

# *Mapk14* Cas9-KO Strategy

**Designer:**

**Jing Jin**

**Reviewer:**

**Yang Zeng**

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# Project Overview

**Project Name**

***Mapk14***

**Project type**

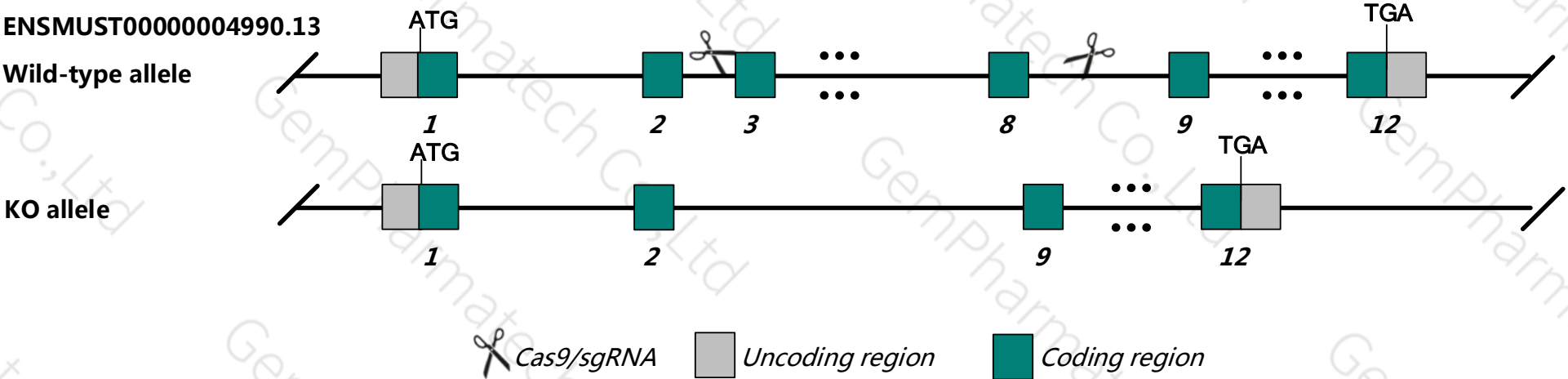
**Cas9-KO**

**Strain background**

**C57BL/6JGpt**

# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Mapk14* gene. The schematic diagram is as follows:



- The *Mapk14* gene has 10 transcripts. According to the structure of *Mapk14* gene, exon3-exon8 of *Mapk14-201* (ENSMUST00000004990.13) transcript is recommended as the knockout region. The region contains 436bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Mapk14* gene. The brief process is as follows: sgRNA was transcribed in vitro. Cas9 and sgRNA 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.

- According to the existing MGI data, Mice homozygous for various null mutations are embryonic to perinatal lethal showing multiple organ system defects. Mice homozygous for a knock-out mutation exhibit abnormal myoblast differentiation and delayed myofiber growth and maturation.
- The *Mapk14* 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.
- The KO region overlaps *Gm4356* gene. Knockout the region may affect the function of *Gm4356* gene. Transcript *Mapk14-207* may not be affected.
- 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)

## Mapk14 mitogen-activated protein kinase 14 [ *Mus musculus* (house mouse) ]

Gene ID: 26416, updated on 15-Aug-2019

### Summary



<b>Official Symbol</b>	Mapk14 provided by <a href="#">MGI</a>
<b>Official Full Name</b>	mitogen-activated protein kinase 14 provided by <a href="#">MGI</a>
<b>Primary source</b>	<a href="#">MGI:MGI:1346865</a>
<b>See related</b>	<a href="#">Ensembl:ENSMUSG00000053436</a>
<b>Gene type</b>	protein coding
<b>RefSeq status</b>	VALIDATED
<b>Organism</b>	<a href="#">Mus musculus</a>
<b>Lineage</b>	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
<b>Also known as</b>	p38; Crk1; Mxi2; p38a; CSBP2; Csbp1; PRKM14; PRKM15; p38MAPK; p38alpha; p38-alpha
<b>Expression</b>	Ubiquitous expression in spleen adult (RPKM 35.2), thymus adult (RPKM 34.4) and 28 other tissues <a href="#">See more</a>
<b>Orthologs</b>	<a href="#">human</a> <a href="#">all</a>

