

# **Pdlim7 Cas9-CKO Strategy**

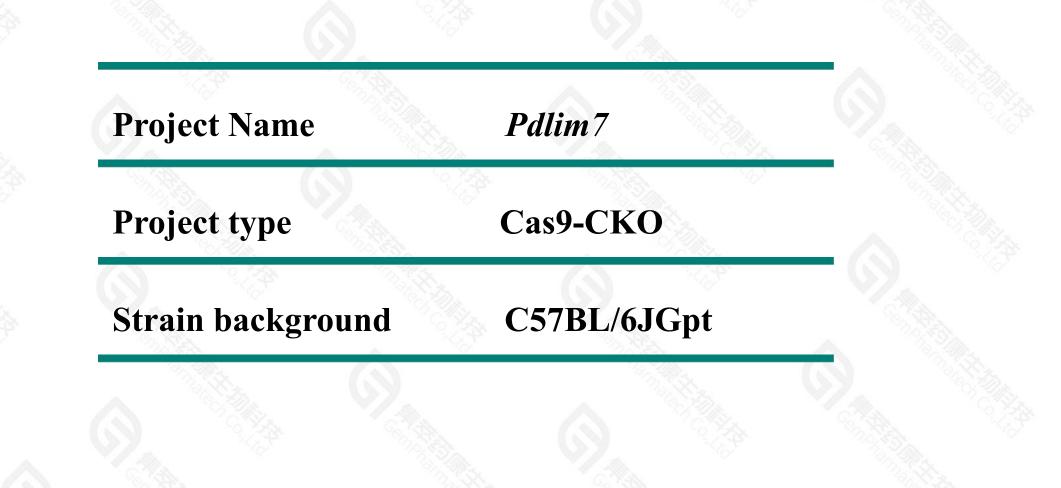
**Designer: Daohua Xu** 

**Reviewer: Xueting Zhang** 

**Design Date: 2021-5-21** 

# **Project Overview**



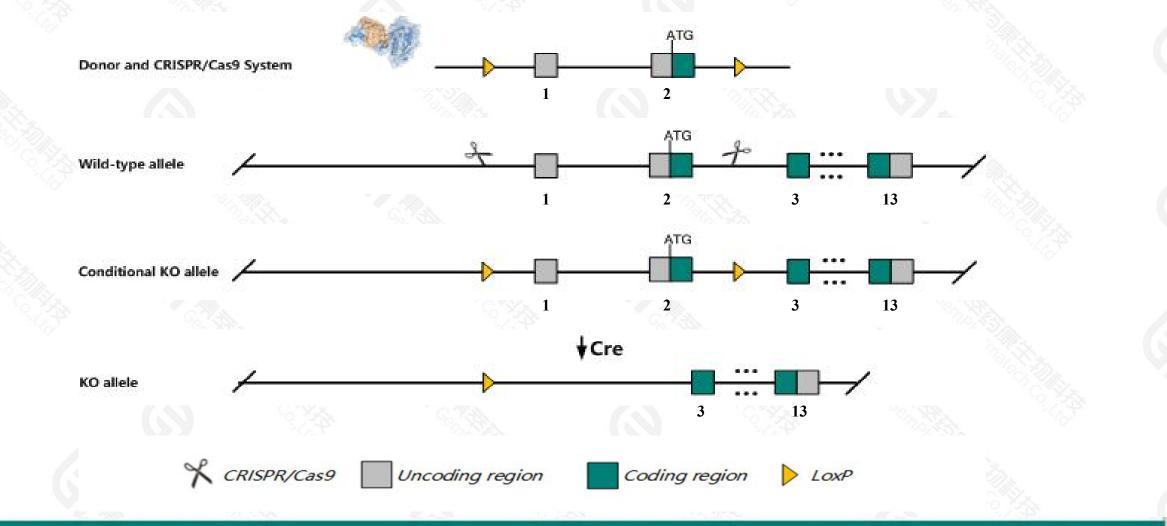


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# **Conditional Knockout strategy**

This model will use CRISPR/Cas9 technology to edit the *Pdlim7* gene. The schematic diagram is as follows:



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The Pdlim7 gene has 11 transcripts. According to the structure of Pdlim7 gene, exon1-exon2 of Pdlim7-201(ENSMUST00000046246.13) transcript is recommended as the knockout region. The region contains start codon ATG coding sequence. Knock out the region will result in disruption of protein function.

