

Agl Cas9-KO Strategy

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Reviewer:

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Design Date:

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Project Overview

Project Name

Agl

Project type

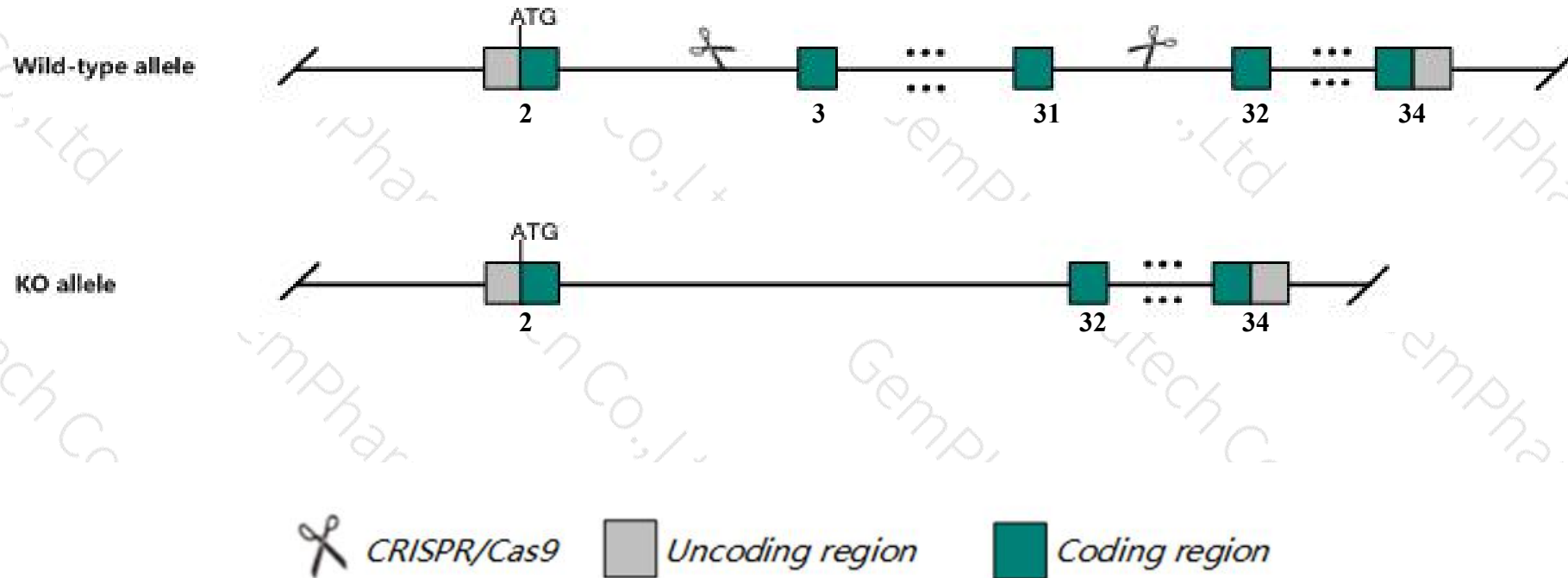
Cas9-KO

Strain background

C57BL/6JGpt

Knockout strategy

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



- The *Agl* gene has 8 transcripts. According to the structure of *Agl* gene, exon3-exon31 of *Agl-201* (ENSMUST00000040603.13) transcript is recommended as the knockout region. The region contains 4177bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Agl* gene. The brief process is as follows: CRISPR/Cas9 system will

- According to the existing MGI data, Homozygous inactivation of this gene leads to hypoglycemia, altered blood biochemistry, severe hepatomegaly, glycogen accumulation in the liver, heart, skeletal muscle and other tissues, motor impairment, and premature death.
- The *Agl* gene is located on the Chr3. 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)

Agl amylo-1,6-glucosidase, 4-alpha-glucanotransferase [Mus musculus (house mouse)]

