B3gnt6 Cas9-CKO Strategy

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Design Date: 2020-3-10

Project Overview



Project Name

B3gnt6

Project type

Cas9-CKO

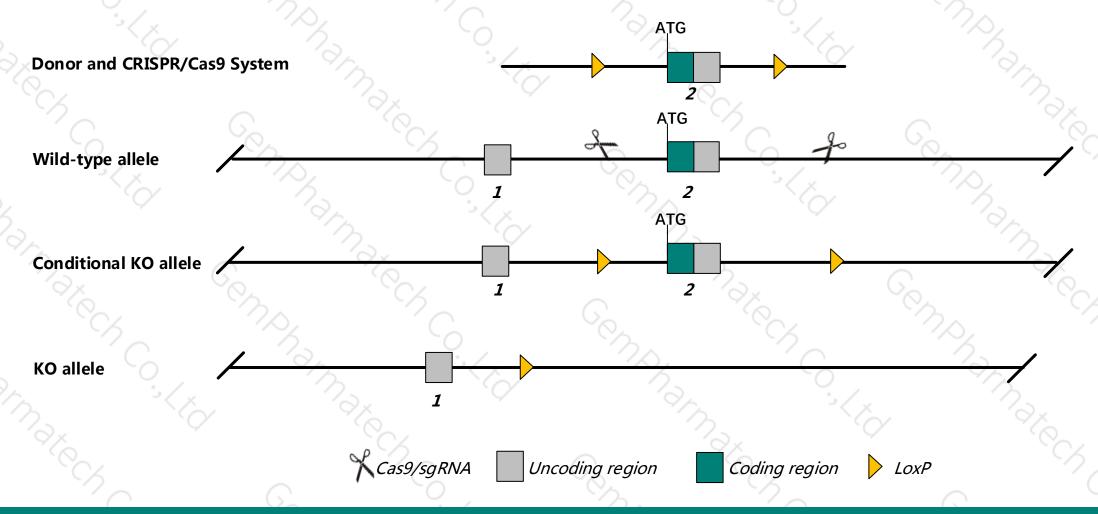
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *B3gnt6* gene. The schematic diagram is as follows:



Technical routes



- ➤ The *B3gnt6* gene has 1 transcript. According to the structure of *B3gnt6* gene, exon 2 of *B3gnt6*-201 (
- ➤ ENSMUST00000098278.3) transcript is recommended as the knockout region. The region contains all of the coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *B3gnt6* gene. The brief process is as follows: sgRNA was transcribed in vitro, donor vector was constructed.Cas9, sgRNA 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 was knocked out after mating with mice expressing Cre recombinase, resulting in the loss of function of the target gene in specific tissues or cell types.

Notice



- According to the existing MGI data, Mice homozgous for a knock-out allele exhibit increased permeability of the intestinal barrier, increased susceptibility to DSS-induced colitis and accelerated colorectal tumorigenesis in mice treated with AOM and DSS.
- ➤ The *B3gnt6* gene is located on the Chr7. 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 loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.

Gene information (NCBI)



B3gnt6 UDP-GlcNAc:betaGal beta-1,3-N-acetylglucosaminyltransferase 6 (core 3 synthase) [Mus musculus (house mouse)]

Gene ID: 272411, updated on 13-Mar-2020

Summary

☆ ?

Official Symbol B3gnt6 provided by MGI

Official Full Name UDP-GlcNAc:betaGal beta-1,3-N-acetylglucosaminyltransferase 6 (core 3 synthase) provided by MGI

Primary source MGI:MGI:3039603

See related Ensembl: ENSMUSG00000074004

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

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

Muroidea; Muridae; Murinae; Mus; Mus

Also known as Core3; Core-3; BC039789

Expression Biased expression in colon adult (RPKM 47.3) and large intestine adult (RPKM 2.1) See more

Orthologs human all

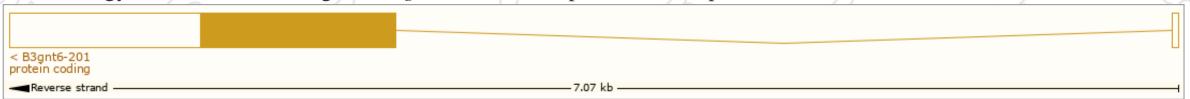
Transcript information (Ensembl)



The gene has 1 transcript, and the transcript is shown below:

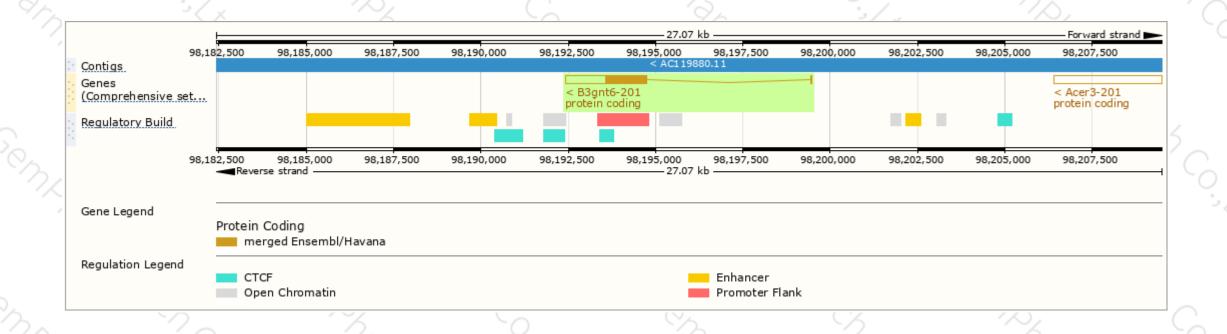
Name 🍦	Transcript ID 🖕	bp 👙	Protein 🍦	Biotype 🍦	CCDS	UniProt ♦		Flags	*
B3gnt6-201	ENSMUST00000098278.3	2371	<u>391aa</u>	Protein coding	CCDS40027 ₺	Q3USF0₽	TSL:1	GENCODE basic	APPRIS P1

The strategy is based on the design of *B3gnt6*-201 transcript, The transcription is shown below



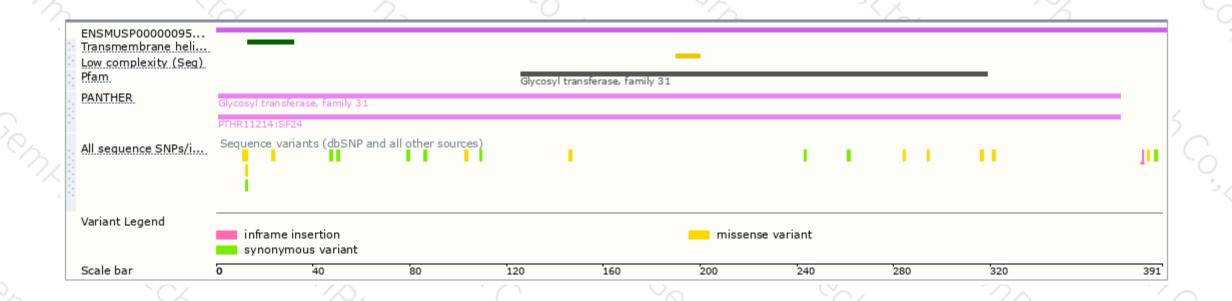
Genomic location distribution





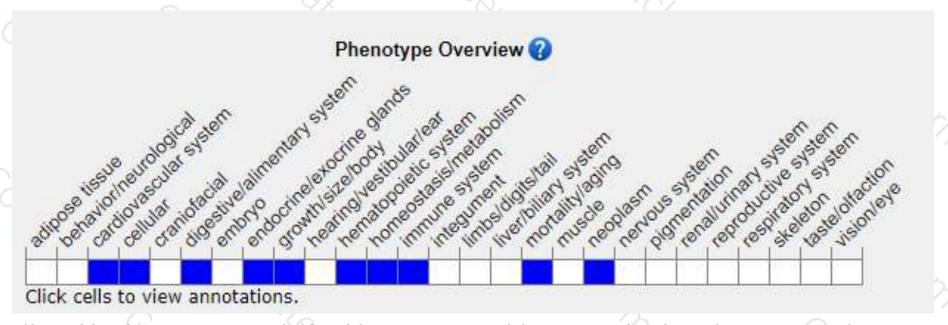
Protein domain





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 homozgous for a knock-out allele exhibit increased permeability of the intestinal barrier, increased susceptibility to DSS-induced colitis and accelerated colorectal tumorigenesis in mice treated with AOM and DSS.

If you have any questions, you are welcome to inquire. Tel: 025-5864 1534





