

# *Scaf11* Cas9-KO Strategy

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# Project Overview

**Project Name**

*Scaf11*

**Project type**

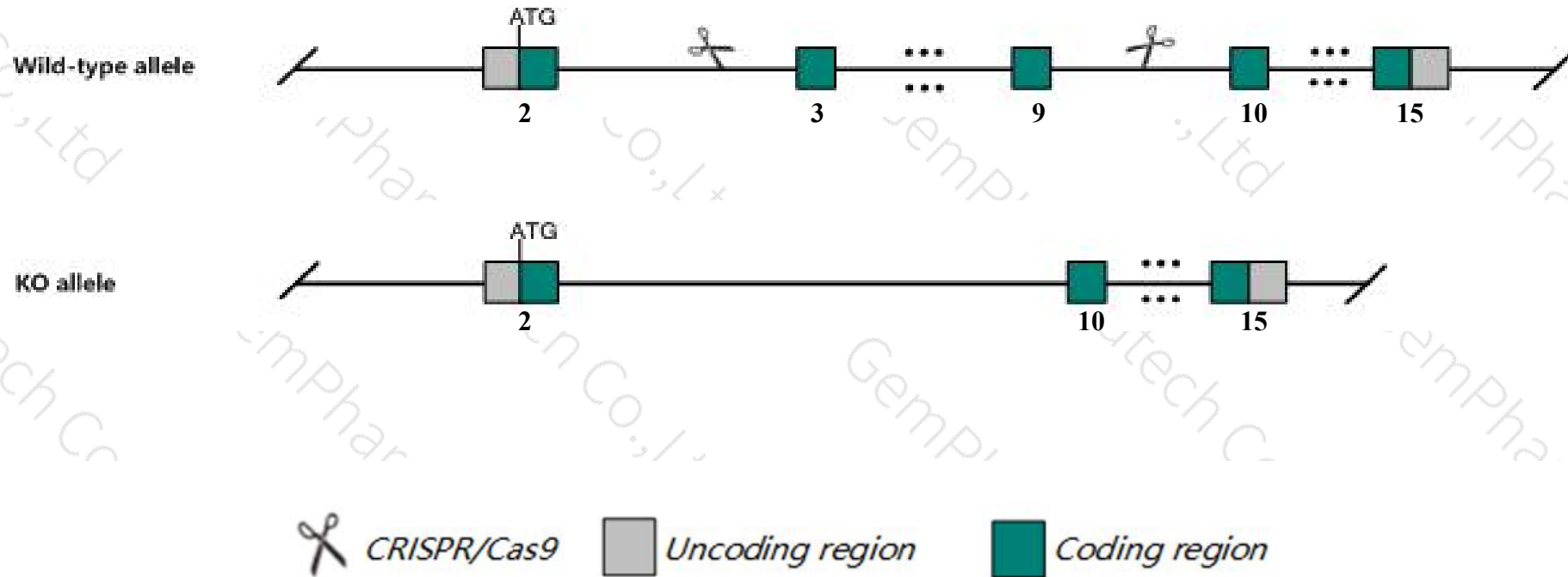
**Cas9-KO**

**Strain background**

**C57BL/6JGpt**

# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Scaf11* gene. The schematic diagram is as follows:



- The *Scaf11* gene has 6 transcripts. According to the structure of *Scaf11* gene, exon3-exon9 of *Scaf11-201* (ENSMUST00000047835.7) transcript is recommended as the knockout region. The region contains 667bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Scaf11* gene. The brief process is as follows: CRISPR/Cas9 system

- Transcript *Scaf11*-203 may not be affected . And the effect on transcript *Scaf11*-204&205&206 is unknown.
- The *Scaf11* gene is located on the Chr15. 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)

## Scaf11 SR-related CTD-associated factor 11 [ *Mus musculus* (house mouse) ]

Gene ID: 72193, updated on 12-Aug-2019

### Summary

**Official Symbol** Scaf11 provided by [MGI](#)  
**Official Full Name** SR-related CTD-associated factor 11 provided by [MGI](#)  
**Primary source** [MGI:MGI:1919443](#)  
**See related** [Ensembl:ENSMUSG00000033228](#)  
**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** SIP1; CASP11; SRRP129; Sfrs2ip; Srsf2ip; AI462454; mKIAA3013; 1110061H03Rik; 2610510E10Rik  
**Expression** Ubiquitous expression in placenta adult (RPKM 13.7), CNS E11.5 (RPKM 12.4) and 28 other tissues [See more](#)  
**Orthologs** [human](#) [all](#)

