

Neu1 Cas9-CKO Strategy

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Project Overview

Project Name

Neu1

Project type

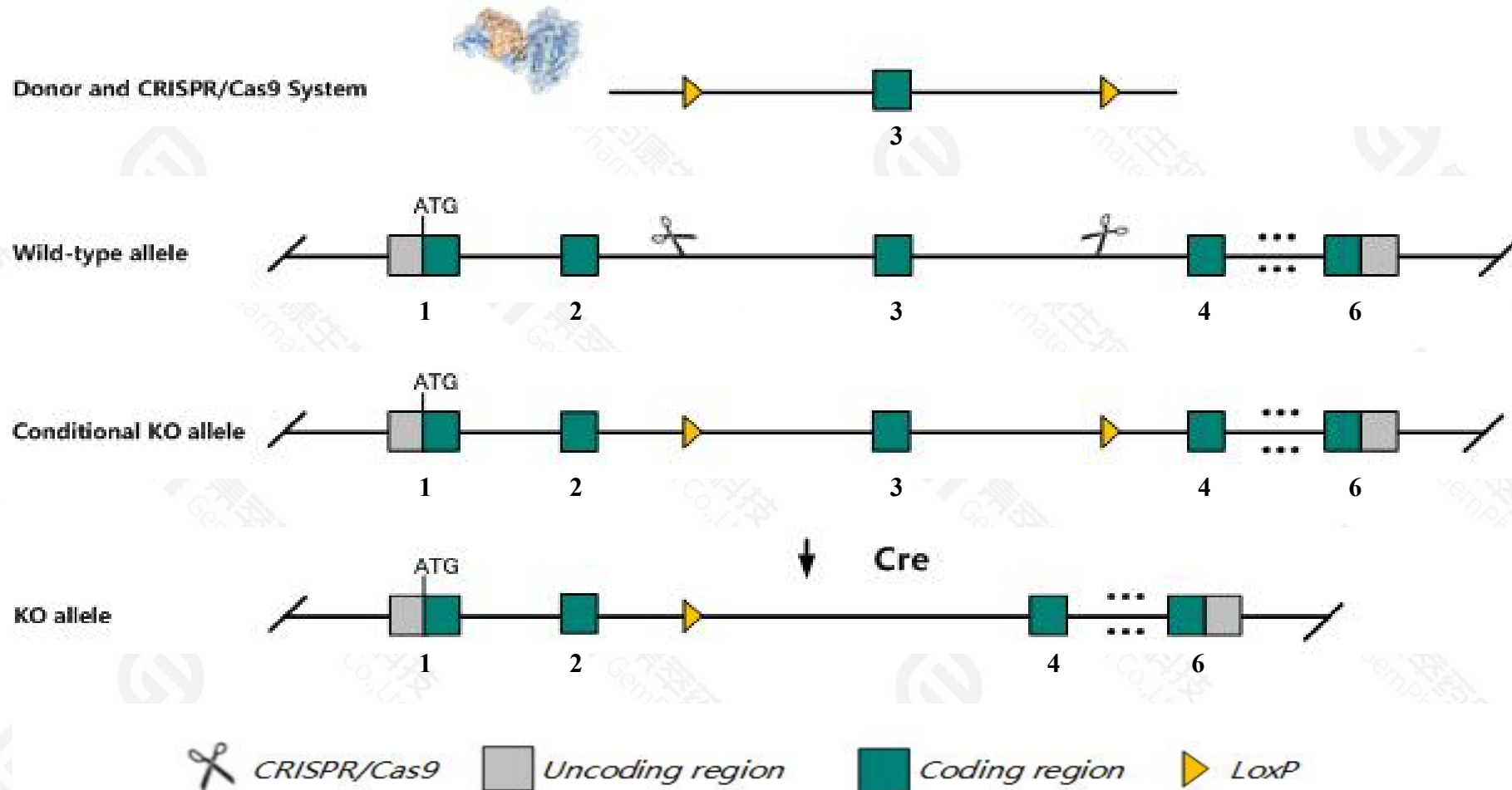
Cas9-CKO

Strain background

C57BL/6JGpt

Conditional Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Neu1* gene. The schematic diagram is as follows:



Technical routes

- The *Neu1* gene has 3 transcripts. According to the structure of *Neu1* gene, exon3 of *Neu1-201*(ENSMUST00000007253.6) transcript is recommended as the knockout region. The region contains 263bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Neu1* 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.