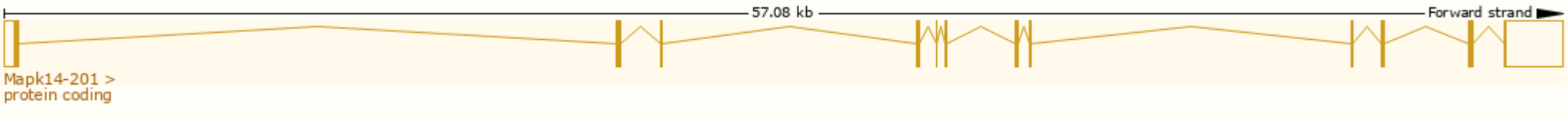


# Transcript information (Ensembl)

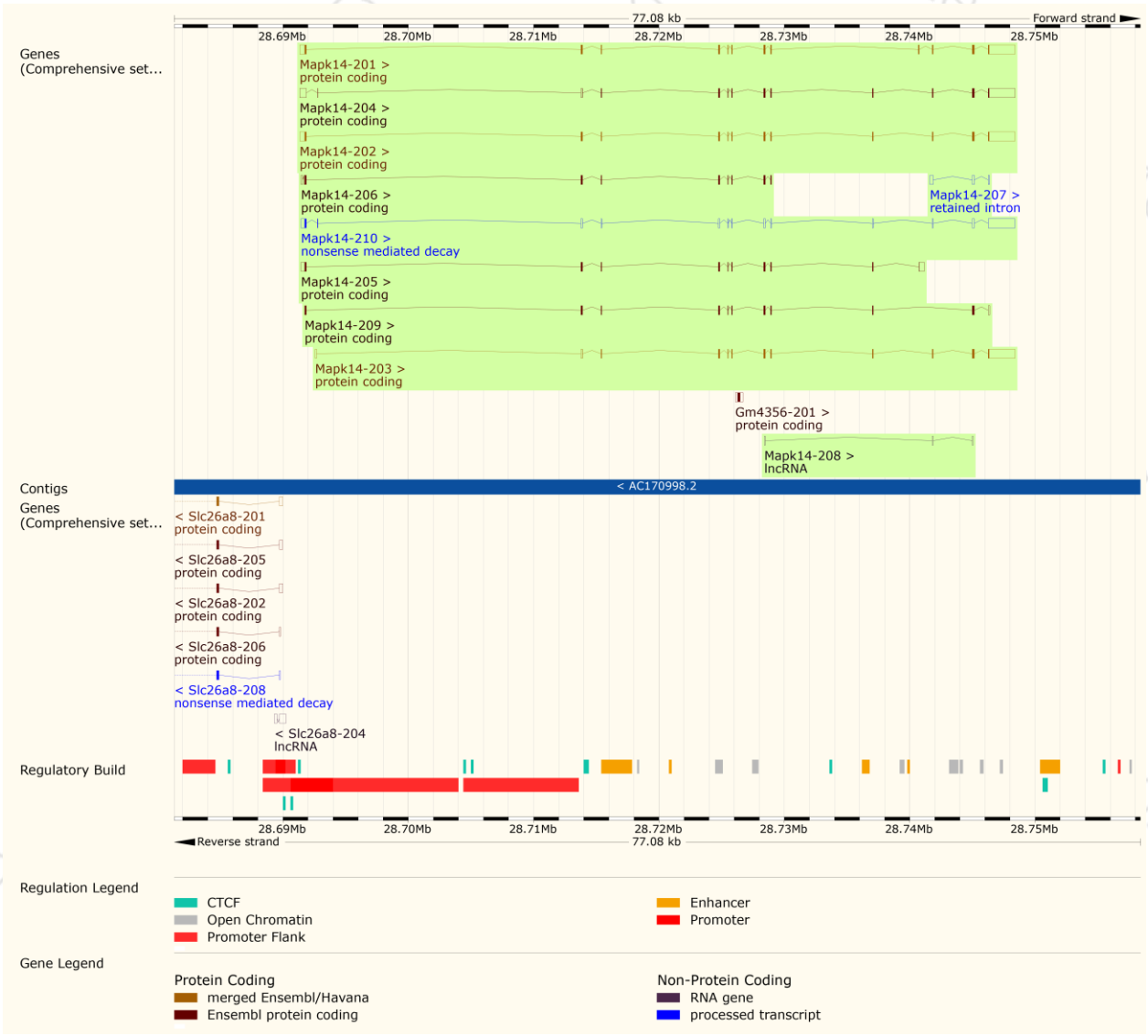
The gene has 10 transcripts, all transcripts are shown below:

