

# Col18a1 Cas9-KO Strategy

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# **Project Overview**



**Project Name** 

Col18a1

**Project type** 

Cas9-KO

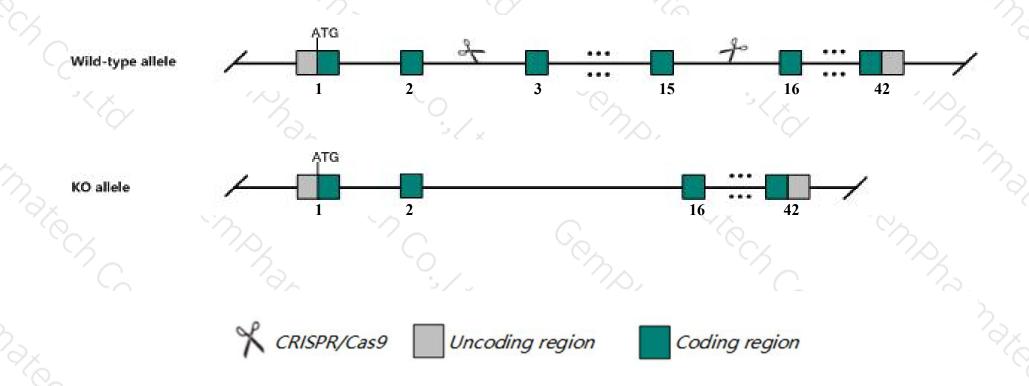
Strain background

C57BL/6JGpt

# **Knockout strategy**



This model will use CRISPR/Cas9 technology to edit the Col18a1 gene. The schematic diagram is as follows:



## **Technical routes**



- ➤ The *Col18a1* gene has 5 transcripts. According to the structure of *Col18a1* gene, exon3-exon15 of *Col18a1-202*(ENSMUST00000081654.12) transcript is recommended as the knockout region. The region contains 1550bp coding sequence Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Col18a1* gene. The brief process is as follows: CRISPR/Cas9 systematically systems.

### **Notice**



- > According to the existing MGI data, Mice homozygous for a knock-out allele exhibit defects in hyaloid vessel regression, attenuated visual function, abnormal electroretinograms, broad proximal tubule basement membrane, podocyte effacement, and softened glomeruli.
- ➤ The effect on transcript *Col18a1*-204&205 is unknown.
- ➤ The strategy may remain the partial function of N-terminal of transcript *Ccna2*-201&203.
- The *Col18a1* gene is located on the Chr10. 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)



#### Col18a1 collagen, type XVIII, alpha 1 [ Mus musculus (house mouse) ]

Gene ID: 12822, updated on 10-Oct-2019

#### Summary

☆ ?

Official Symbol Col18a1 provided by MGI

Official Full Name collagen, type XVIII, alpha 1 provided by MGI

Primary source MGI:MGI:88451

See related Ensembl: ENSMUSG00000001435

Gene type protein coding
RefSeq status VALIDATED
Organism <u>Mus musculus</u>

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

Muridae; Murinae; Mus; Mus

Expression Broad expression in liver adult (RPKM 128.4), liver E18 (RPKM 40.0) and 21 other tissues See more

Orthologs human all

#### Genomic context



**Location:** 10 C1; 10 39.72 cM

See Col18a1 in Genome Data Viewer

Exon count: 44

Annotation release	Status	Assembly	Chr	Location
108	current	GRCm38.p6 (GCF_000001635.26)	10	NC_000076.6 (7705217877166530, complement)
Build 37.2	previous assembly	MGSCv37 (GCF_000001635.18)	10	NC_000076.5 (7651492476629275, complement)

# Transcript information (Ensembl)



The gene has 5 transcripts, all transcripts are shown below:

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Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Col18a1-202	ENSMUST00000081654.12	5063	<u>1315aa</u>	Protein coding	CCDS23953	P39061	TSL:1 GENCODE basic
Col18a1-203	ENSMUST00000105409.7	4791	<u>1527aa</u>	Protein coding	CCDS48604	P39061	TSL:1 GENCODE basic APPRIS P2
Col18a1-201	ENSMUST00000072755.11	5595	<u>1774aa</u>	Protein coding	120	E9QPX1	TSL:5 GENCODE basic APPRIS ALT2
Col18a1-204	ENSMUST00000156009.1	717	<u>161aa</u>	Protein coding	7925	D3Z556	CDS 3' incomplete TSL:3
Col18a1-205	ENSMUST00000218407.1	675	225aa	Protein coding	127	A0A1W2P6N8	5' and 3' truncations in transcript evidence prevent annotation of the start and the end of the CDS. CDS 5' and 3' incomplete TSL:3
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The strategy is based on the design of Col18a1-202 transcript, The transcription is shown below