- According to the existing MGI data, nullizygous mice develop features of early-onset lysosomal storage disease (sialidosis), including severe nephropathy, edema, splenomegaly, kyphosis and oligosacchariduria, and display myoclonus, lordosis, extramedullary hematopoiesis, dyspnea, weight loss, gait defects, tremors and premature death.
- The Intron2 and Intron3 are only 457bp and 1174bp, loxp insertion may affect mRNA splicing.
- The *Neu1* gene is located on the Chr17. 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)

Neu1 neuraminidase 1 [Mus musculus (house mouse)]

Gene ID: 18010, updated on 13-Mar-2020

Summary

Official Symbol Neu1 provided by [MGI](#)

Official Full Name neuraminidase 1 provided by [MGI](#)

Primary source [MGI:MGI:97305](#)

See related [Ensembl:ENSMUSG00000007038](#)

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 AA407268, AA407316, Aglp, Apl, Bat-7, Bat7, G9, Map-2, Neu, Neu-1

Expression Broad expression in kidney adult (RPKM 114.3), colon adult (RPKM 62.8) and 20 other tissues [See more](#)

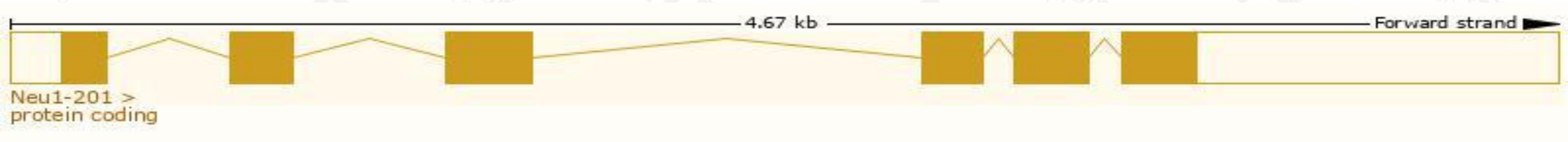
Orthologs [human](#) [all](#)

Transcript information (Ensembl)

