

B4galt6 Cas9-KO Strategy

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Project Overview

Project Name

B4galt6

Project type

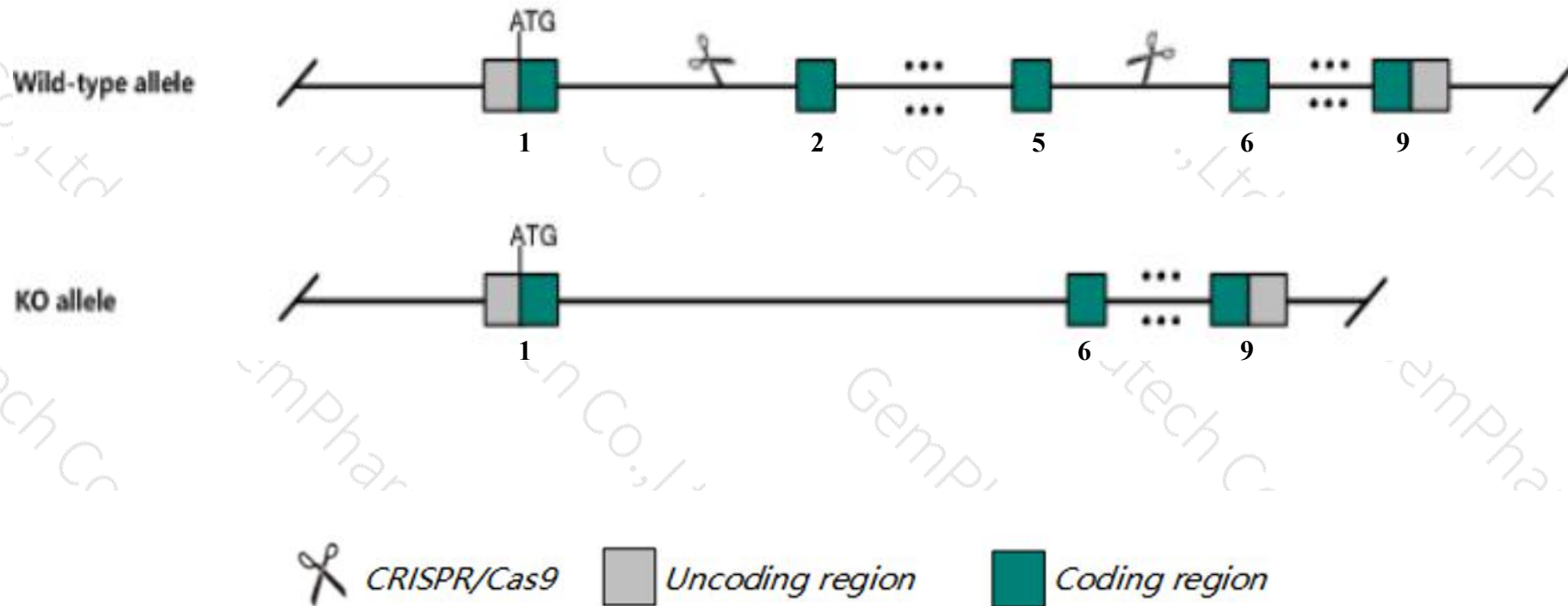
Cas9-KO

Strain background

C57BL/6JGpt

Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *B4galt6* gene. The schematic diagram is as follows:



- The *B4galt6* gene has 1 transcript. According to the structure of *B4galt6* gene, exon2-exon5 of *B4galt6-201* (ENSMUST00000070080.5) transcript is recommended as the knockout region. The region contains 473bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *B4galt6* gene. The brief process is as follows: CRISPR/Cas9 system

- According to the existing MGI data, Mice homozygous for disruptions in this gene display a normal phenotype with reduced lactosylceramide synthase in MEFs.
- The *B4galt6* gene is located on the Chr18. 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)

B4galt6 UDP-Gal:betaGlcNAc beta 1,4-galactosyltransferase, polypeptide 6 [*Mus musculus* (house mouse)]

