

# *Colec11* Cas9-KO Strategy

**Designer:** Daohua Xu

**Reviewer:** Huimin Su

**Design Date:** 2020-3-30

# Project Overview



**Project Name**

***Colec11***

**Project type**

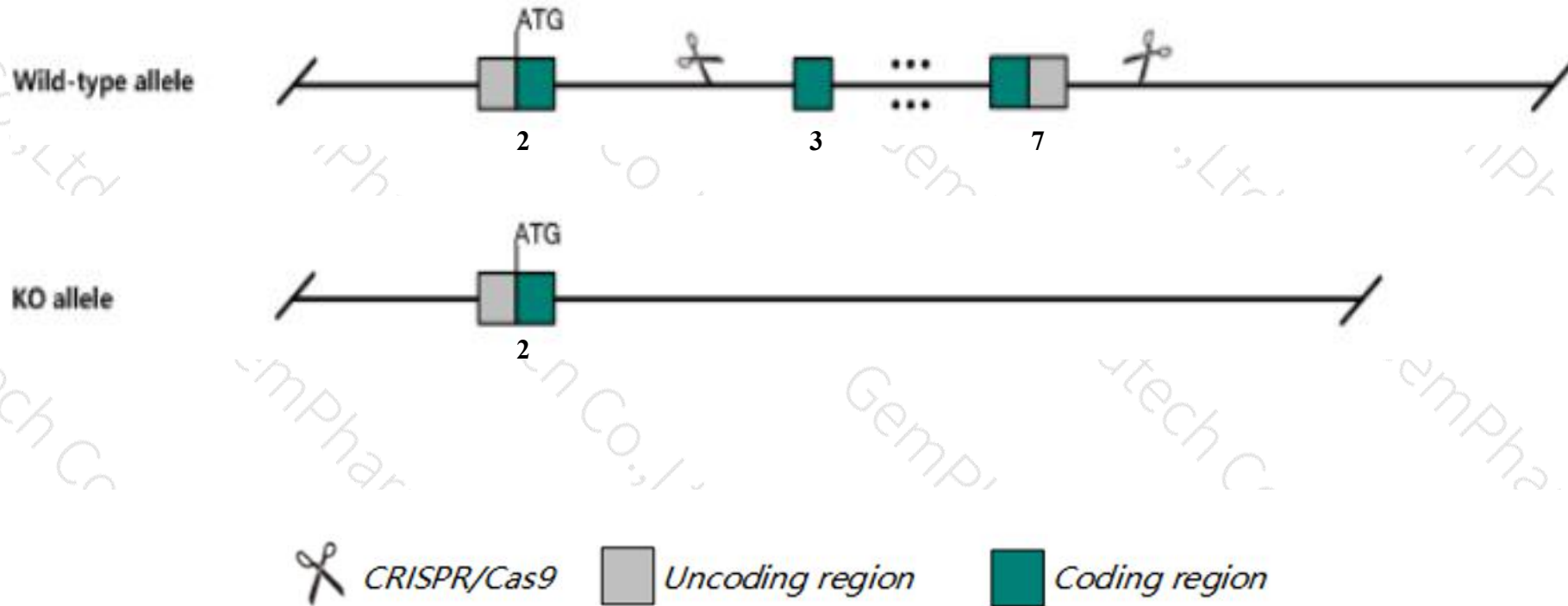
**Cas9-KO**

**Strain background**

**C57BL/6JGpt**

# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Colec11* gene. The schematic diagram is as follows:



- The *Colec11* gene has 3 transcripts. According to the structure of *Colec11* gene, exon3-exon7 of *Colec11-201* (ENSMUST00000036136.8) transcript is recommended as the knockout region. The region contains 686bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Colec11* gene. The brief process is as follows: CRISPR/Cas9 system

- According to the existing MGI data, Mice homozygous for a knockout allele exhibit decreased susceptibility to kidney reperfusion injury.
- The *Colec11* gene is located on the Chr12. 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)