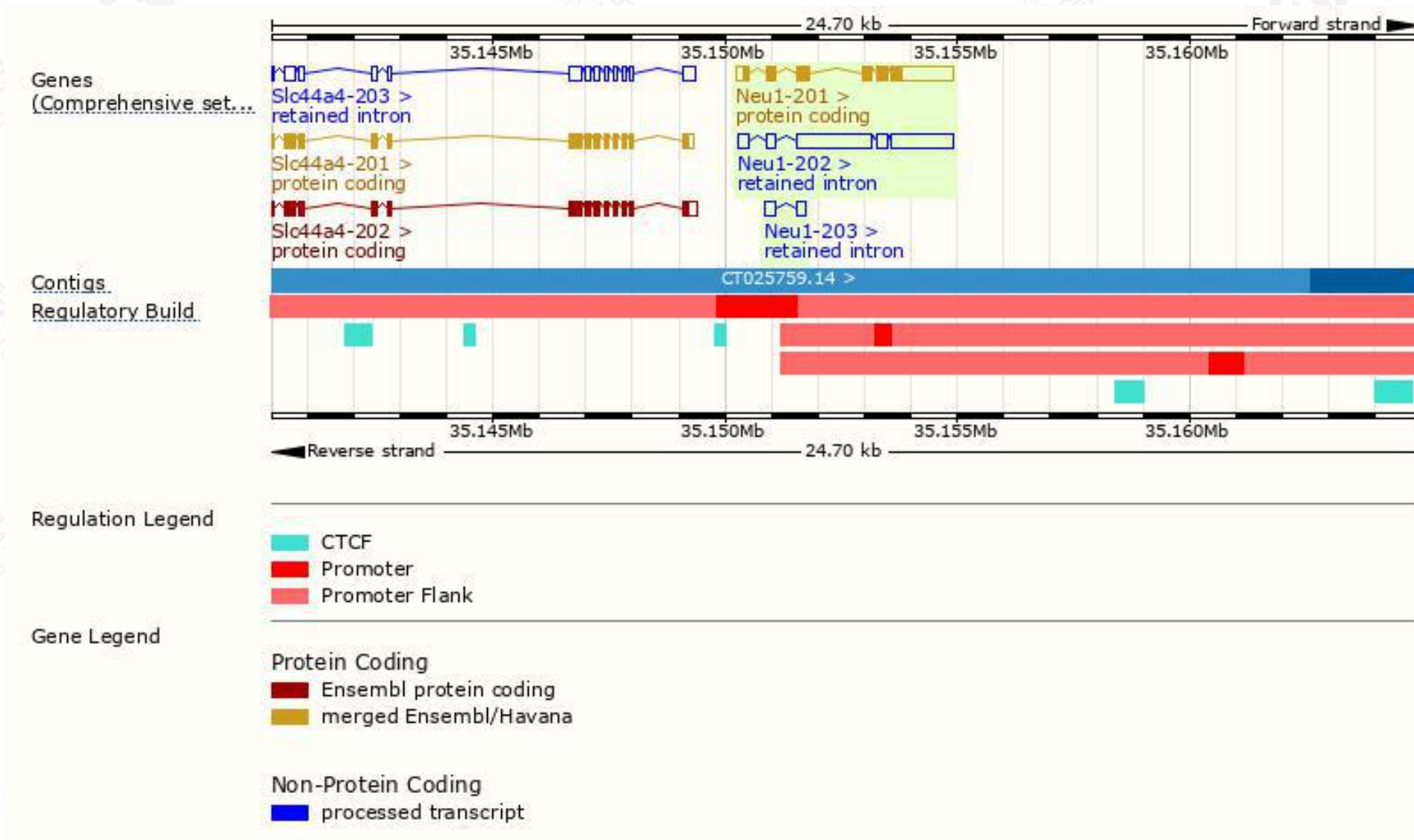
The gene has 3 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Neu1-201	ENSMUST00000007253.5	2474	409aa	Protein coding	CCDS28668	Q35657 Q3UL64	TSL:1 GENCODE basic APPRIS is a system to annotate alternatively spliced transcripts based on a range of computational methods to identify the most functionally important transcript(s) of a gene. APPRIS P1
Neu1-202	ENSMUST00000173269.1	3619	No protein	Retained intron	-	-	TSL:1
Neu1-203	ENSMUST00000174715.1	446	No protein	Retained intron	-	-	TSL:2

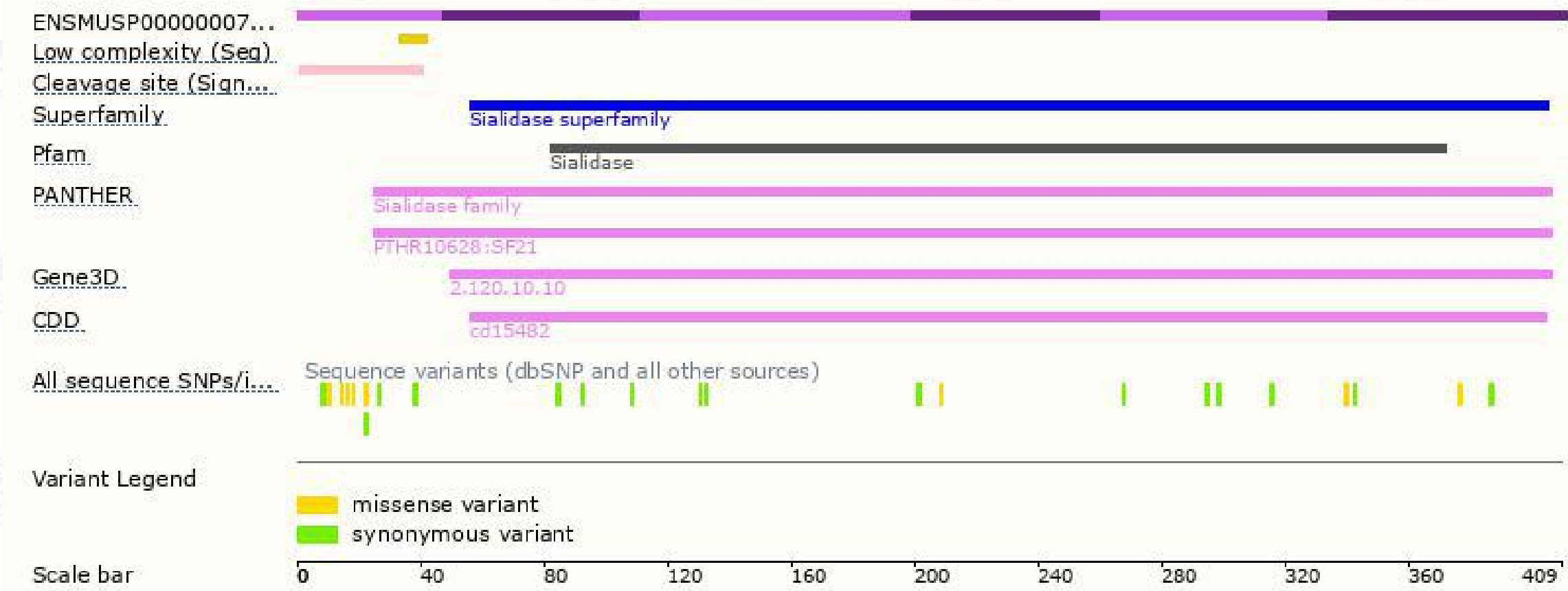
The strategy is based on the design of *Neu1-201* transcript,the transcription is shown below:



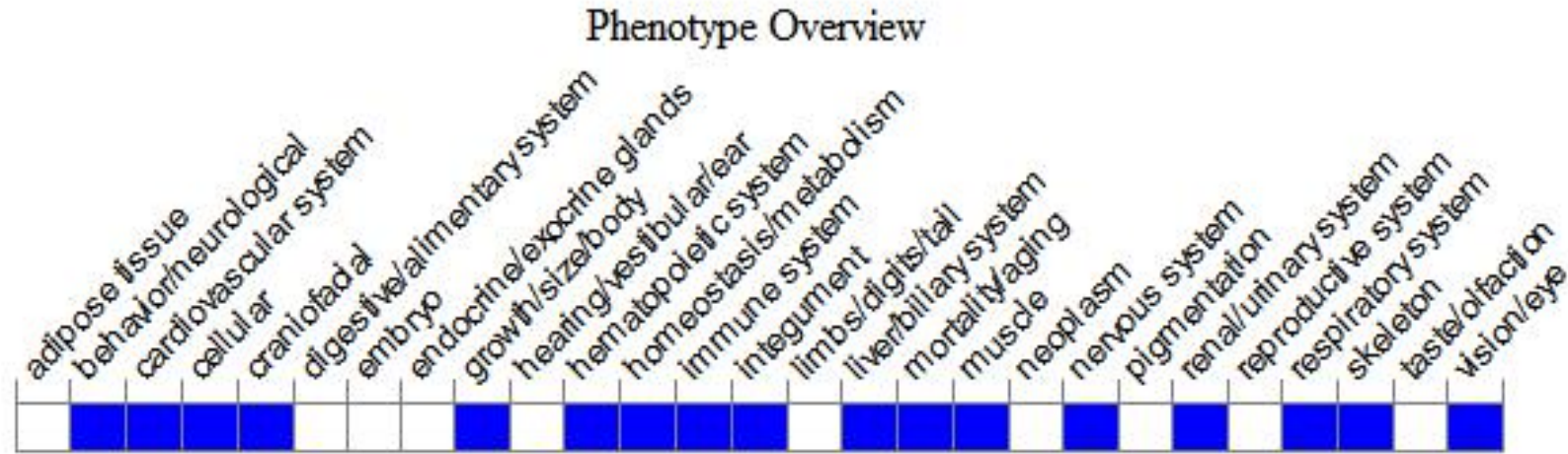
Genomic location distribution



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, nullizygous mice develop features of early-onset lysosomal storage disease (sialidosis), including severe nephropathy, edema, splenomegaly, kyphosis and oligosacchariduria, and display myoclonus, lordosis, extramedullary hematopoiesis, dyspnea, weight loss, gait defects, tremors and premature death.

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