Name ▲	Transcript ID ▲	bp ▲	Protein ▲	Translation ID ▲	Biotype ▲	CCDS ▲	UniProt ▲	Flags ▲
Mapk14-201	<a href="#">ENSMUST00000004990.13</a>	3562	<a href="#">360aa</a>	<a href="#">ENSMUSP000000004990.6</a>	Protein coding	<a href="#">CCDS50048</a>	<a href="#">P47811</a> <a href="#">Q5U421</a>	TSL:1 GENCODE basic APPRIS ALT1
Mapk14-202	<a href="#">ENSMUST000000062694.15</a>	3548	<a href="#">360aa</a>	<a href="#">ENSMUSP000000061958.8</a>	Protein coding	<a href="#">CCDS28583</a>	<a href="#">P47811</a>	TSL:1 GENCODE basic APPRIS P3
Mapk14-203	<a href="#">ENSMUST000000114752.2</a>	3148	<a href="#">283aa</a>	<a href="#">ENSMUSP000000110400.1</a>	Protein coding	<a href="#">CCDS50049</a>	<a href="#">P47811</a>	TSL:1 GENCODE basic
Mapk14-204	<a href="#">ENSMUST000000114754.7</a>	3592	<a href="#">283aa</a>	<a href="#">ENSMUSP000000110402.1</a>	Protein coding	<a href="#">CCDS50049</a>	<a href="#">P47811</a>	TSL:1 GENCODE basic
Mapk14-205	<a href="#">ENSMUST000000114758.8</a>	1478	<a href="#">258aa</a>	<a href="#">ENSMUSP000000110406.1</a>	Protein coding	-	<a href="#">B2KF35</a> <a href="#">P47811</a>	TSL:1 GENCODE basic
Mapk14-206	<a href="#">ENSMUST000000124886.8</a>	866	<a href="#">227aa</a>	<a href="#">ENSMUSP000000116914.2</a>	Protein coding	-	<a href="#">B2KF34</a>	CDS 3' incomplete TSL:3
Mapk14-207	<a href="#">ENSMUST000000151613.1</a>	509	No protein	-	Retained intron	-	-	TSL:2
Mapk14-208	<a href="#">ENSMUST000000233095.1</a>	223	No protein	-	lncRNA	-	-	-
Mapk14-209	<a href="#">ENSMUST000000233250.1</a>	1066	<a href="#">307aa</a>	<a href="#">ENSMUSP000000156692.1</a>	Protein coding	-	<a href="#">A0A3B2WB60</a>	GENCODE basic
Mapk14-210	<a href="#">ENSMUST000000233811.1</a>	3490	<a href="#">50aa</a>	<a href="#">ENSMUSP000000156603.1</a>	Nonsense mediated decay	-	<a href="#">A0A3B2WAZ7</a>	-

The strategy is based on the design of *Mapk14-201* transcript, The transcription is shown below