Gene ID: 56386, updated on 12-Aug-2019

Summary

- Official Symbol** B4galt6 provided by [MGI](#)
- Official Full Name** UDP-Gal:betaGlcNAc beta 1,4-galactosyltransferase, polypeptide 6 provided by [MGI](#)
- Primary source** [MGI:MGI:1928380](#)
- See related** [Ensembl:ENSMUSG00000056124](#)
- 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** AA536803; AU022389
- Expression** Broad expression in small intestine adult (RPKM 16.5), duodenum adult (RPKM 16.3) and 23 other tissues [See more](#)
- Orthologs** [human](#) [all](#)

Genomic context

Location: 18; 18 A2 See B4galt6 in [Genome Data Viewer](#)

Exon count: 10

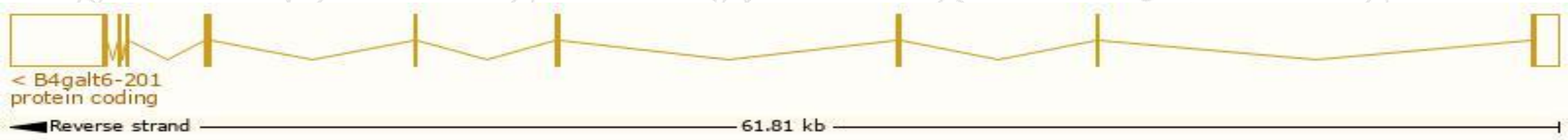
Annotation release	Status	Assembly	Chr	Location
108	current	GRCm38.p6 (GCF_000001635.26)	18	NC_000084.6 (20684599..20746404, complement)
Build 37.2	previous assembly	MGSCv37 (GCF_000001635.18)	18	NC_000084.5 (20843100..20904905, complement)

Transcript information (Ensembl)

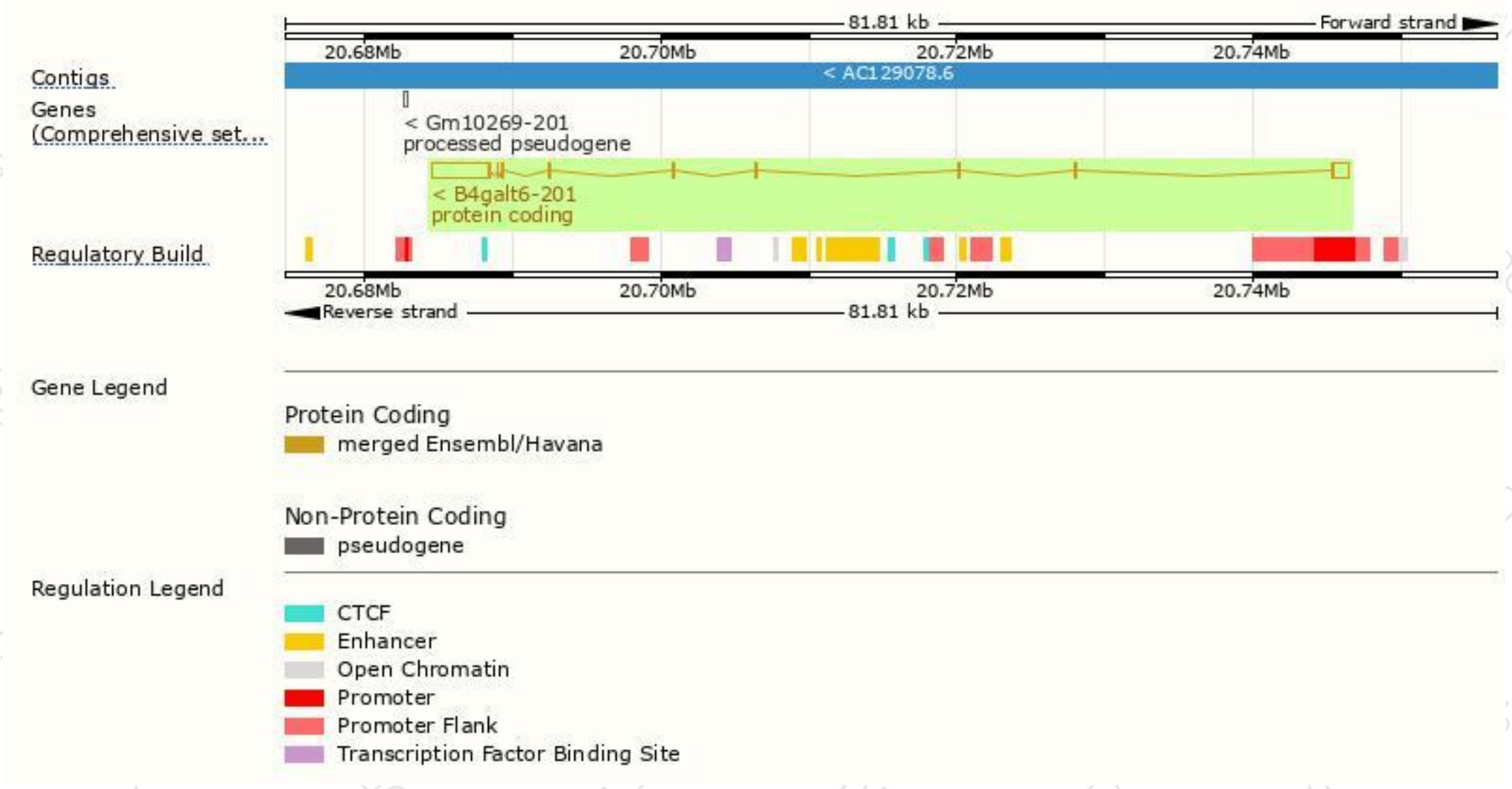
The gene has 1 transcript, and the transcript is shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
B4galt6-201	ENSMUST00000070080.5	5808	382aa	Protein coding	CCDS29086	Q3UUA9 Q9WVK5	TSL:1 GENCODE basic APPRIS P1