Gene ID: 77559, updated on 31-Jan-2019

Summary



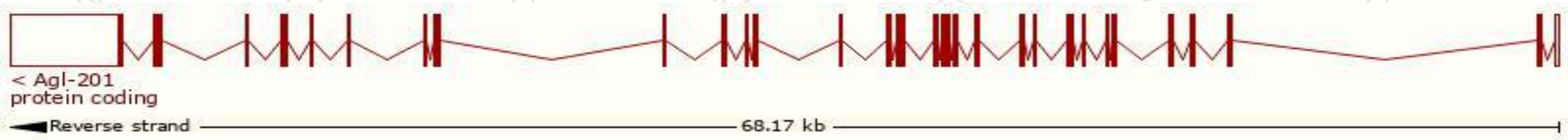
| | |
|---------------------------|---|
| Official Symbol | Agl provided by MGI |
| Official Full Name | amylo-1,6-glucosidase, 4-alpha-glucanotransferase provided by MGI |
| Primary source | MGI:MGI:1924809 |
| See related | Ensembl:ENSMUSG00000033400 |
| Gene type | protein coding |
| RefSeq status | 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 | 1110061O17Rik, 9430004C13Rik, 9630046L06Rik, AI850929, C77197 |
| Expression | Ubiquitous expression in heart adult (RPKM 13.2), liver E18 (RPKM 6.5) and 26 other tissues See more |
| Orthologs | human all |

Transcript information (Ensembl)

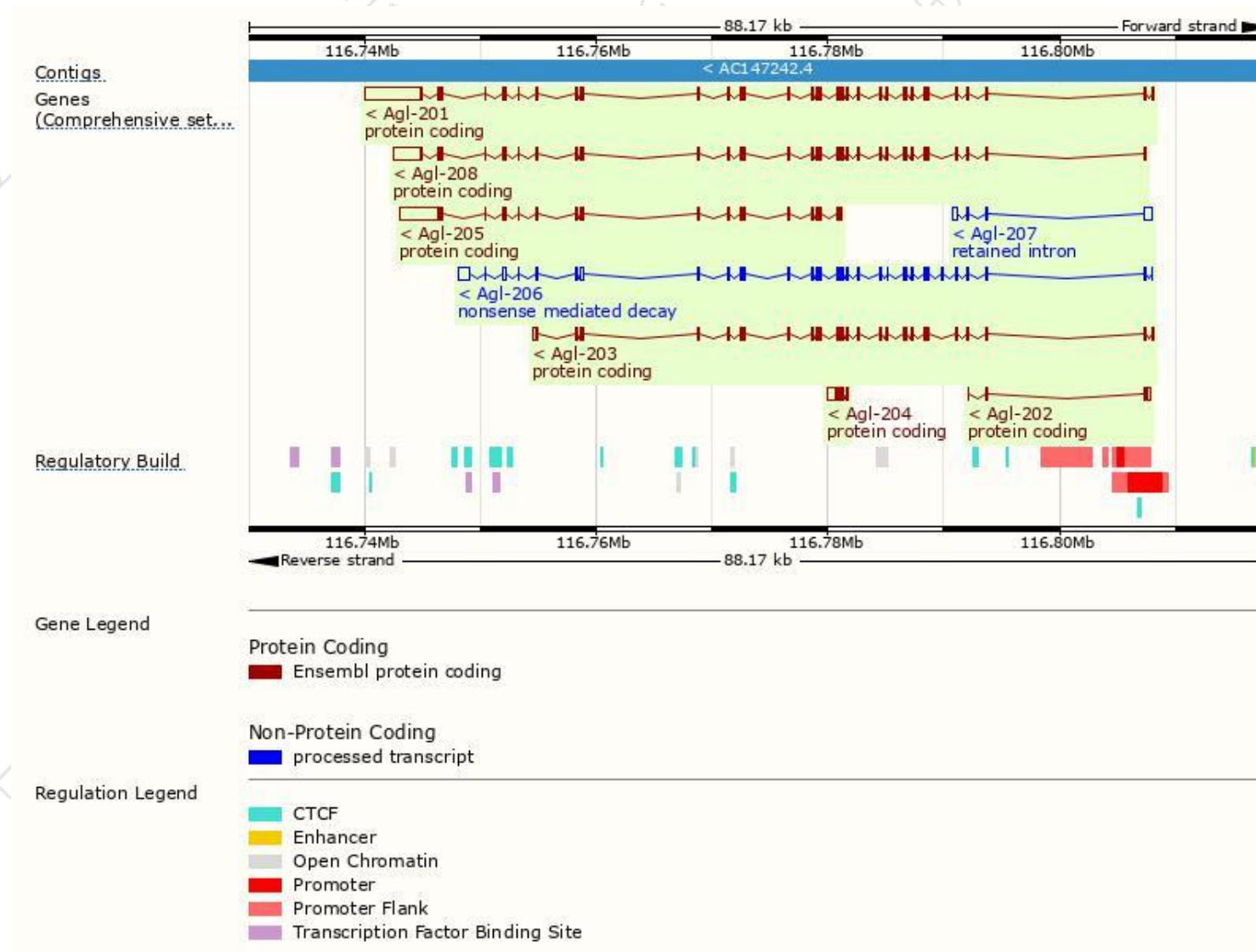
The gene has 8 transcripts,all transcripts are shown below:

| Name | Transcript ID | bp | Protein | Biotype | CCDS | UniProt | Flags |
|---------|---------------------------------------|------|------------------------|-------------------------|---------------------------|----------------------------|-------------------------------|
| Agl-201 | ENSMUST00000040603.13 | 9625 | 1532aa | Protein coding | CCDS38613 | F8VPN4 | TSL:5 GENCODE basic APPRIS P1 |
| Agl-208 | ENSMUST00000162792.7 | 6978 | 1532aa | Protein coding | CCDS38613 | F8VPN4 | TSL:2 GENCODE basic APPRIS P1 |
| Agl-205 | ENSMUST00000160484.5 | 5817 | 830aa | Protein coding | - | F6XXE6 | CDS 5' incomplete TSL:1 |
| Agl-203 | ENSMUST00000159742.7 | 4344 | 1279aa | Protein coding | - | A0A0G2JGI9 | TSL:1 GENCODE basic |
| Agl-204 | ENSMUST00000159995.1 | 1285 | 198aa | Protein coding | - | F7CSZ6 | CDS 5' incomplete TSL:1 |
| Agl-202 | ENSMUST00000159670.2 | 722 | 116aa | Protein coding | - | E0CX86 | CDS 3' incomplete TSL:5 |
| Agl-206 | ENSMUST00000161336.7 | 5365 | 233aa | Nonsense mediated decay | - | E0CYU6 | TSL:5 |
| Agl-207 | ENSMUST00000162040.1 | 1572 | No protein | Retained intron | - | - | TSL:1 |

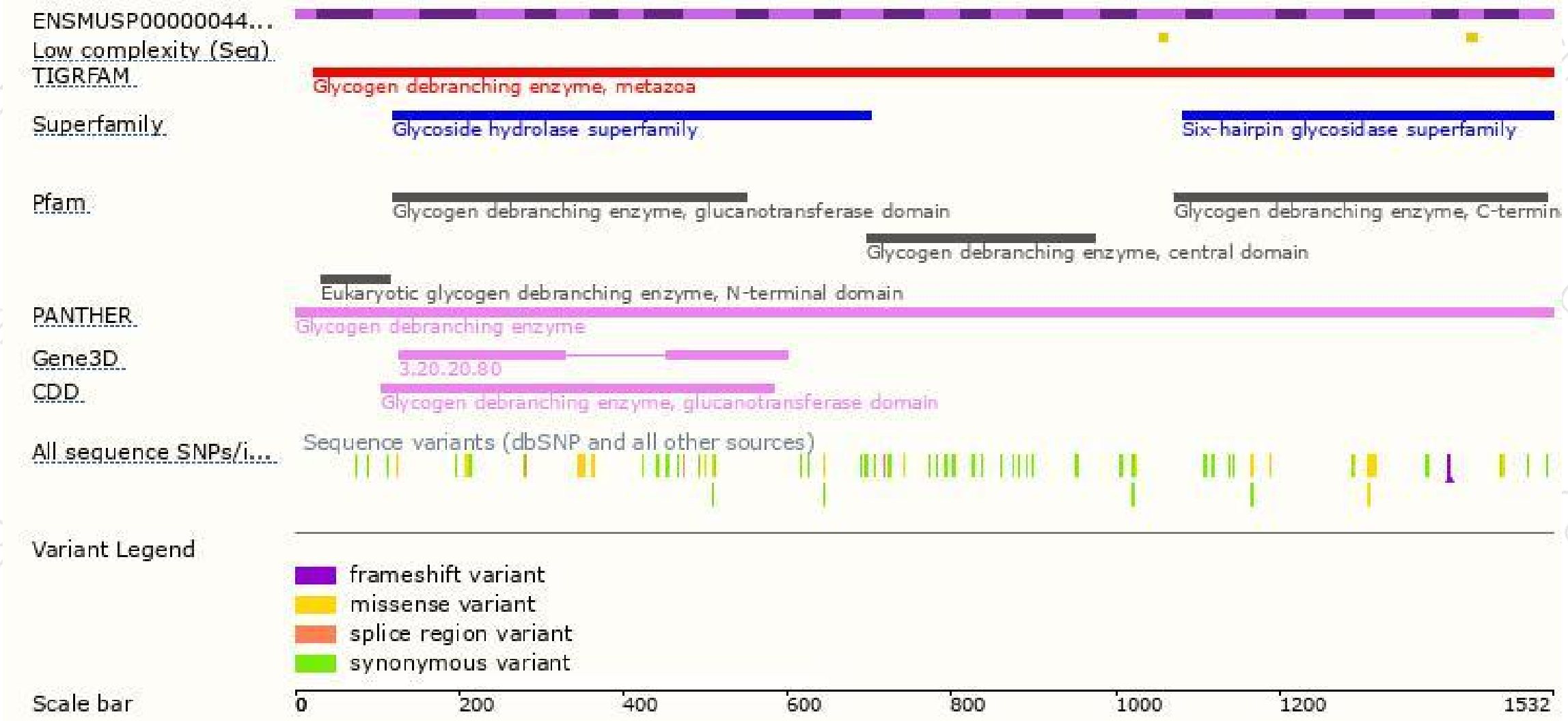
The strategy is based on the design of *Agl-201* transcript,The transcription is shown below



