Slc10a2 Cas9-CKO Strategy

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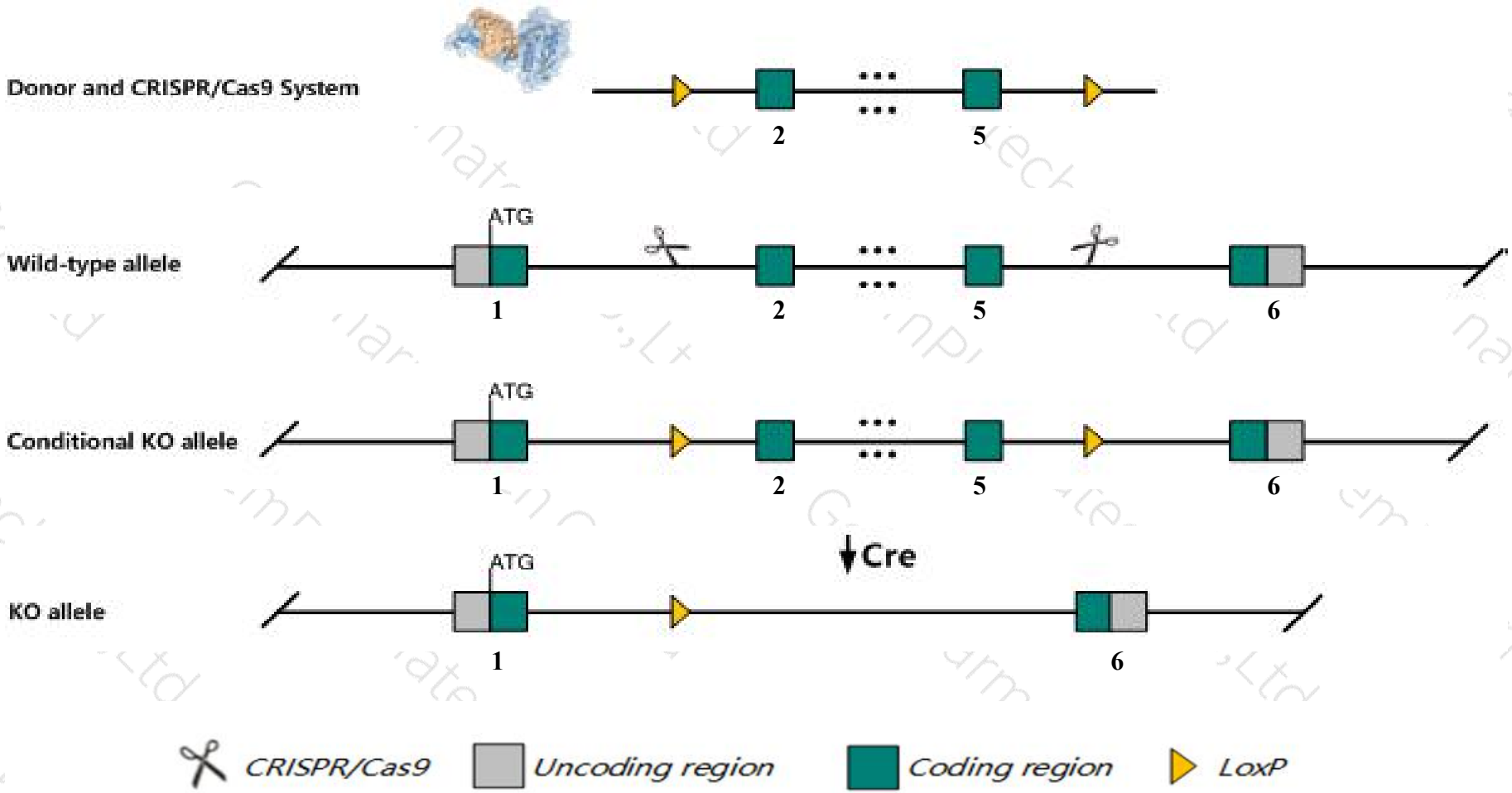
Project Overview

Project Name	***Slc10a2***
Project type	**Cas9-CKO**
Strain background	**C57BL/6JGpt**

Conditional Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Slc10a2* gene. The schematic diagram is as follows:



Technical routes

- The *Slc10a2* gene has 1 transcript. According to the structure of *Slc10a2* gene, exon2-exon5 of *Slc10a2-201* (ENSMUST00000023835.2) transcript is recommended as the knockout region.The region contains 542bp coding sequence. Knock out the region will result in disruption of protein function.

- In this project we use CRISPR/Cas9 technology to modify *Slc10a2* 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.

