

# *Mgat5b* Cas9-CKO Strategy

**Designer:**

**JiaYu**

**Reviewer:**

**Xiaojing Li**

**Design Date:**

**2020-2-18**

# Project Overview

**Project Name**

***Mgat5b***

**Project type**

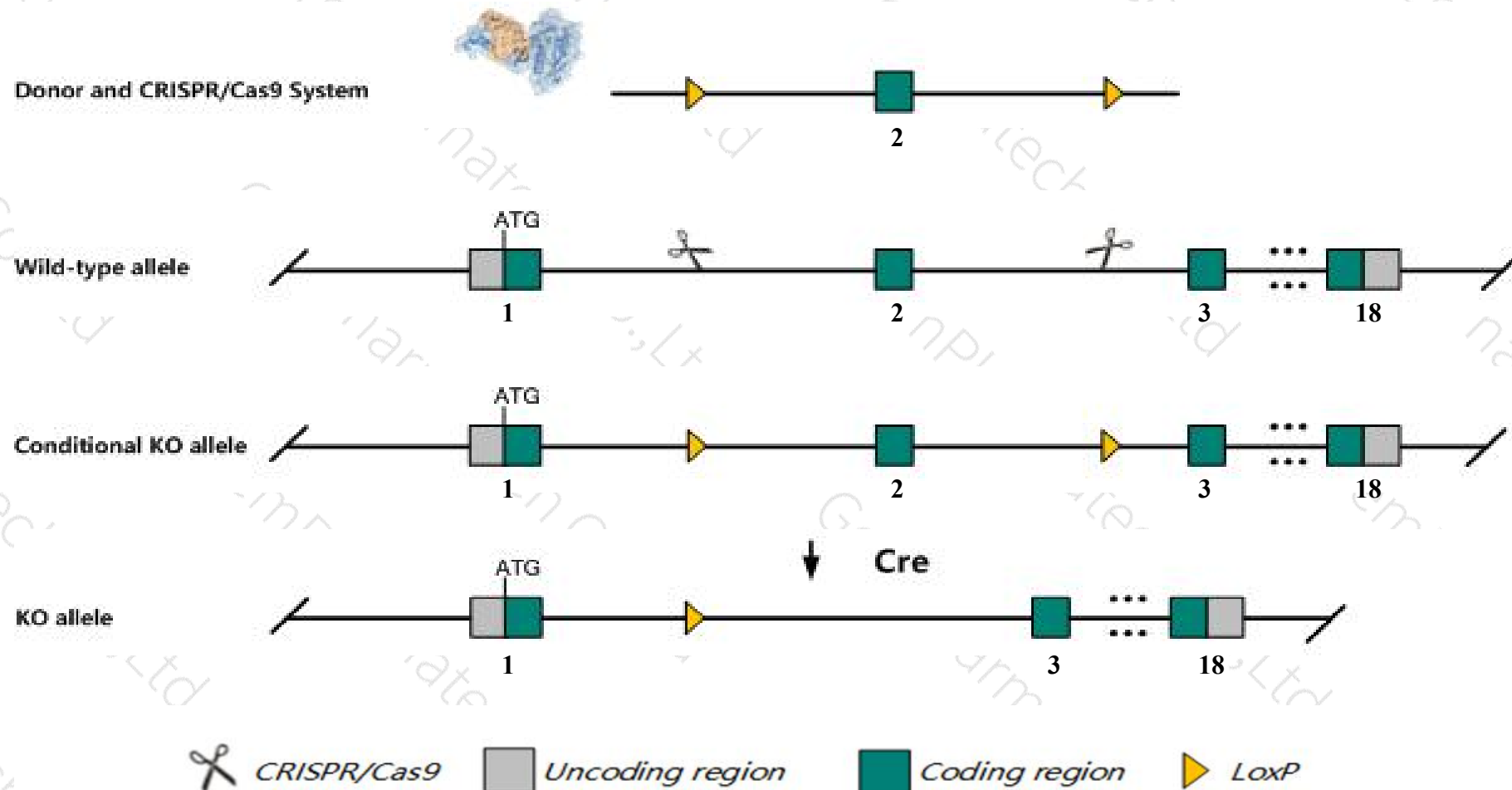
**Cas9-CKO**

**Strain background**

**C57BL/6JGpt**

# Conditional Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Mgat5b* gene. The schematic diagram is as follows:



# Technical routes

- The *Mgat5b* gene has 4 transcripts. According to the structure of *Mgat5b* gene, exon2 of *Mgat5b-201* (ENSMUST00000103027.9) transcript is recommended as the knockout region. The region contains 113bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Mgat5b* 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.
- 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 knock-out allele exhibit low levels of O-man-linked beta1,6-branched glycans. Mice homozygous for a different knock-out allele exhibit decreased susceptibility to cuprizone induced injury.
- The *Mgat5b* 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 loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at existing technological level.



# Gene information (NCBI)

## Mgat5b mannoside acetylglucosaminyltransferase 5, isoenzyme B [Mus musculus (house mouse)]

Gene ID: 268510, updated on 31-Jan-2019

### Summary



<b>Official Symbol</b>	Mgat5b provided by <a href="#">MGI</a>
<b>Official Full Name</b>	mannoside acetylglucosaminyltransferase 5, isoenzyme B provided by <a href="#">MGI</a>
<b>Primary source</b>	<a href="#">MGI:MGI:3606200</a>
<b>See related</b>	<a href="#">Ensembl:ENSMUSG00000043857</a>
<b>Gene type</b>	protein coding
<b>RefSeq status</b>	PROVISIONAL
<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>	C330018B01, GnT-IX, mGnTVB
<b>Expression</b>	Biased expression in whole brain E14.5 (RPKM 22.3), CNS E18 (RPKM 20.1) and 8 other tissues <a href="#">See more</a>
<b>Orthologs</b>	<a href="#">human</a> <a href="#">all</a>