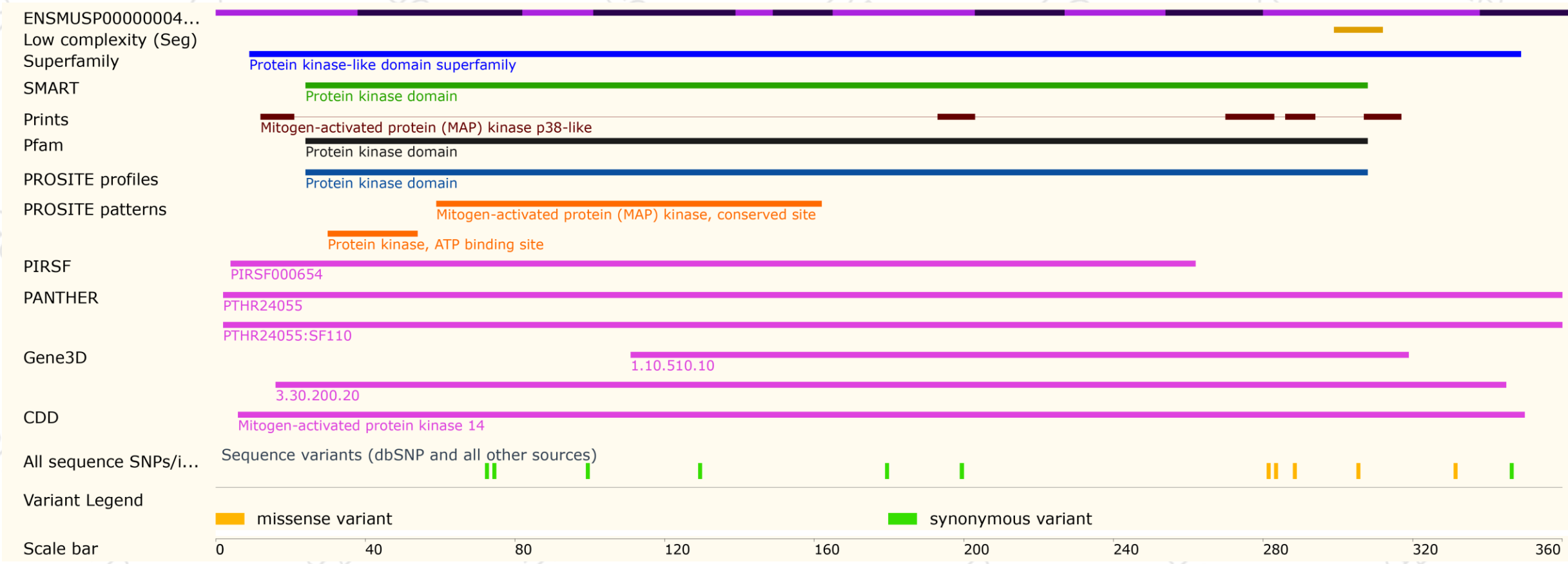


# Genomic location distribution



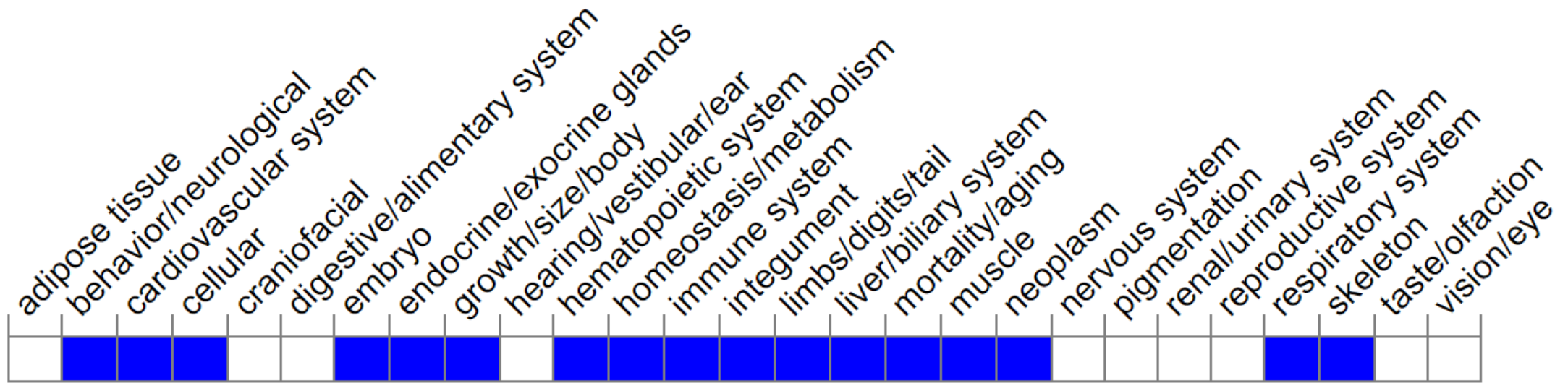


# Protein domain



# Mouse phenotype description(MGI)

## Phenotype Overview ?



*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 various null mutations are embryonic to perinatal lethal showing multiple organ system defects. Mice homozygous for a knock-out mutation exhibit abnormal myoblast differentiation and delayed myofiber growth and maturation.

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

Tel: 025-5864 1534