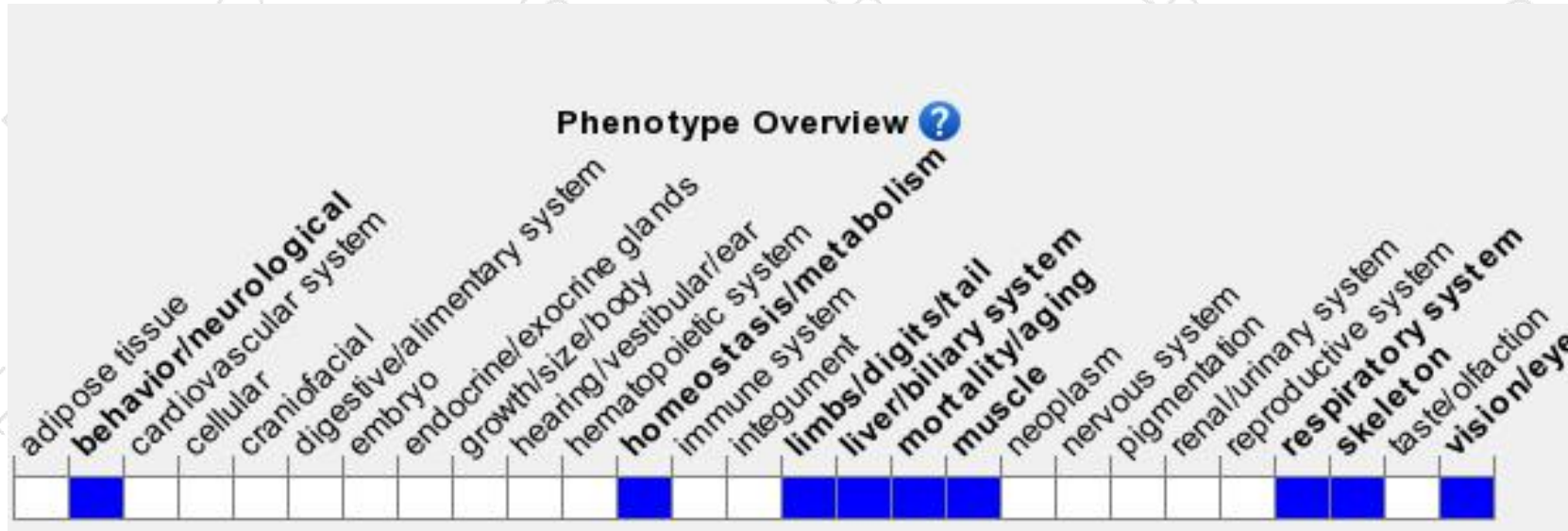
Genomic location distribution



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 inactivation of this gene leads to hypoglycemia, altered blood biochemistry, severe hepatomegaly, glycogen accumulation in the liver, heart, skeletal muscle and other tissues, motor impairment, and premature death.

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

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