## Colec11 collectin sub-family member 11 [ *Mus musculus* (house mouse) ]

Gene ID: 71693, updated on 13-Mar-2020

### Summary

<b>Official Symbol</b>	Colec11 provided by <a href="#">MGI</a>
<b>Official Full Name</b>	collectin sub-family member 11 provided by <a href="#">MGI</a>
<b>Primary source</b>	<a href="#">MGI:MGI:1918943</a>
<b>See related</b>	<a href="#">Ensembl:ENSMUSG00000036655</a>
<b>Gene type</b>	protein coding
<b>RefSeq status</b>	REVIEWED
<b>Organism</b>	<a href="#">Mus musculus</a>
<b>Lineage</b>	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
<b>Also known as</b>	CL-K1
<b>Summary</b>	This gene encodes a member of the collectin family of C-type lectins that possess collagen-like sequences and carbohydrate recognition domains. Collectins are secreted proteins that play important roles in the innate immune system by binding to carbohydrate antigens on microorganisms, facilitating their recognition and removal. The encoded protein binds to multiple sugars with a preference for fucose and mannose. Mutations in the human gene are a cause of 3MC syndrome-2. Alternative splicing results in multiple transcript variants encoding different isoforms. [provided by RefSeq, Sep 2015]
<b>Expression</b>	Biased expression in liver adult (RPKM 12.7), liver E18 (RPKM 9.2) and 11 other tissues <a href="#">See more</a>
<b>Orthologs</b>	<a href="#">human</a> <a href="#">all</a>

# Transcript information (Ensembl)

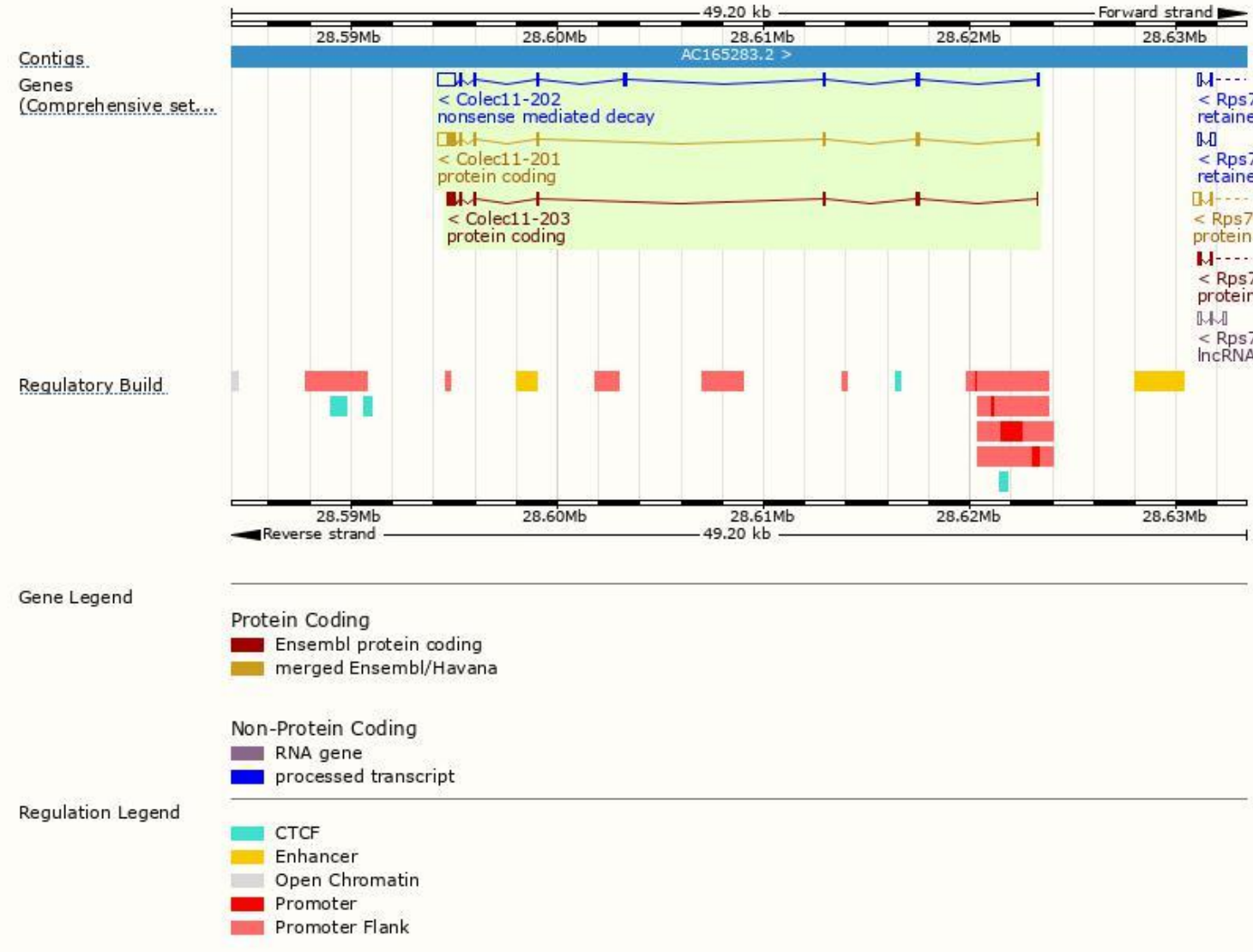
The gene has 3 transcripts, all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Colec11-201	<a href="#">ENSMUST00000036136.8</a>	1423	<a href="#">272aa</a>	Protein coding	<a href="#">CCDS36426</a>	<a href="#">A0A0R4J0M6</a>	TSL:1 GENCODE basic APPRIS P1
Colec11-203	<a href="#">ENSMUST00000220836.1</a>	887	<a href="#">278aa</a>	Protein coding	-	<a href="#">A0A1Y7VKG6</a>	TSL:1 GENCODE basic
Colec11-202	<a href="#">ENSMUST00000220655.1</a>	1603	<a href="#">91aa</a>	Nonsense mediated decay	-	<a href="#">A0A1Y7VLI6</a>	TSL:5

