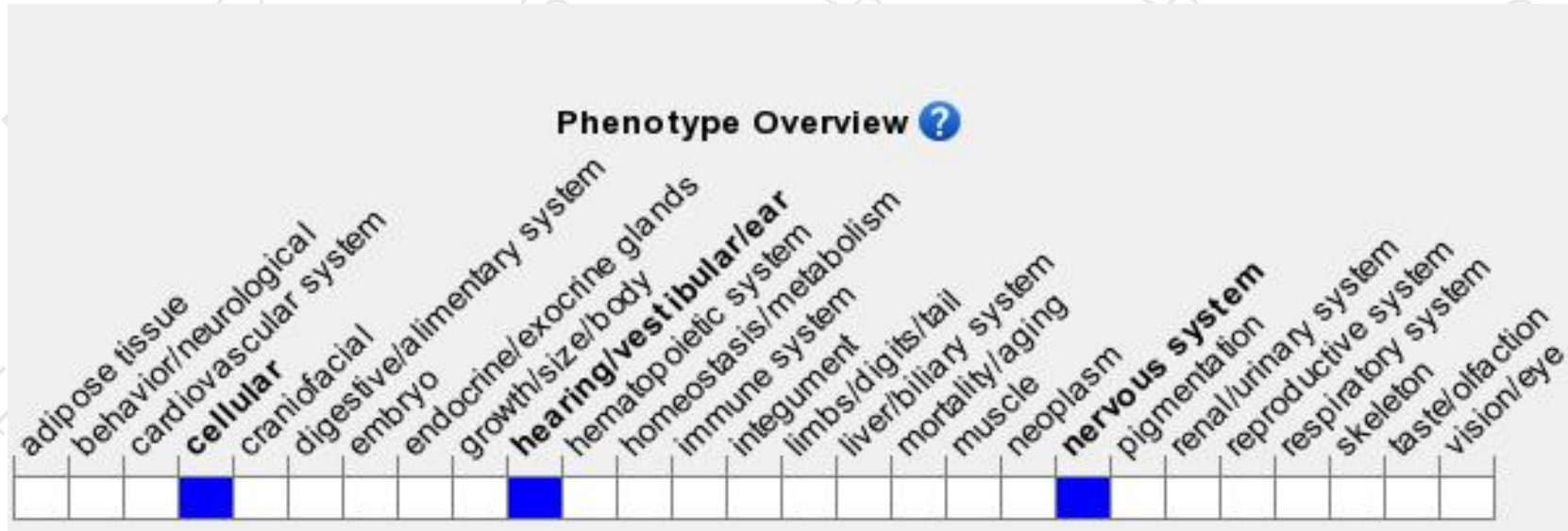




# Protein domain



# 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.

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