Notice

- According to the existing MGI data,Mice homozygous for disruptions in this gene are essentially indistinguishable from wild-type in terms of survival, gross appearance and behavior. However, they do have defects in lipid absorption from the intestine.
- The N-terminal of *Slc10a2* gene will remain several amino acids ,it may remain the partial function of *Slc10a2* gene.
- The *Slc10a2* gene is located on the Chr8. 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）

集萃药康 GemPharmatech

Slc10a2 solute carrier family 10, member 2 [*Mus musculus* (house mouse)]

Gene ID: 20494, updated on 22-Oct-2019

Summary

Official Symbol	Slc10a2 provided by MGI
Official Full Name	solute carrier family 10, member 2 provided by MGI
Primary source	MGI:MGI:1201406
See related	Ensembl:ENSMUSG00000023073
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	ASBT; IBAT; ISBT; AI605518; 9130221J18Rik
Expression	Biased expression in large intestine adult (RPKM 20.9), kidney adult (RPKM 7.5) and 2 other tissues See more
Orthologs	human all

Genomic context

Location: 8 A1.1; 8 2.16 cM

See Slc10a2 in Genome Data Viewer

Exon count: 6

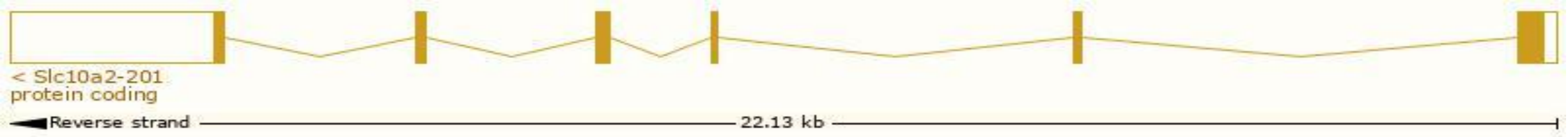
Annotation release	Status	Assembly	Chr	Location
108	current	GRCm38.p6 (GCF_000001635.26)	8	NC_000074.6 (5083219..5105287, complement)
Build 37.2	previous assembly	MGSCv37 (GCF_000001635.18)	8	NC_000074.5 (5085623..5105232, complement)

Transcript information（Ensembl）

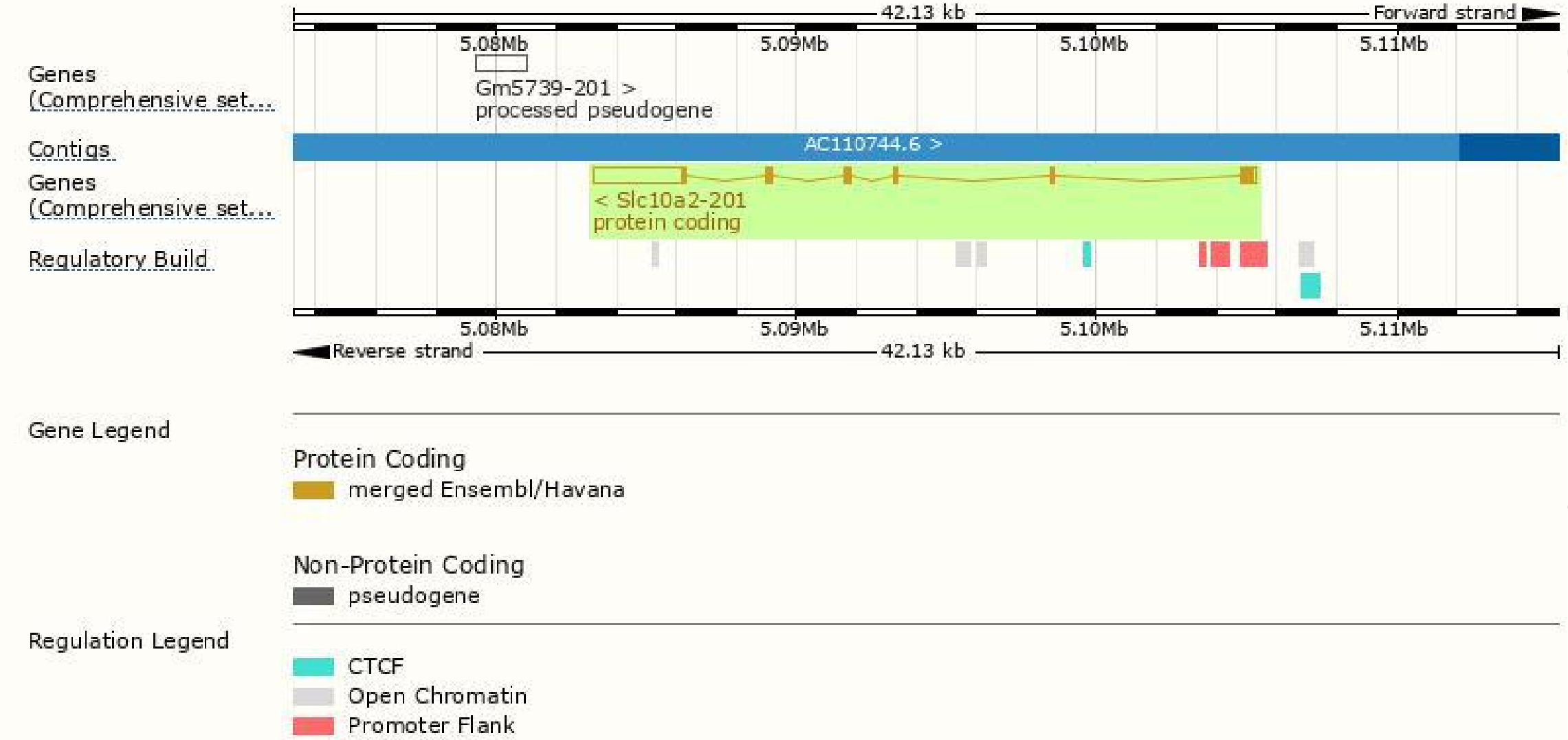
The gene has 1 transcript,and the transcript is shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Slc10a2-201	ENSMUST00000023835.2	4152	348aa	Protein coding	CCDS22089	P70172 Q0VBB8	TSL:1 GENCODE basic APPRIS P1

