

# Collal Cas9-KO Strategy

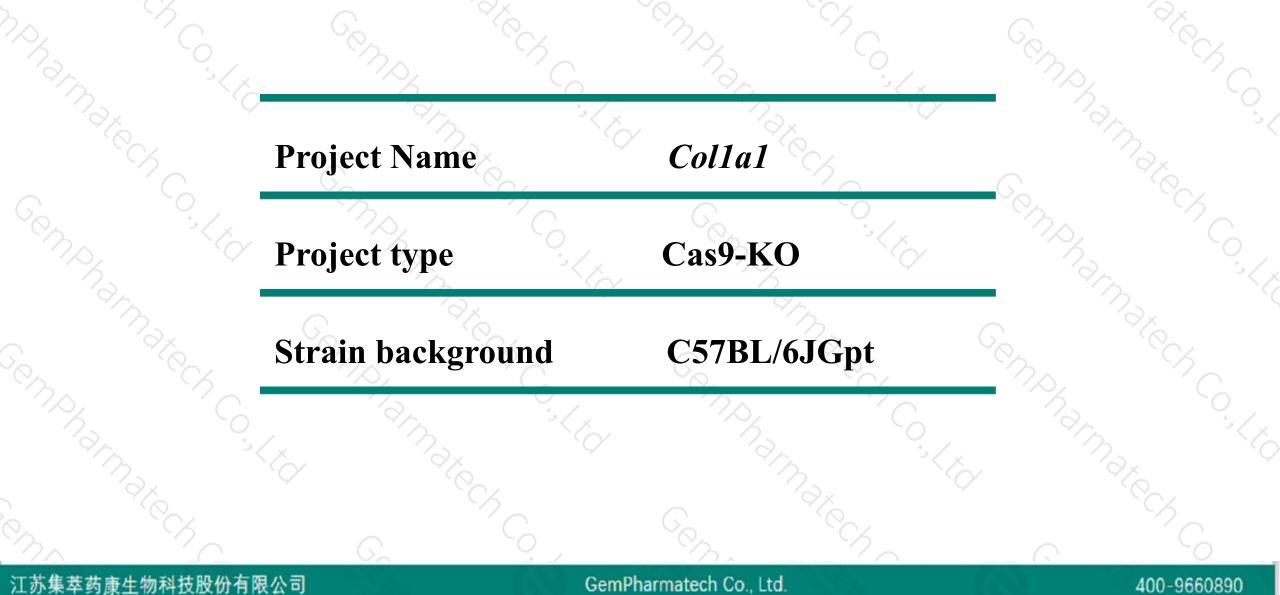
Designer: Reviewer:

**Design Date:** 

Daohua Xu Huimin Su 2019-10-17

## **Project Overview**

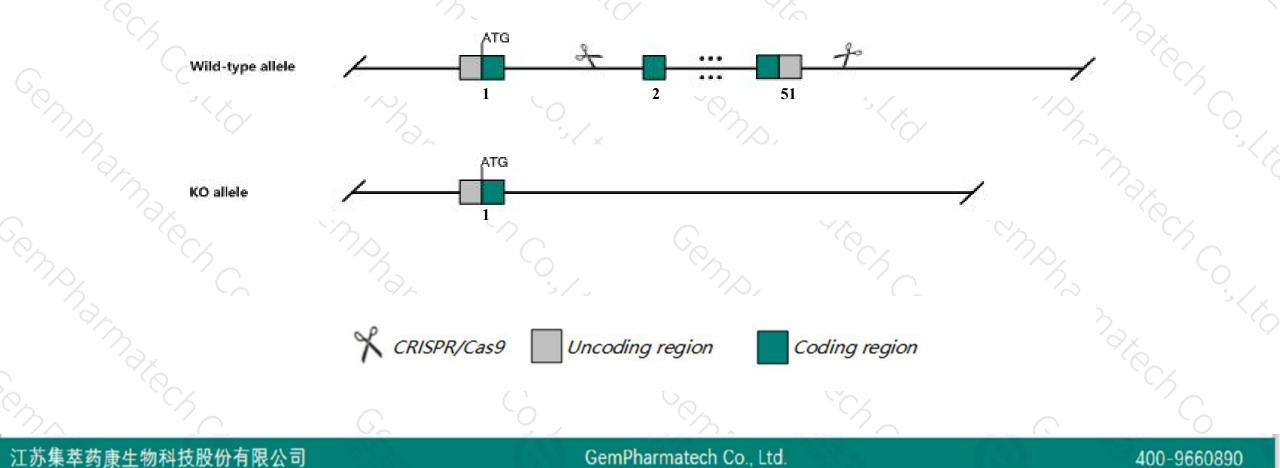




# **Knockout strategy**



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





- The Collal gene has 4 transcripts. According to the structure of Collal gene, exon2-exon51 of Collal-201 (ENSMUST0000001547.7) transcript is recommended as the knockout region. The region contains 4286bp coding sequence. Knock out the region will result in disruption of protein function.
- > In this project we use CRISPR/Cas9 technology to modify Collal gene. The brief process is as follows: CRISPR/Cas9 system



- According to the existing MGI data, Mutations in this locus cause variable phenotype, from embryonic lethal to viable/fertile with altered fibrillogenesis. Homozygotes can show impaired bone formation and fragility, osteoporosis, dermal fibrosis, impaired uterine postpartum involution, and aortic dissection.
- The Collal gene is located on the Chr11. 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)**