The strategy is based on the design of *Colec11-201* transcript, The transcription is shown below

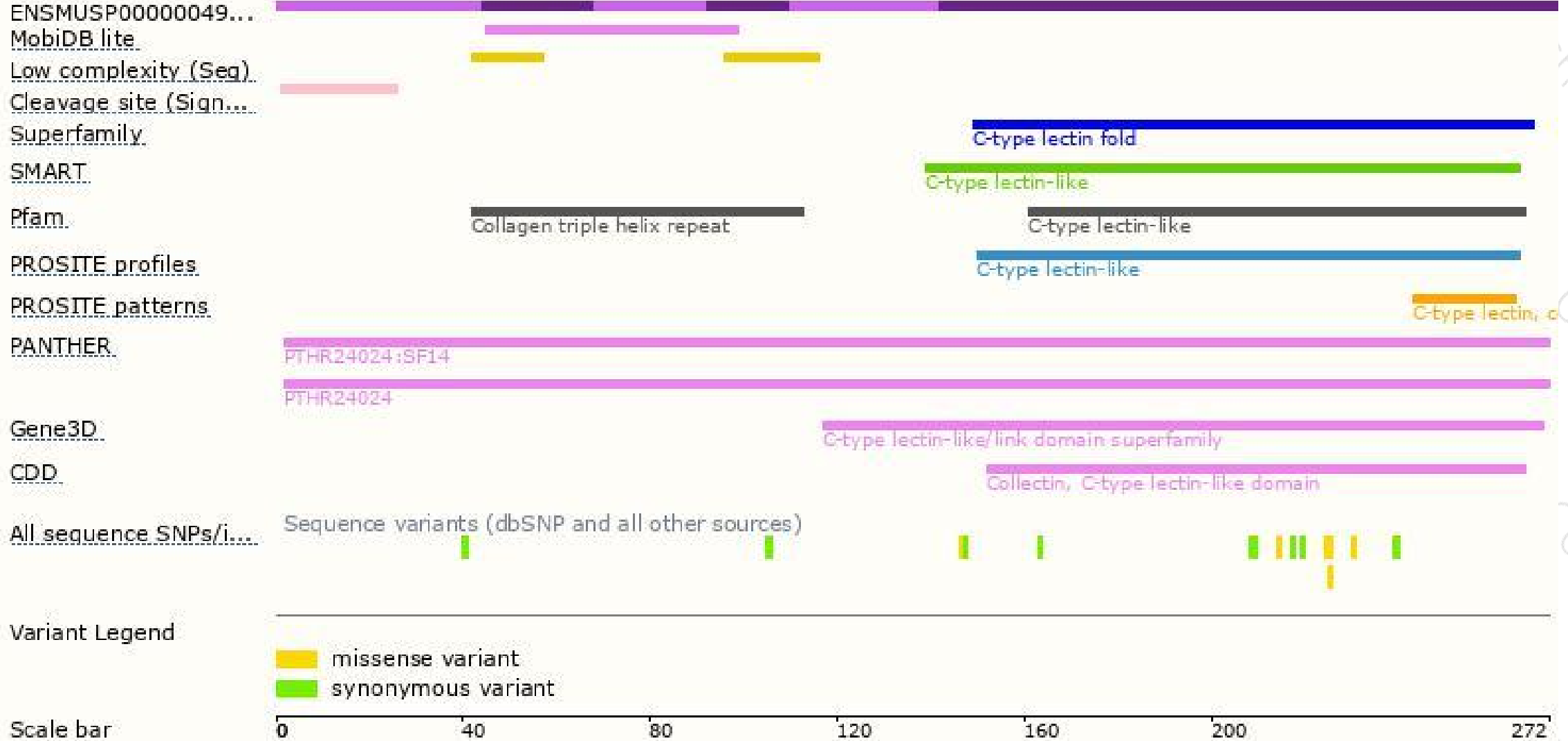


# Genomic location distribution

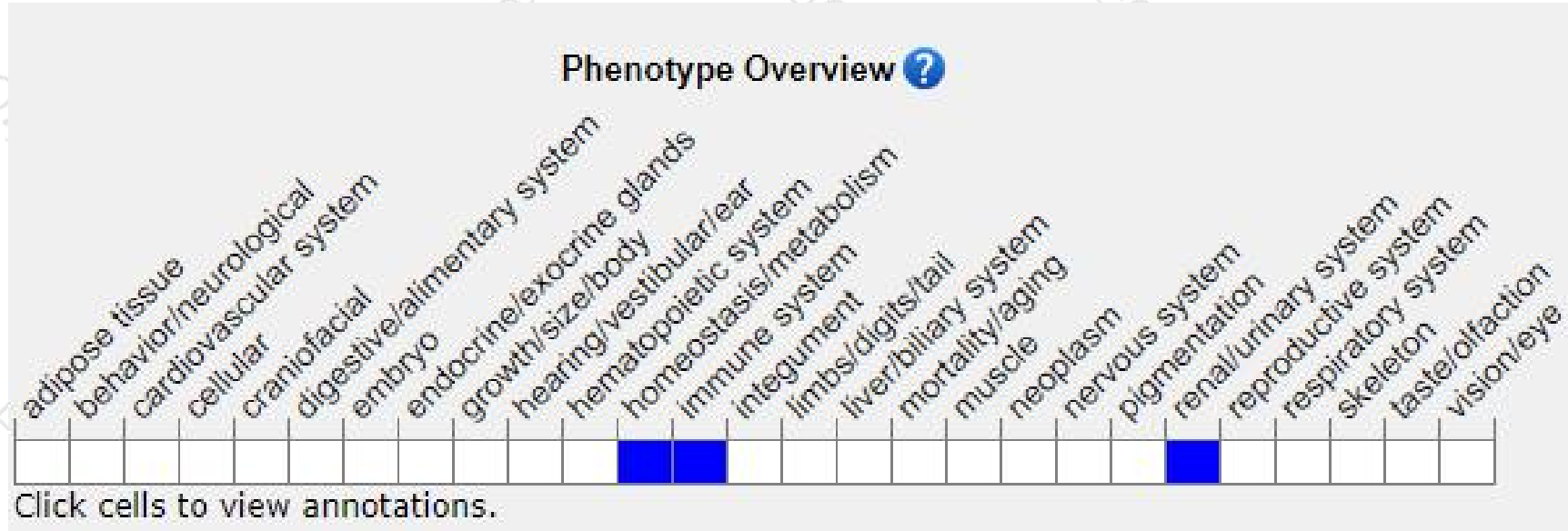




# Protein domain



# 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 knockout allele exhibit decreased susceptibility to kidney reperfusion injury.

If you have any questions, you are welcome to inquire.

Tel: 400-9660890