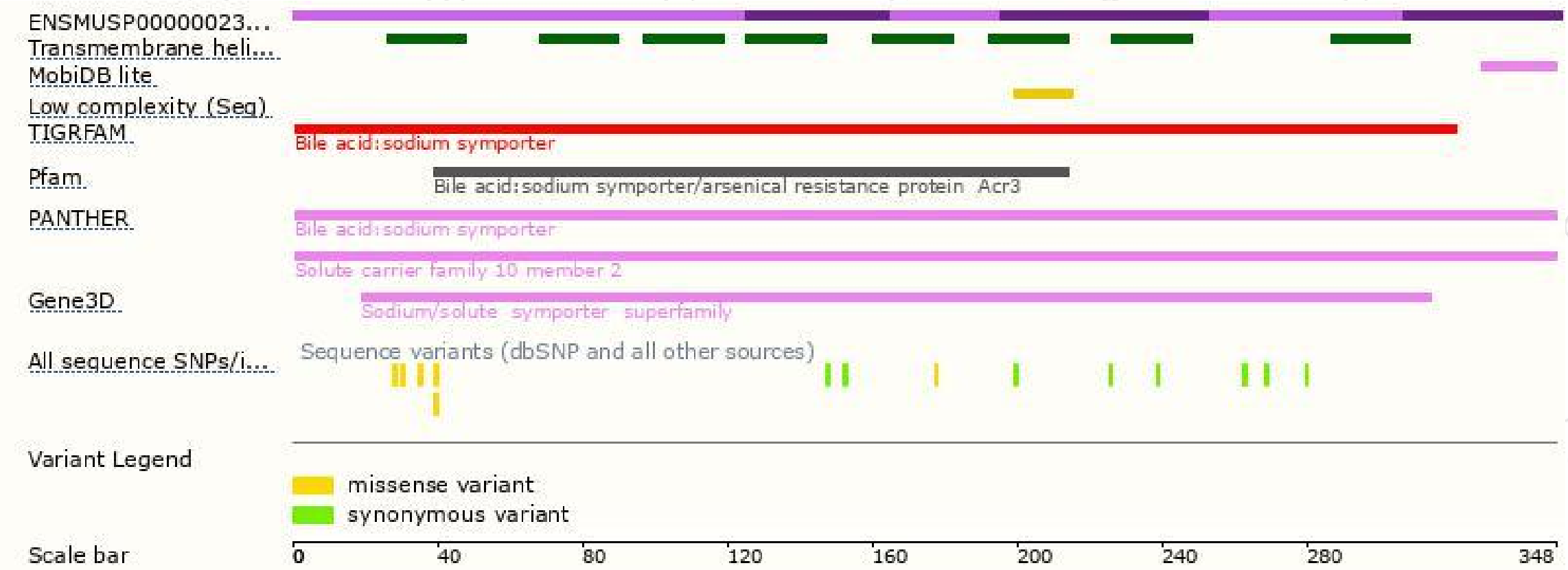
The strategy is based on the design of *Slc10a2-201* transcript,The transcription is shown below



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/).

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

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