# Transcript information (Ensembl)

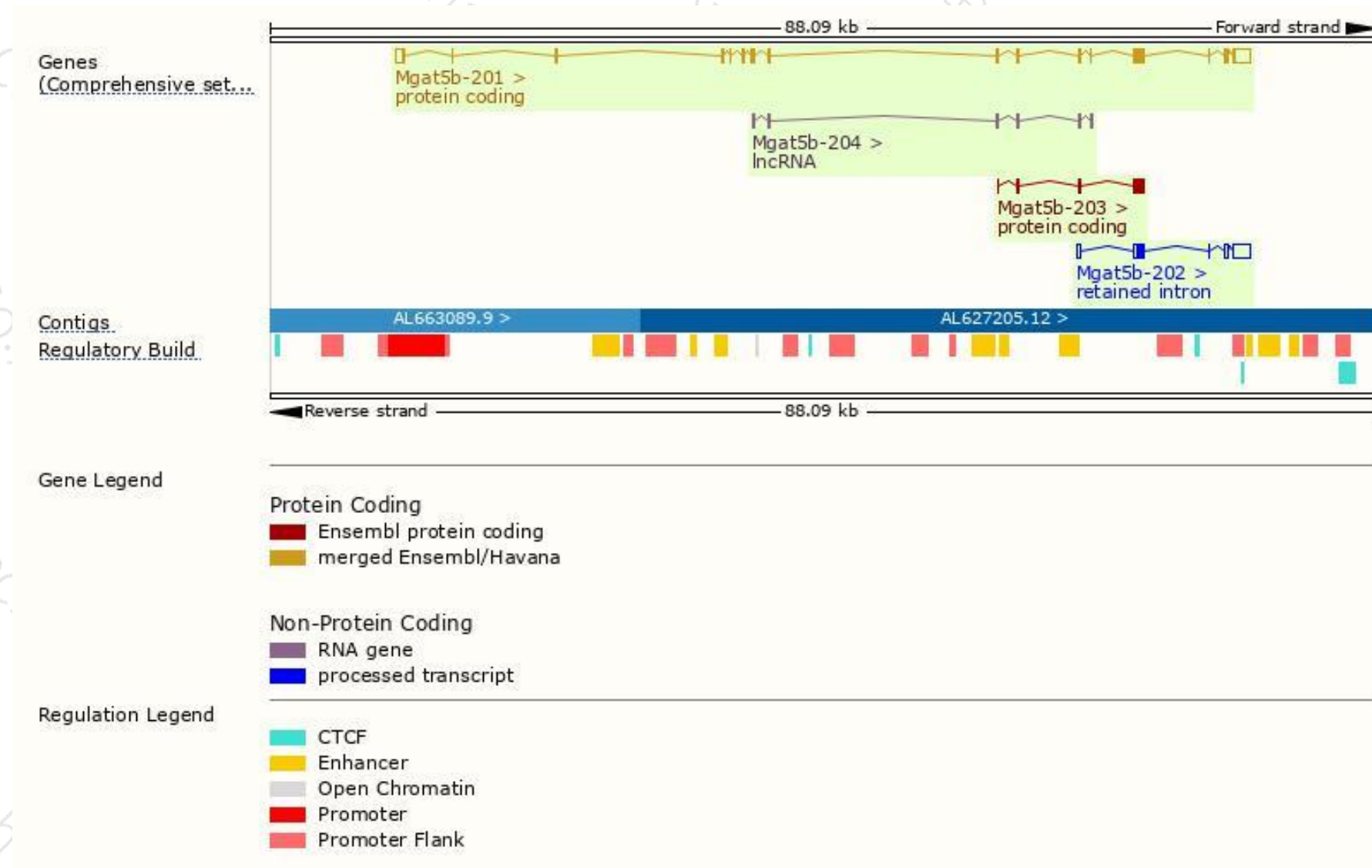
The gene has 4 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Mgat5b-201	<a href="#">ENSMUST00000103027.9</a>	4258	<a href="#">792aa</a>	Protein coding	<a href="#">CCDS25682</a>	<a href="#">Q765H6</a>	TSL:5 GENCODE basic APPRIS P1
Mgat5b-203	<a href="#">ENSMUST00000136584.1</a>	641	<a href="#">214aa</a>	Protein coding	-	<a href="#">F7BPB5</a>	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
Mgat5b-202	<a href="#">ENSMUST00000126757.1</a>	2568	No protein	Retained intron	-	-	TSL:1
Mgat5b-204	<a href="#">ENSMUST00000139905.1</a>	725	No protein	lncRNA	-	-	TSL:3

The strategy is based on the design of *Mgat5b-201* transcript,The transcription is shown below