➤ In this project we use CRISPR/Cas9 technology to modify *Pdlim7* 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.

 $\succ$  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.



- > According to the existing MGI data, mice homozygous for a gene trap allele exhibit heart defects and hemostatic dysfunction.
- KO region is close to the 5-terminal of *Mir6945* gene, which may affect the 5-terminal regulation of *Mir6945* gene.
  The *Pdlim7* gene is located on the Chr13. 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)**



☆ ?

### Pdlim7 PDZ and LIM domain 7 [Mus musculus (house mouse)]

Gene ID: 67399, updated on 31-Jan-2019

#### Summary

Official Symbol	Pdlim7 provided by MGI
Official Full Name	PDZ and LIM domain 7 provided by MGI
Primary source	MGI:MGI:1914649
See related	Ensembl:ENSMUSG0000021493
Gene type	protein coding
<b>RefSeq status</b>	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	LMP
Expression	Broad expression in bladder adult (RPKM 74.8), ovary adult (RPKM 59.7) and 26 other tissues See more
Orthologs	human all

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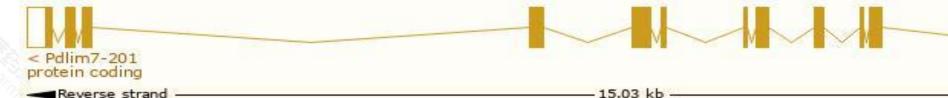
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# **Transcript information (Ensembl)**

#### Transcript ID Flags Name Biotype CCDS UniProt bp Protein ENSMUST00000046246.12 1697 Protein coding CCDS49275 TSL:1 GENCODE basic APPRIS P1 Pdlim7-201 457aa Q3TJD7 222aa Protein coding Pdlim7-203 ENSMUST0000069968.12 1006 CCDS49276 Q8BVJ7 TSL:1 GENCODE basic Q3TJD7 Pdlim7-202 ENSMUST0000069929.12 899 188aa Protein coding CCDS26547 TSL:1 GENCODE basic Pdlim7-206 ENSMUST00000131306.7 1915 219aa Protein coding B8JJB3 TSL:2 GENCODE basic Pdlim7-208 ENSMUST00000144288.7 Protein coding CDS 3' incomplete TSL:5 457 124aa B8JJB2 100 ENSMUST00000153426.1 365 Protein coding CDS 3' incomplete TSL:3 Pdlim7-210 B8JJB1 58aa -Nonsense mediated decay Q3TJD7 Pdlim7-211 ENSMUST00000155098.7 1756 457aa CCDS49275 TSL:1 Pdlim7-209 3573 No protein TSL:2 ENSMUST00000153230.7 Retained intron 1 ENSMUST00000128910.7 1694 No protein Pdlim7-204 Retained intron TSL:2 10 Pdlim7-205 ENSMUST00000128911.1 450 No protein Retained intron TSL:3 -14 396 No protein TSL:2 Pdlim7-207 ENSMUST00000136583.1 Retained intron

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

The strategy is based on the design of *Pdlim7-201* transcript, the transcription is shown below:

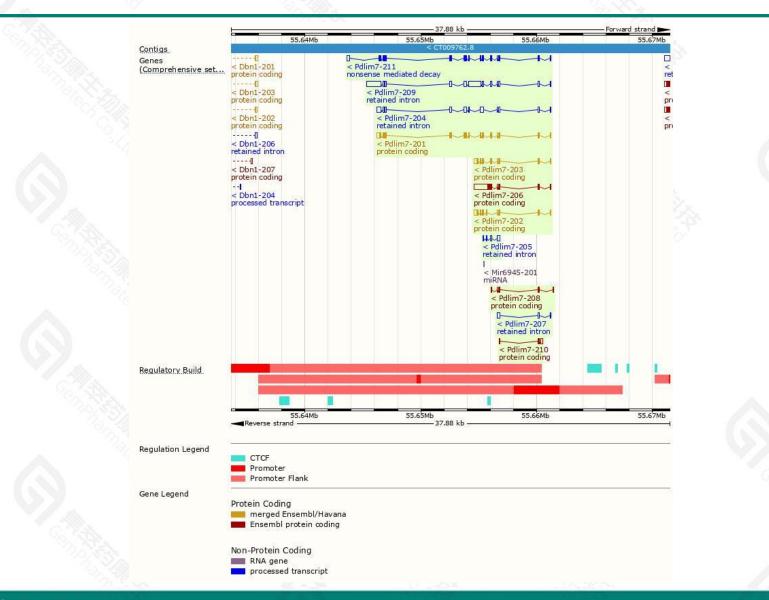


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## **Genomic location distribution**



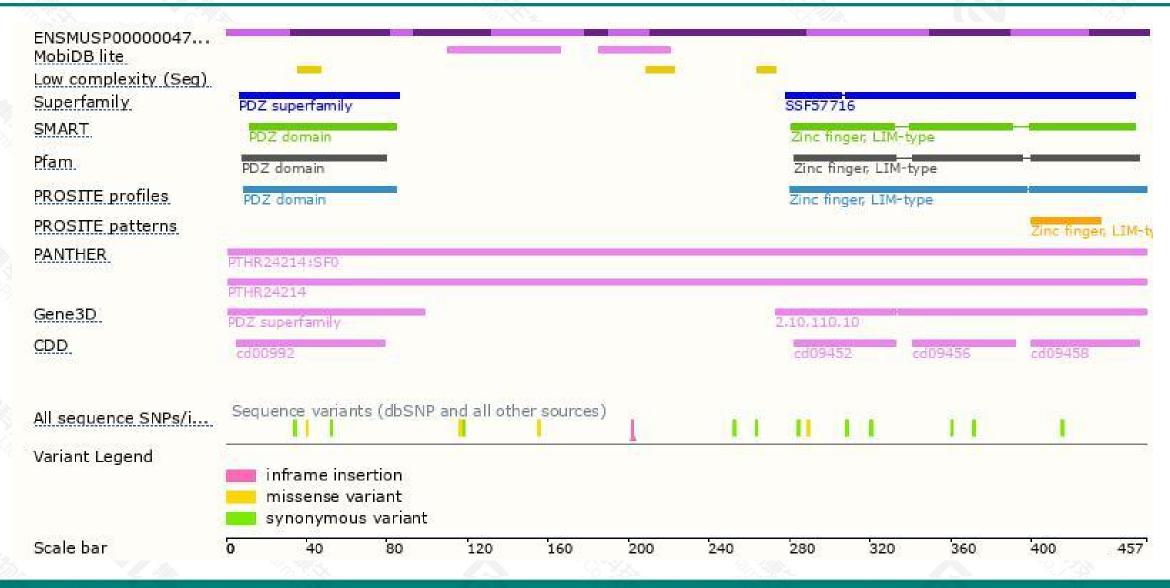


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# **Protein domain**

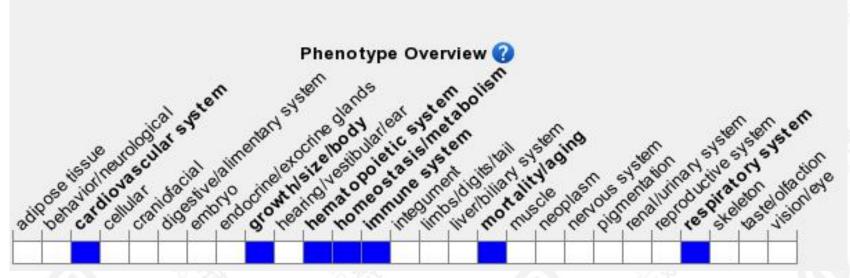




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# Mouse phenotype description(MGI)



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 gene trap allele exhibit heart defects and hemostatic dysfunction.



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



