

# B3gat3 Cas9-CKO Strategy

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## **Project Overview**



**Project Name** 

B3gat3

**Project type** 

Cas9-CKO

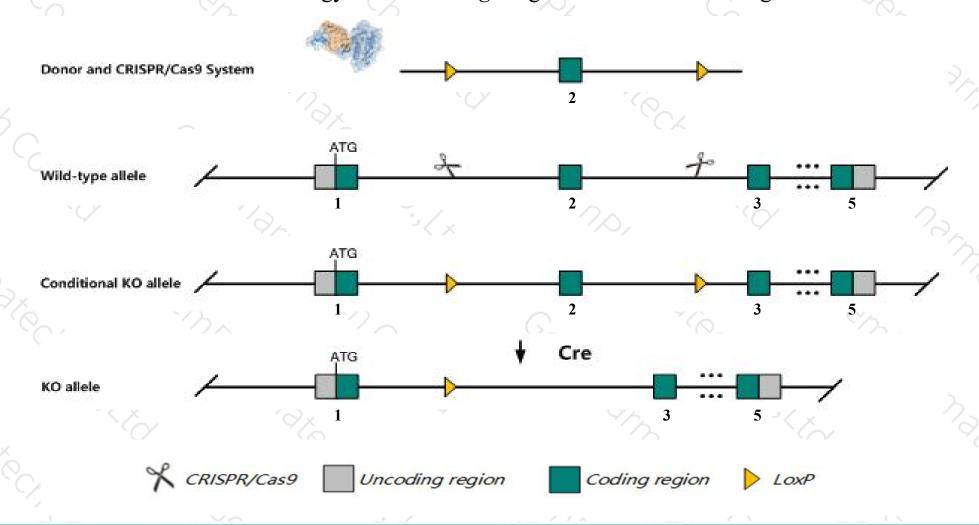
Strain background

C57BL/6JGpt

## Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *B3gat3* gene. The schematic diagram is as follows:



## Technical routes



- ➤ The *B3gat3* gene has 3 transcripts. According to the structure of *B3gat3* gene, exon2 of *B3gat3-201*(ENSMUST00000096243.6) transcript is recommended as the knockout region. The region contains 175bp coding sequence.

  Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *B3gat3* 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.

## **Notice**



- ➤ According to the existing MGI data, Homozygous mutants die prenatally before the 8-cell stage due to failed cytokinesis, and show reduction of the synthesis of chondroitin sulfate and heparan sulfate glycosaminoglycans.
- > The *B3gat3* gene is located on the Chr19. 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)



#### B3gat3 beta-1,3-glucuronyltransferase 3 (glucuronosyltransferase I) [ Mus musculus (house mouse) ]

Gene ID: 72727, updated on 12-Aug-2019

#### Summary

↑ ?

Official Symbol B3gat3 provided by MGI

Official Full Name beta-1,3-glucuronyltransferase 3 (glucuronosyltransferase I) provided by MGI

Primary source MGI:MGI:1919977

See related Ensembl: ENSMUSG00000071649

Gene type protein coding
RefSeq status 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 2810405M13Rik

Expression Ubiquitous expression in adrenal adult (RPKM 58.2), cerebellum adult (RPKM 55.6) and 28 other tissues See more

Orthologs human all

#### **Genomic context**

Location: 19; 19 A

Exon count: 5

☆ ?

See B3gat3 in Genome Data Viewer

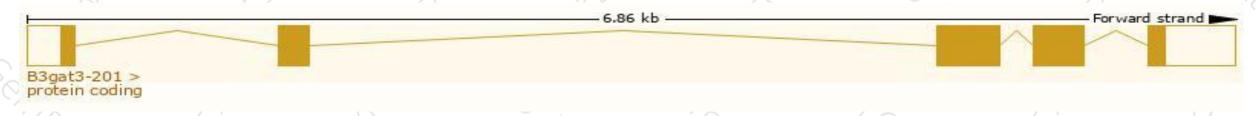
## Transcript information (Ensembl)