### Genomic context

Location: 15; 15 F1

See Scaf11 in [Genome Data Viewer](#)

Exon count: 16

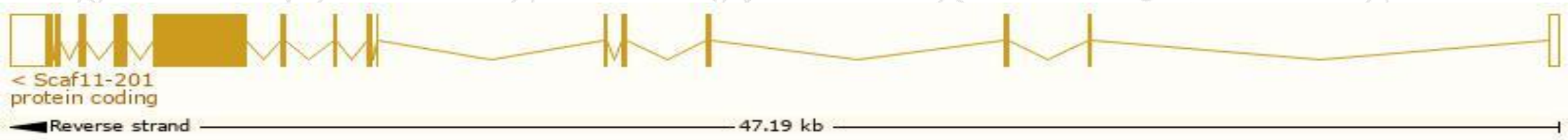
Annotation release	Status	Assembly	Chr	Location
<a href="#">108</a>	current	GRCm38.p6 ( <a href="#">GCF_000001635.26</a> )	15	NC_000081.6 (96411698..96460988, complement)
Build 37.2	previous assembly	MGSCv37 ( <a href="#">GCF_000001635.18</a> )	15	NC_000081.5 (96242129..96291274, complement)

# Transcript information (Ensembl)

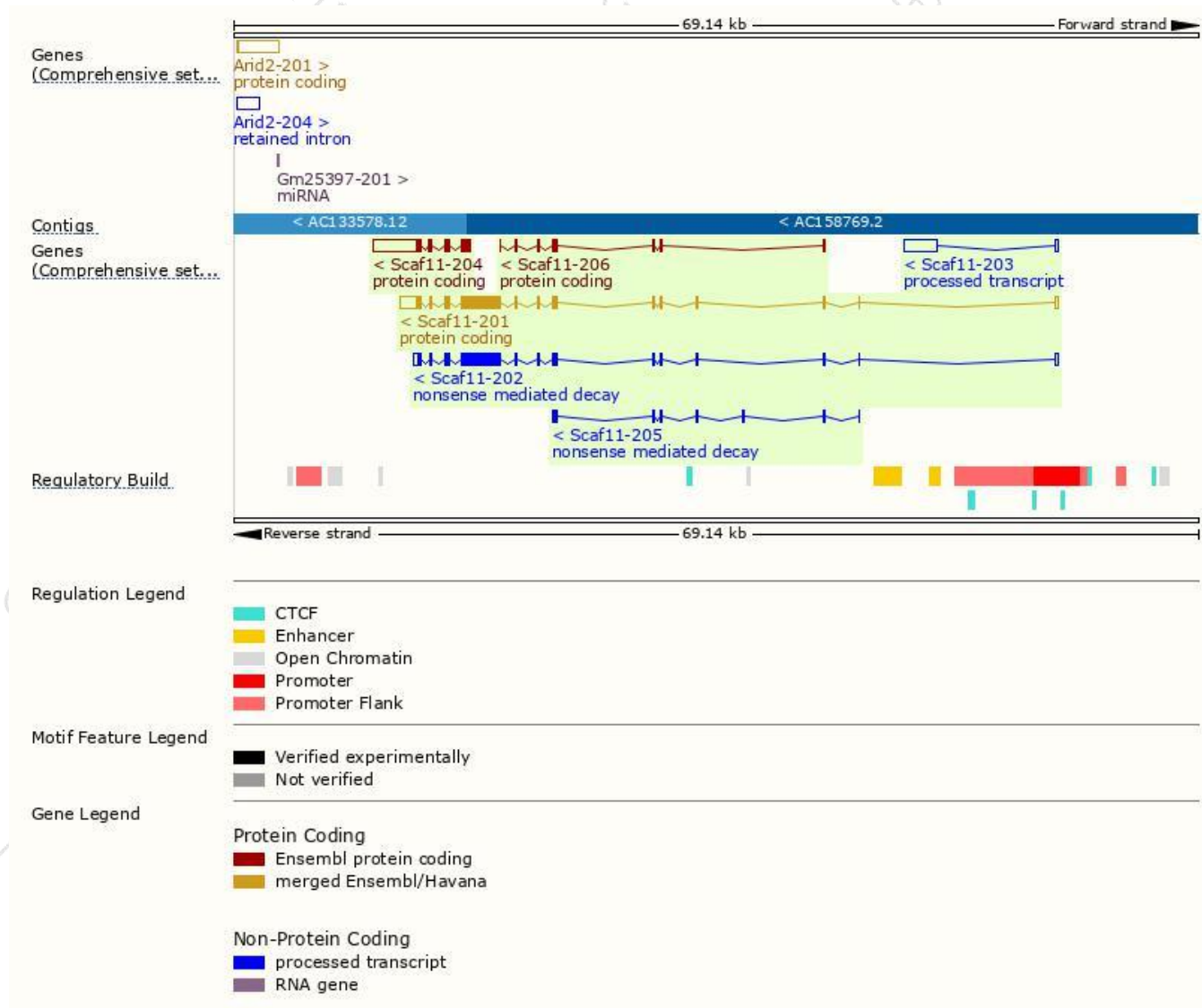
The gene has 6 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Scaf11-201	<a href="#">ENSMUST00000047835.7</a>	5766	<a href="#">1456aa</a>	Protein coding	<a href="#">CCDS37186</a>	<a href="#">E9PZM7</a>	TSL:5 GENCODE basic APPRIS P1
Scaf11-204	<a href="#">ENSMUST00000228072.1</a>	4519	<a href="#">476aa</a>	Protein coding	-	<a href="#">A0A2I3BRQ7</a>	CDS 5' incomplete
Scaf11-206	<a href="#">ENSMUST00000228535.1</a>	696	<a href="#">232aa</a>	Protein coding	-	<a href="#">A0A2I3BQX1</a>	5' and 3' truncations in transcript evidence prevent annotation of the start and the end of the CDS. CDS 5' and 3' incomplete
Scaf11-202	<a href="#">ENSMUST00000227069.1</a>	4829	<a href="#">1389aa</a>	Nonsense mediated decay	-	<a href="#">A0A2I3BQU5</a>	
Scaf11-205	<a href="#">ENSMUST00000228260.1</a>	796	<a href="#">84aa</a>	Nonsense mediated decay	-	<a href="#">A0A2I3BQ68</a>	CDS 5' incomplete
Scaf11-203	<a href="#">ENSMUST00000227133.1</a>	2599	No protein	Processed transcript	-	-	