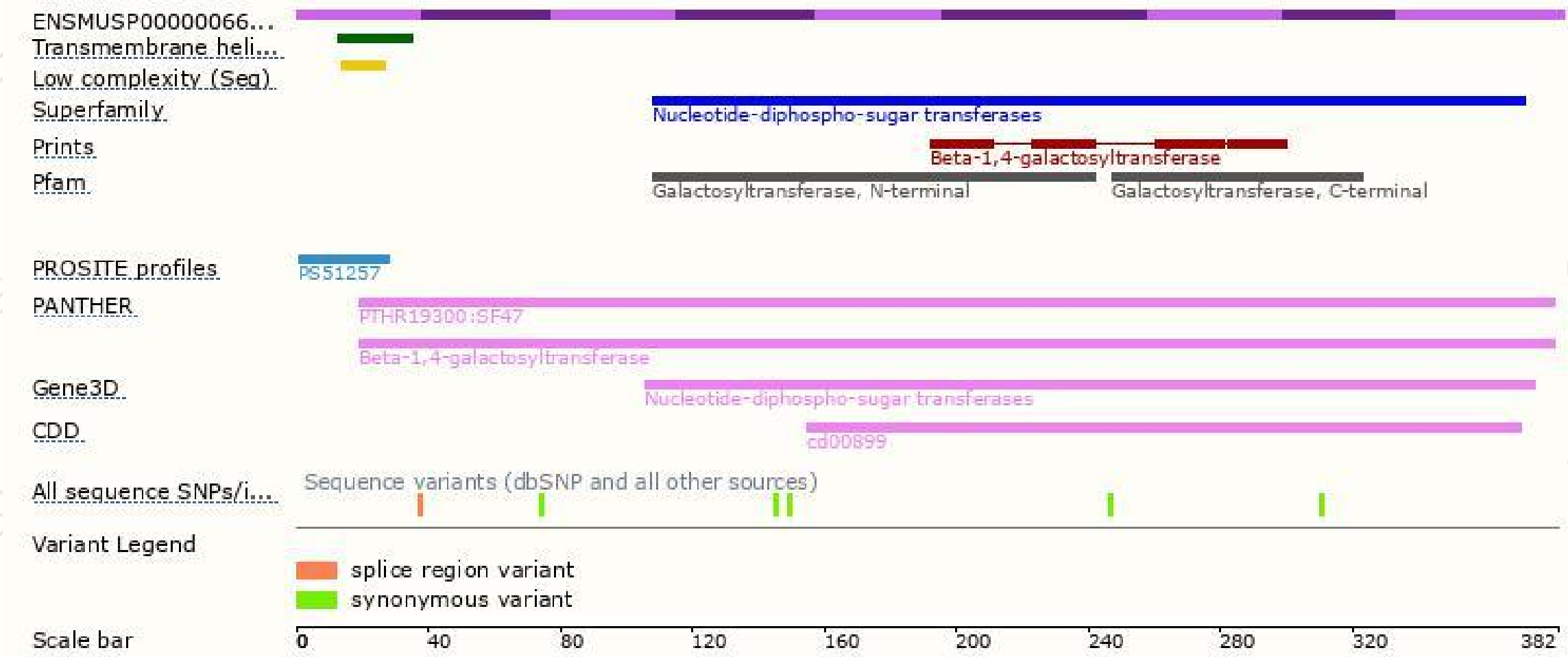
The strategy is based on the design of *B4galt6-201* transcript, The transcription is shown below