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

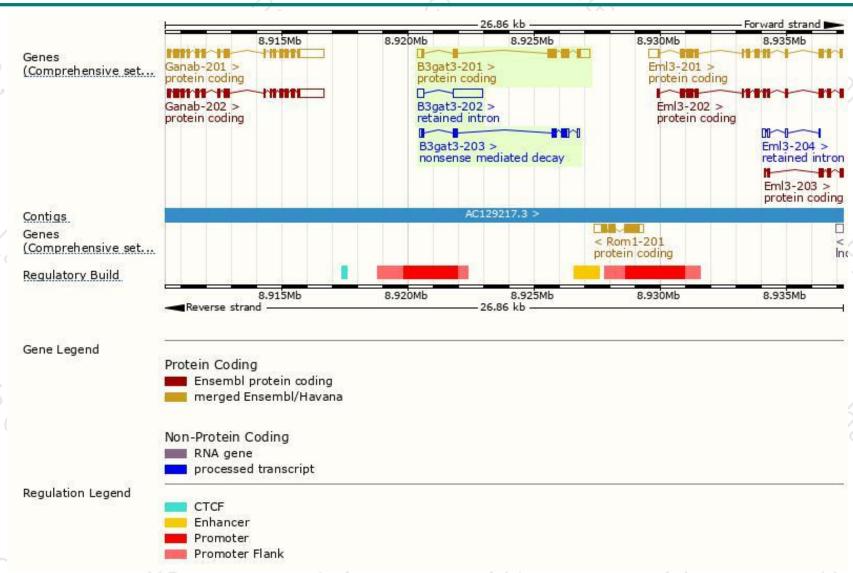
| Name       | Transcript ID        | bp   | Protein      | Biotype                 | CCDS      | UniProt | Flags                         |
|------------|----------------------|------|--------------|-------------------------|-----------|---------|-------------------------------|
| B3gat3-201 | ENSMUST00000096243.6 | 1603 | <u>335aa</u> | Protein coding          | CCDS29558 | P58158  | TSL:1 GENCODE basic APPRIS P1 |
| B3gat3-203 | ENSMUST00000237071.1 | 909  | <u>229aa</u> | Nonsense mediated decay |           |         |                               |
| B3gat3-202 | ENSMUST00000235193.1 | 1414 | No protein   | Retained intron         | -         | -       |                               |

The strategy is based on the design of B3gat3-201 transcript, The transcription is shown below



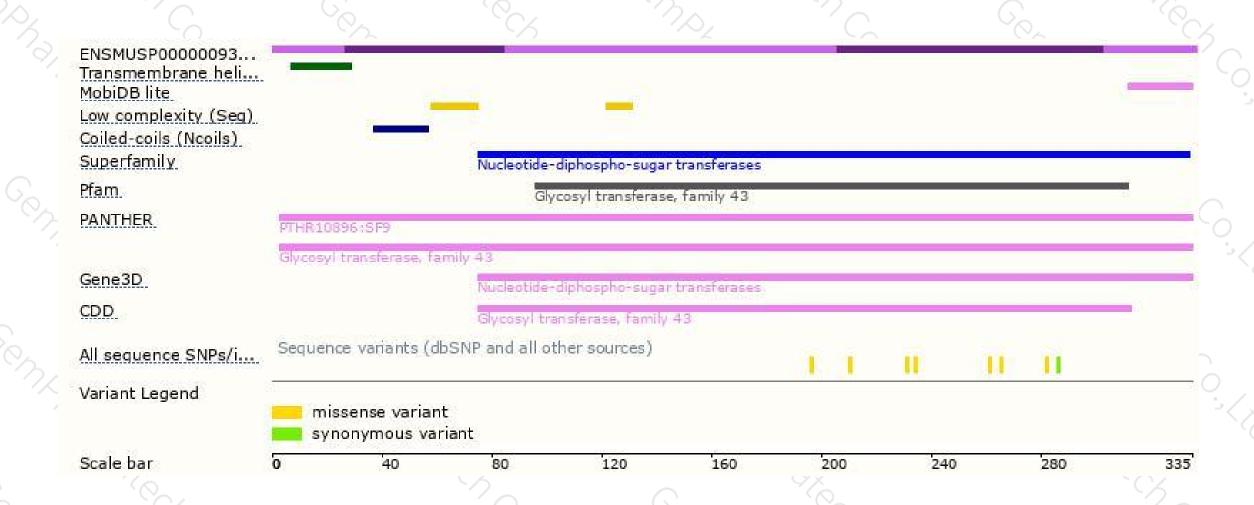
## Genomic location distribution





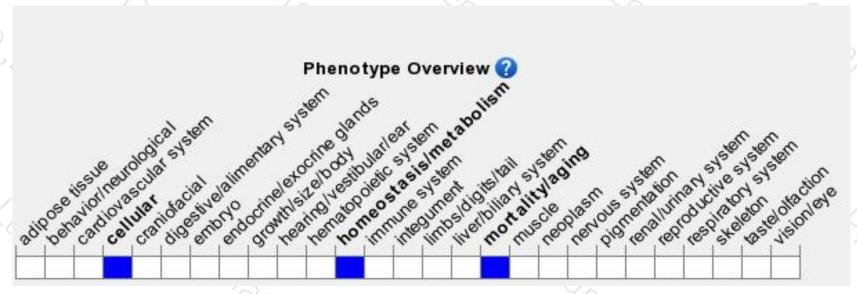
### Protein domain





## 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, Homozygous mutants die prenatally before the 8-cell stage due to failed cytokinesis, and show reduction of the synthesis of chondroitin sulfate and heparan sulfate glycosaminoglycans.



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