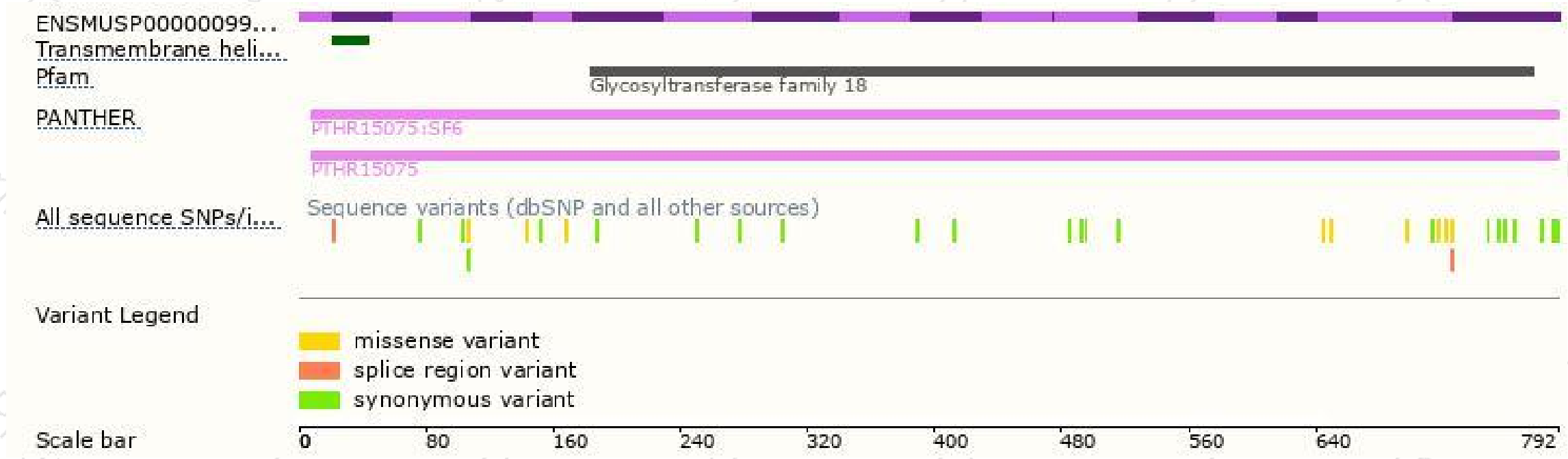


# Genomic location distribution

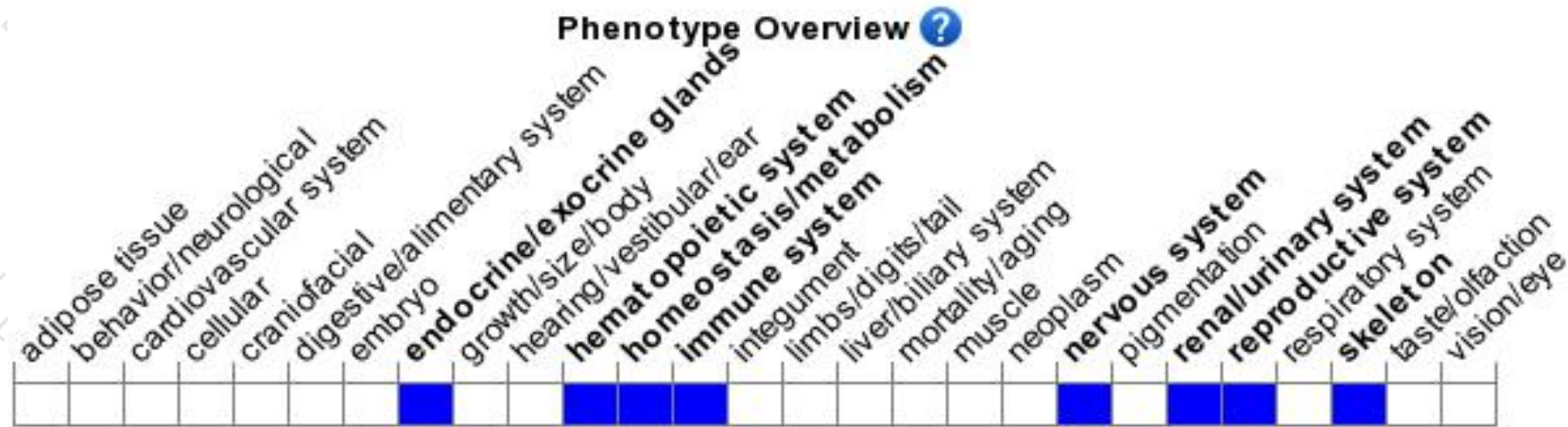




# Protein domain



# 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 knock-out allele exhibit low levels of O-man-linked beta1,6-branched glycans. Mice homozygous for a different knock-out allele exhibit decreased susceptibility to cuprizone induced injury.

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

Tel: 400-9660890

