

# ***B4galt2* Cas9-KO Strategy**

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# Project Overview

**Project Name**

***B4galt2***

**Project type**

**Cas9-KO**

**Strain background**

**C57BL/6JGpt**

# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *B4galt2* gene. The schematic diagram is as follows:



- The *B4galt2* gene has 7 transcripts. According to the structure of *B4galt2* gene, exon5-exon7 of *B4galt2-201* (ENSMUST00000030266.11) transcript is recommended as the knockout region. The region contains 419bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *B4galt2* gene. The brief process is as follows: CRISPR/Cas9 system

- According to the existing MGI data, Mice homozygous for a null allele exhibit decreased brain weight, ectopic Purkinje cells in the cerebellum, and impaired spatial learning and coordination.
- Some amino acids will remain at the N-terminus and some functions may be retained.
- The effect of transcripts 205,207 is unknown.
- The *B4galt2* gene is located on the Chr4. 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)

## B4galt2 UDP-Gal:betaGlcNAc beta 1,4- galactosyltransferase, polypeptide 2 [ *Mus musculus* (house mouse) ]

Gene ID: 53418, updated on 12-Aug-2019

### Summary

Official Symbol	B4galt2 provided by <a href="#">MGI</a>
Official Full Name	UDP-Gal:betaGlcNAc beta 1,4- galactosyltransferase, polypeptide 2 provided by <a href="#">MGI</a>
Primary source	<a href="#">MGI:MGI:1858493</a>
See related	<a href="#">Ensembl:ENSMUSG00000028541</a>
Gene type	protein coding
RefSeq status	VALIDATED
Organism	<a href="#">Mus musculus</a>
Lineage	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
Also known as	Ggtb2
Expression	Broad expression in CNS E18 (RPKM 21.0), ovary adult (RPKM 20.1) and 23 other tissues <a href="#">See more</a>
Orthologs	<a href="#">human</a> <a href="#">all</a>

### Genomic context

Location: 4; 4 D2.1

See B4galt2 in [Genome Data Viewer](#)