The strategy is based on the design of *Scaf11-201* transcript,The transcription is shown below

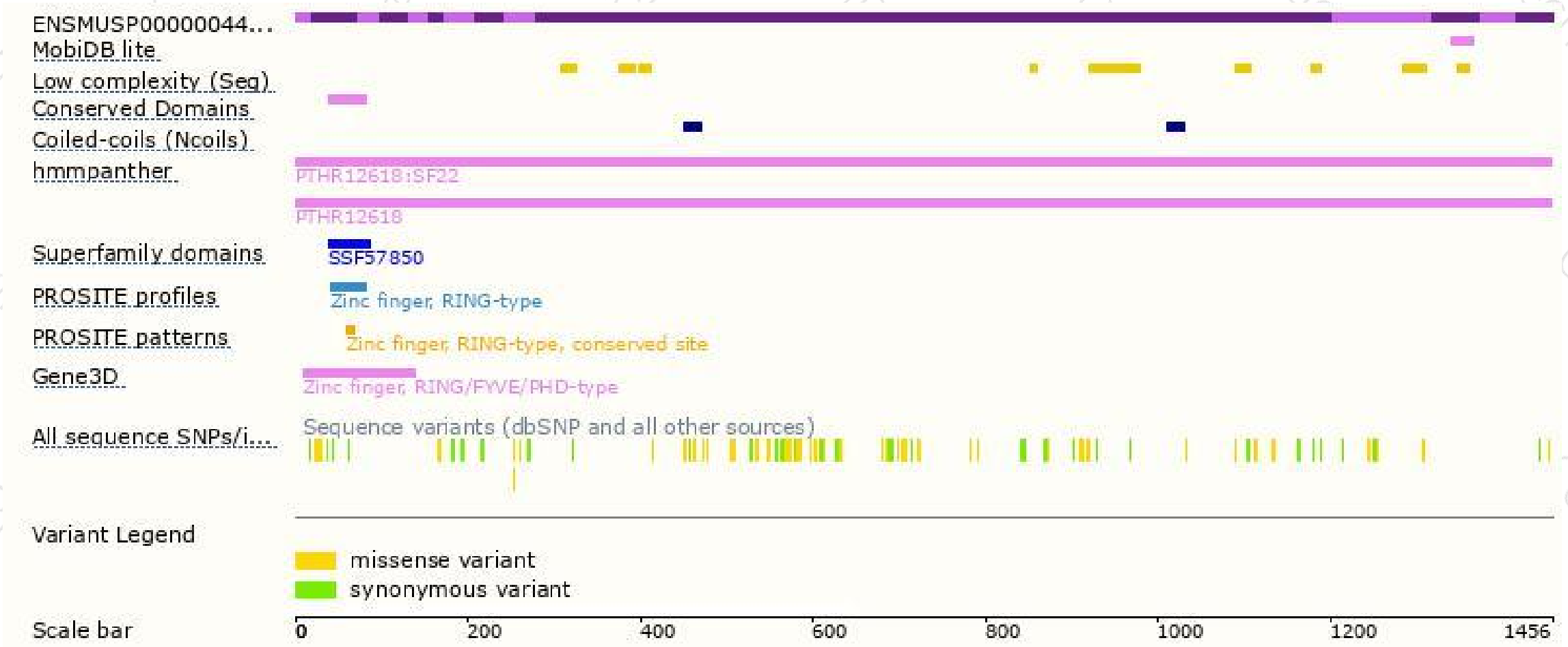


# Genomic location distribution

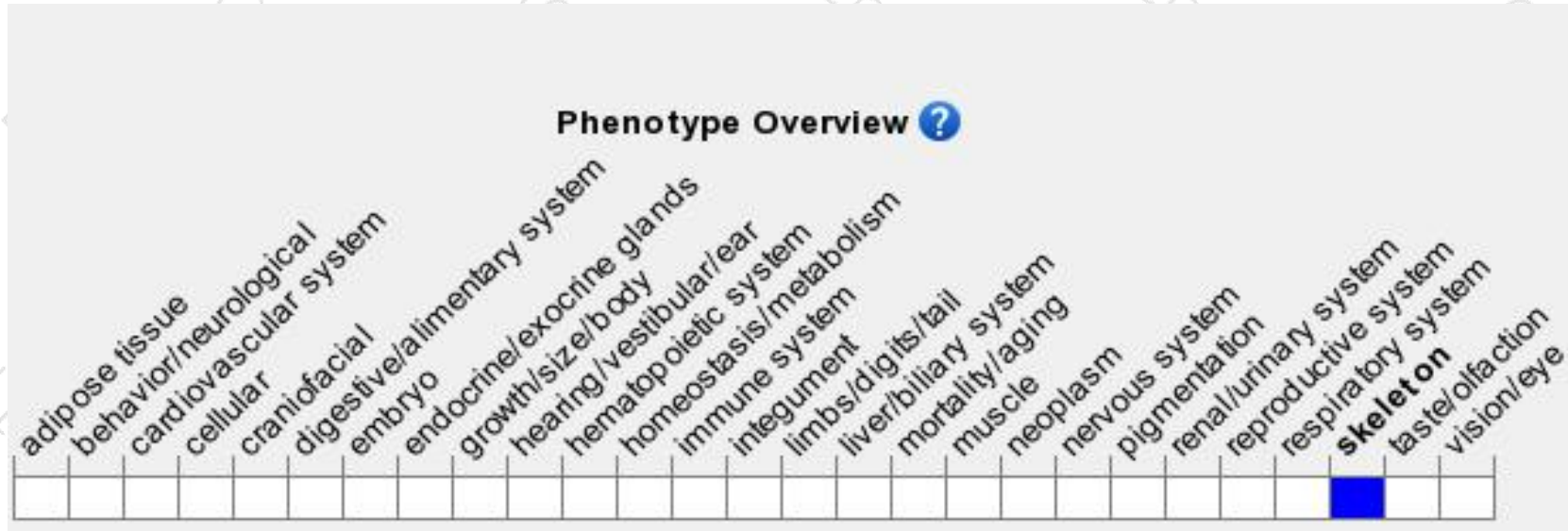




# Protein domain



# Mouse phenotype description(MGI )



Phenotypes affected by the gene are marked in blue. Data quoted from MGI database(<http://www.informatics.jax.org/>).

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

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