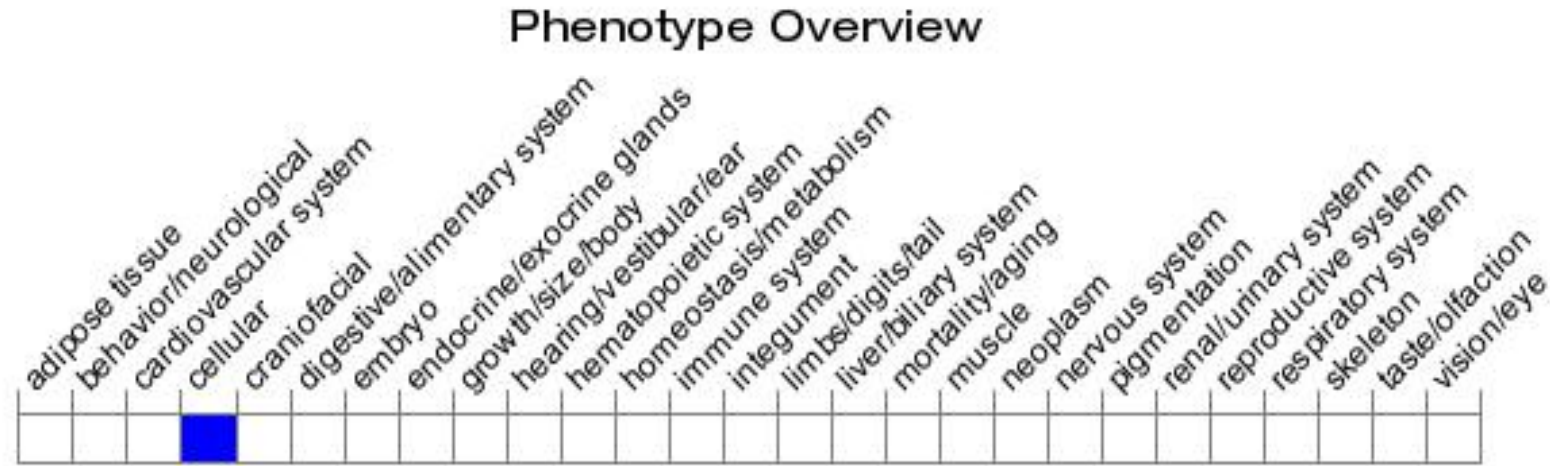
Genomic location distribution



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 disruptions in this gene display a normal phenotype with reduced lactosylceramide synthase in MEFs.

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

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