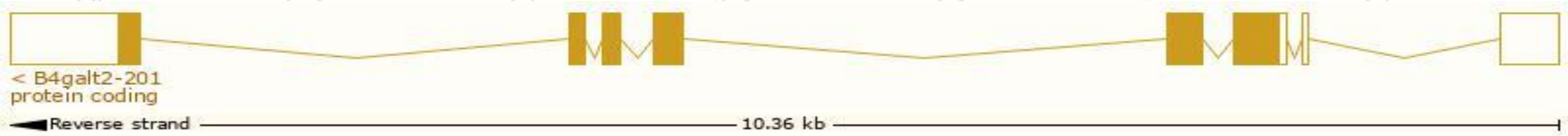
Exon count: 10

# Transcript information (Ensembl)

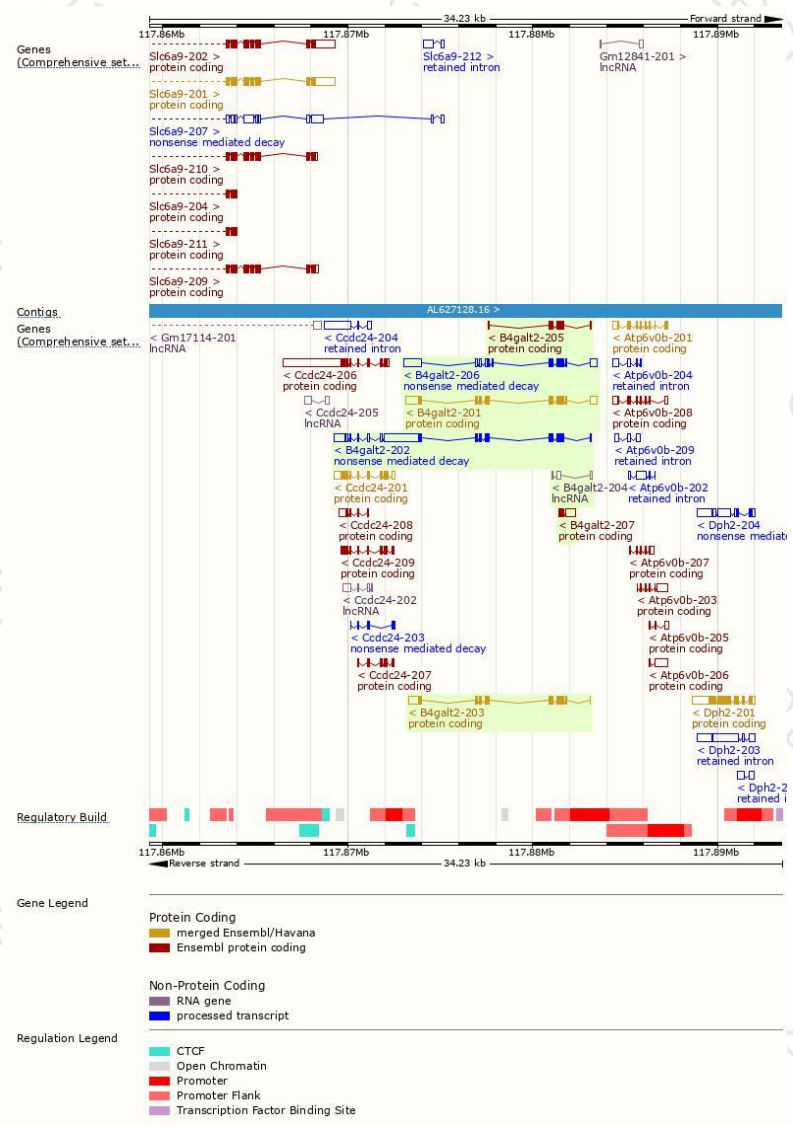
The gene has 7 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
B4galt2-201	<a href="#">ENSMUST00000030266.11</a>	2312	<a href="#">369aa</a>	Protein coding	<a href="#">CCDS18539</a>	<a href="#">B1ASJ2 Q9Z2Y2</a>	TSL:1 GENCODE basic APPRIS P1
B4galt2-203	<a href="#">ENSMUST00000106421.8</a>	1801	<a href="#">369aa</a>	Protein coding	<a href="#">CCDS18539</a>	<a href="#">B1ASJ2 Q9Z2Y2</a>	TSL:1 GENCODE basic APPRIS P1
B4galt2-207	<a href="#">ENSMUST00000171548.1</a>	794	<a href="#">75aa</a>	Protein coding	-	<a href="#">E9PY89</a>	CDS 3' incomplete TSL:3
B4galt2-205	<a href="#">ENSMUST00000153358.2</a>	696	<a href="#">196aa</a>	Protein coding	-	<a href="#">A0A0A0MQK1</a>	CDS 3' incomplete TSL:3
B4galt2-202	<a href="#">ENSMUST00000084325.9</a>	4093	<a href="#">369aa</a>	Nonsense mediated decay	<a href="#">CCDS18539</a>	<a href="#">B1ASJ2 Q9Z2Y2</a>	TSL:2
B4galt2-206	<a href="#">ENSMUST00000167443.7</a>	2466	<a href="#">224aa</a>	Nonsense mediated decay	-	<a href="#">E9Q8H8</a>	TSL:1
B4galt2-204	<a href="#">ENSMUST00000137016.1</a>	396	No protein	lncRNA	-	-	TSL:3