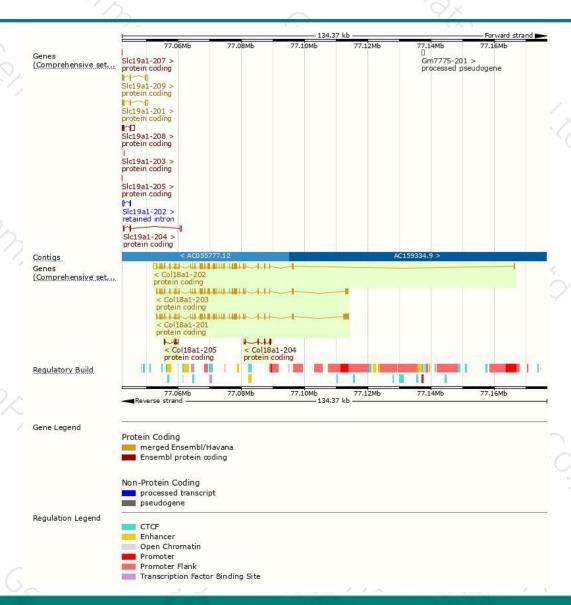


Reverse strand

-114.37 kb

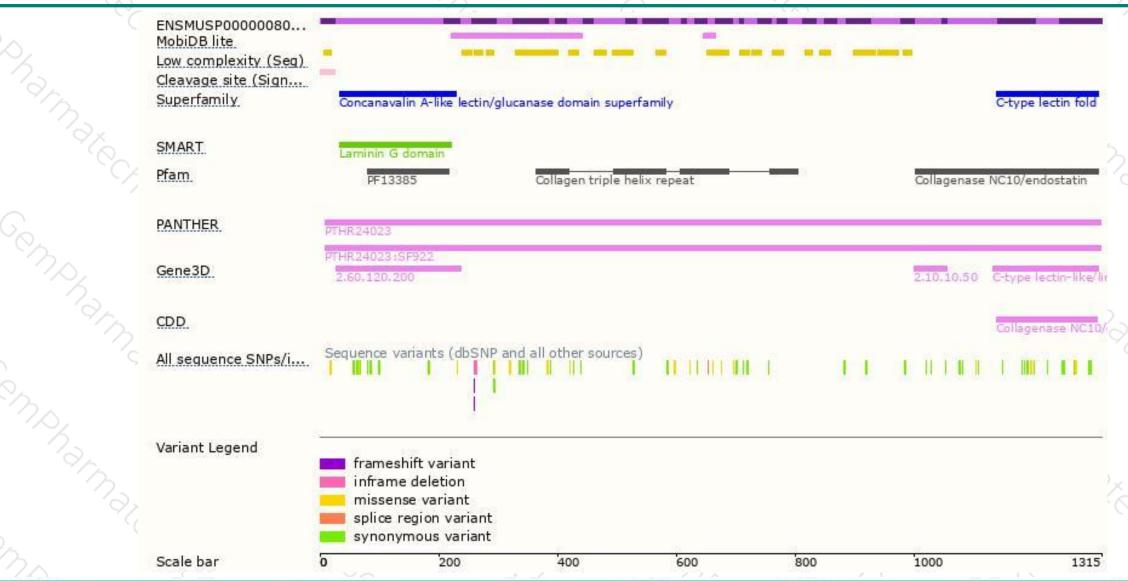
### Genomic location distribution





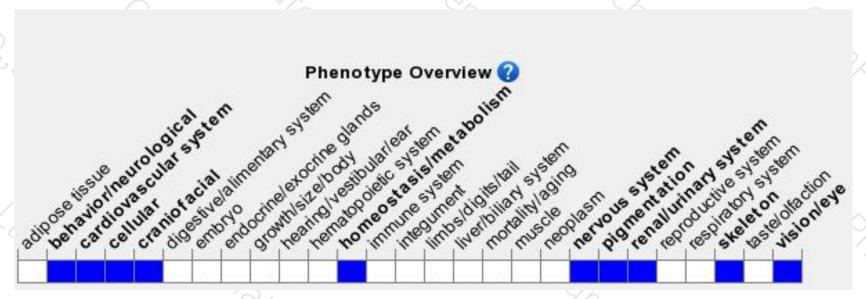
## Protein domain





# 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 defects in hyaloid vessel regression, attenuated visual function, abnormal electroretinograms, broad proximal tubule basement membrane, podocyte effacement, and softened glomeruli.



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