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#### Col1a1 collagen, type I, alpha 1 [Mus musculus (house mouse)]

Gene ID: 12842, updated on 9-Apr-2019

#### Summary

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Official Symbol	Col1a1 provided by MGI	
Official Full Name	collagen, type I, alpha 1 provided by <u>MGI</u>	
Primary source	MGI:MGI:88467	
See related	Ensembl:ENSMUSG0000001506	
Gene type	protein coding	
<b>RefSeq status</b>	REVIEWED	
Organism	Mus musculus	
Lineage	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha;	
	Muroidea; Muridae; Murinae; Mus; Mus	
Also known as	Col1a-1, Cola-1, Cola1, Mov-13, Mov13	
Summary	This gene encodes the alpha-1 subunit of the fibril-forming type I collagen, the most abundant protein of bone, skin and tendon extracellular	
	matrices. The encoded protein, in association with alpha-2 subunit, forms heterotrimeric type I procollagen that undergoes proteolytic	
	processing during fibril formation. Mice lacking the encoded protein die in utero caused by the rupture of a major blood vessel. Transgenic	
	mice expressing significantly lower levels of this gene exhibit morphological and functional defects in mineralized and non-mineralized	
	connective tissue and, progressive loss of hearing. [provided by RefSeq, Nov 2015]	
Expression	Biased expression in limb E14.5 (RPKM 527.2), bladder adult (RPKM 493.7) and 12 other tissues See more	
Orthologs	human all	

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## **Transcript information (Ensembl)**



The gene has 4 transcripts, all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Col1a1-201	ENSMUST0000001547.7	5930	<u>1453aa</u>	Protein coding	CCDS25265	P11087	TSL:1 GENCODE basic APPRIS P1
Col1a1-202	ENSMUST00000139974.1	337	No protein	Processed transcript	-	87	TSL:3
Col1a1-204	ENSMUST00000148593.1	493	No protein	Retained intron	-	84	TSL:3
Col1a1-203	ENSMUST00000148046.1	363	No protein	Retained intron	92	1 <u>11</u>	TSL:3
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		- Str			~~~		and the second sec

16.82 kb

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

Collal-201 > protein coding

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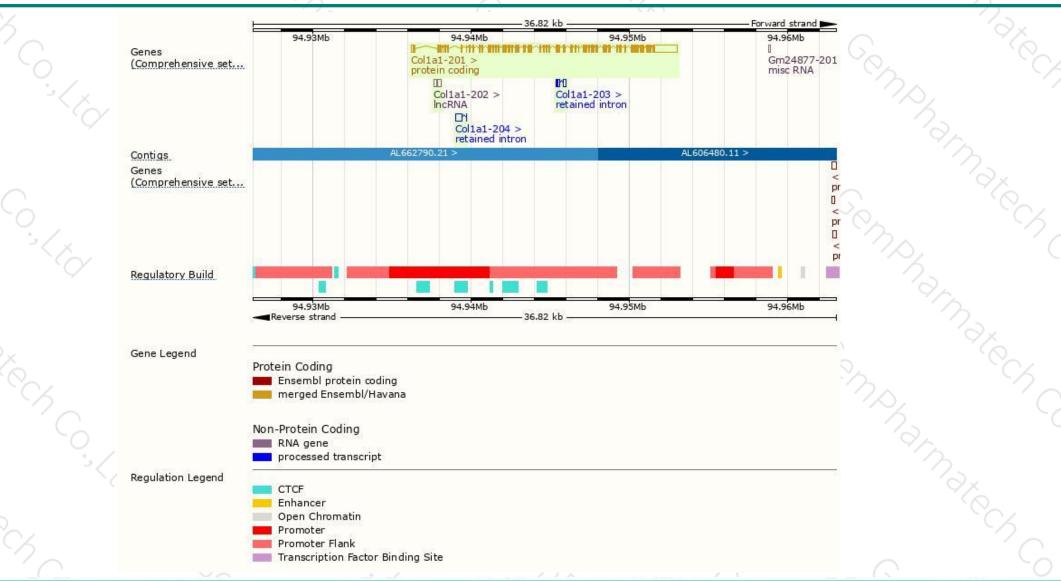
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Forward strand

### **Genomic location distribution**



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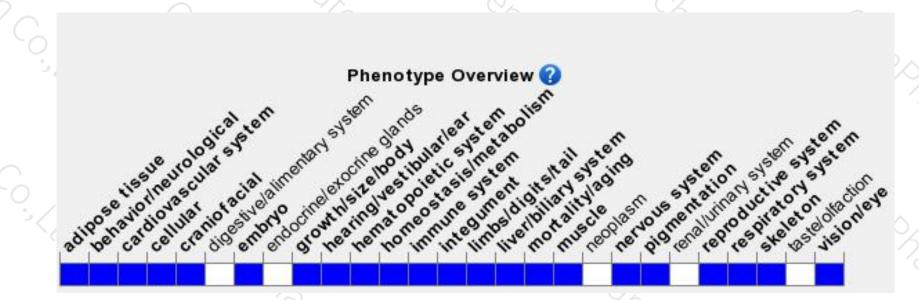
### **Protein domain**





### 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, Mutations in this locus cause variable phenotype, from embryonic lethal to viable/fertile with altered fibrillogenesis. Homozygotes can show impaired bone formation and fragility, osteoporosis, dermal fibrosis, impaired uterine postpartum involution, and a dissection.

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