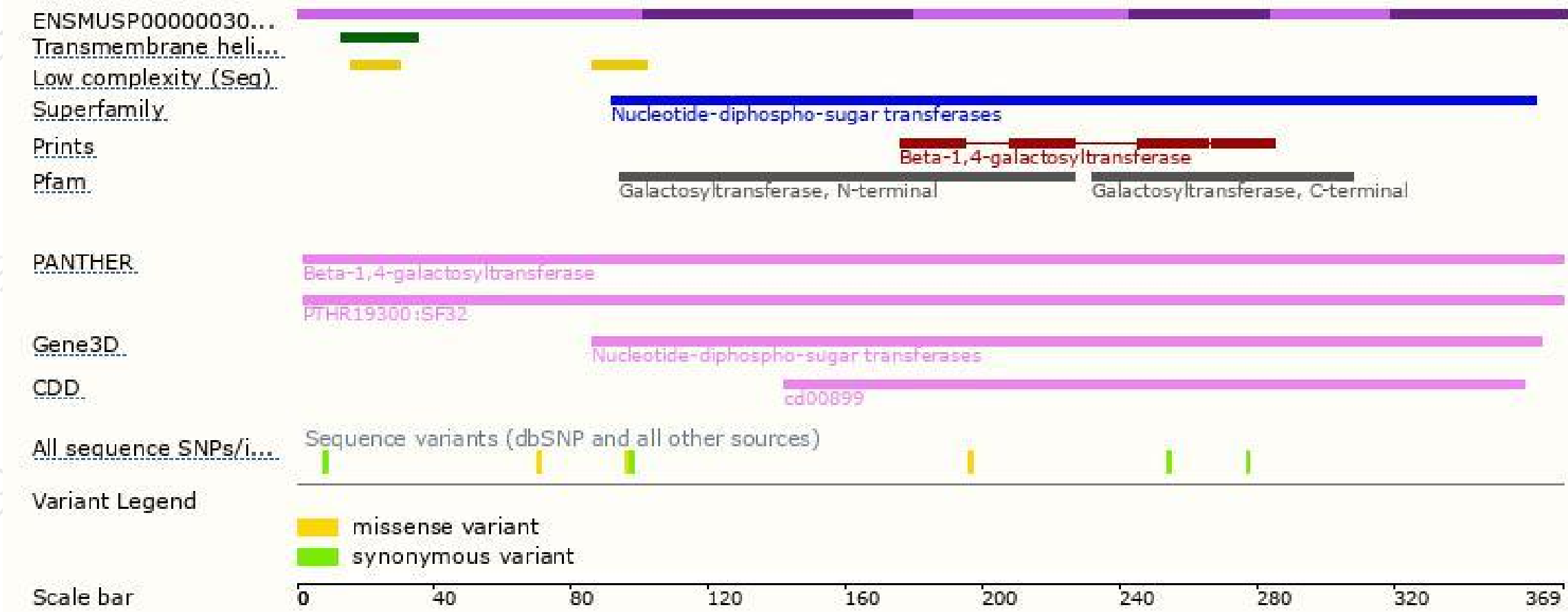
The strategy is based on the design of *B4galt2-201* transcript,The transcription is shown below



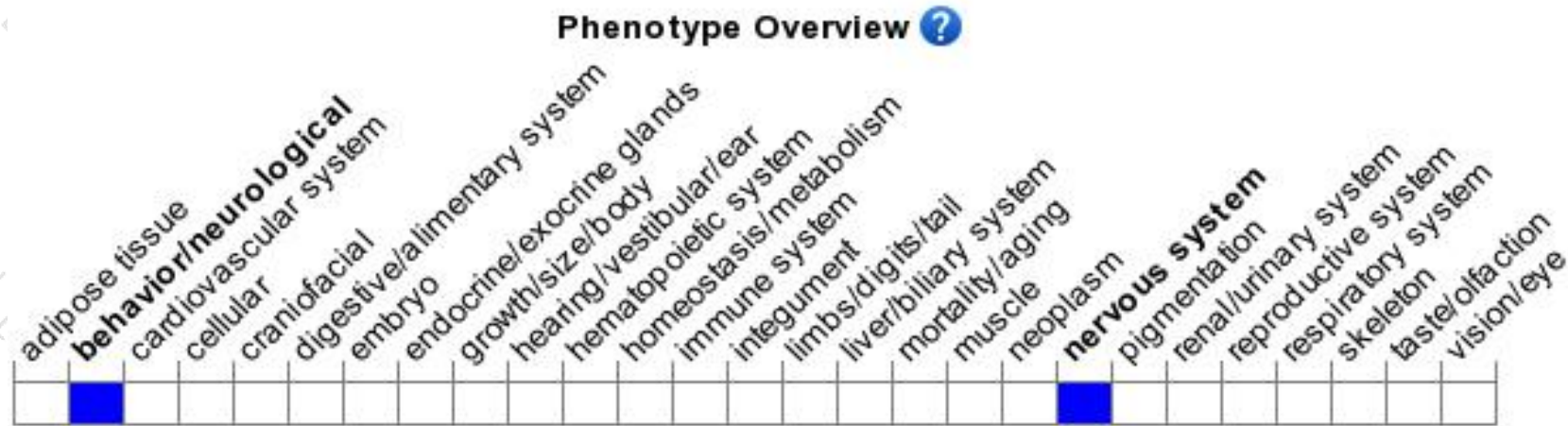
# Genomic location distribution



# 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 null allele exhibit decreased brain weight, ectopic Purkinje cells in the cerebellum, and impaired spatial learning and coordination.

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

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