

# ***Stat3* Cas9-CKO Strategy**

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Design Date: 2019-8-15

# Project Overview

**Project Name**

*Stat3*

**Project type**

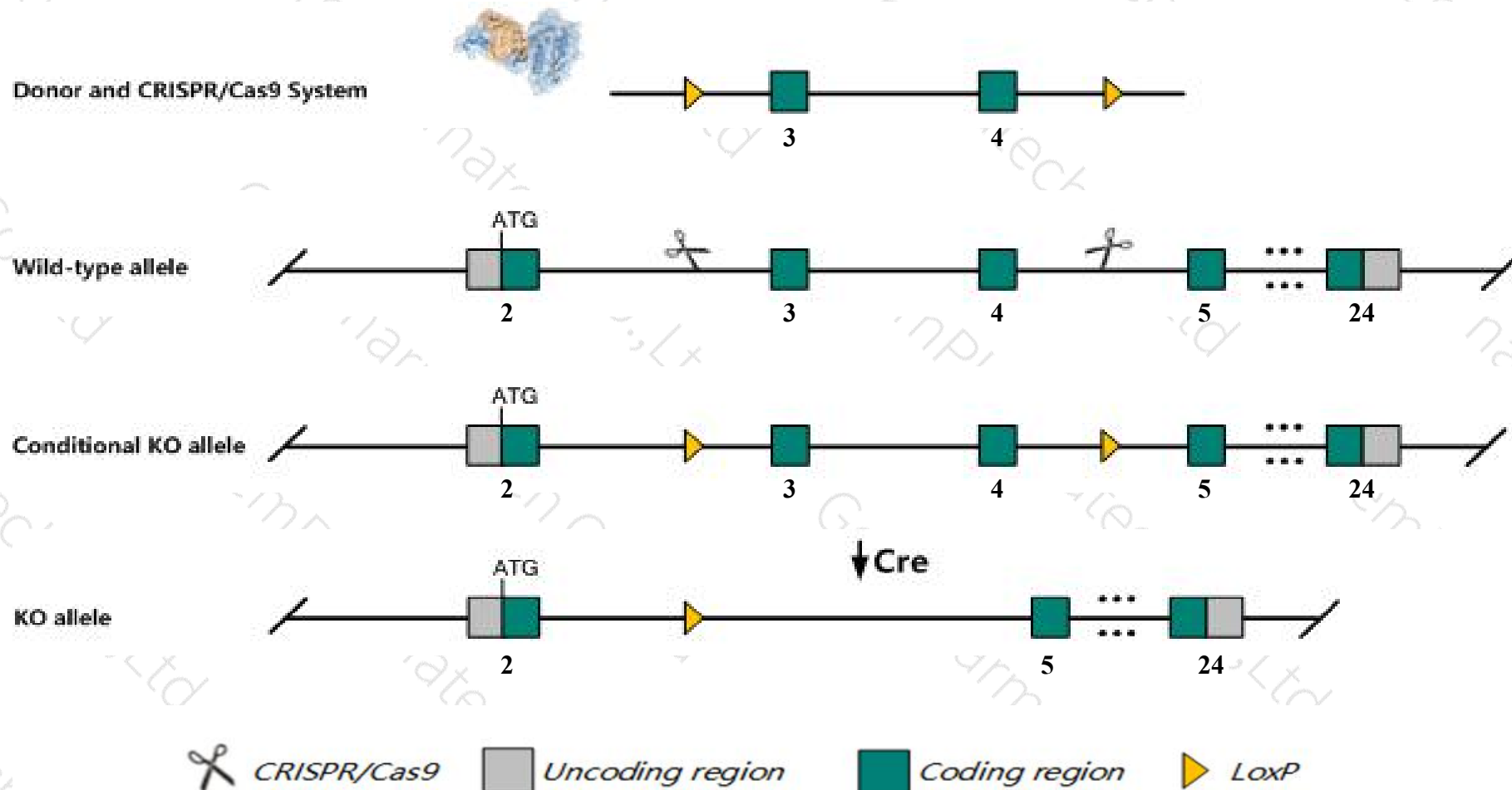
**Cas9-CKO**

**Strain background**

**C57BL/6JGpt**

# Conditional Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Stat3* gene. The schematic diagram is as follows:



- The *Stat3* gene has 14 transcripts. According to the structure of *Stat3* gene, exon3-exon4 of *Stat3-203* (ENSMUST00000127638.7) transcript is recommended as the knockout region. The region contains 244bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Stat3* gene. The brief process is as follows: gRNA was transcribed in vitro, donor was constructed. Cas9, gRNA 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, Homozygotes for targeted null mutations die at embryonic day 6.5-7.5. Conditional, tissue specific mutants are variably viable and show diverse defects including obesity, diabetes, thermal dysregulation and infertility.
- The *Stat3* gene is located on the Chr11. 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)

## Stat3 signal transducer and activator of transcription 3 [Mus musculus (house mouse)]

Gene ID: 20848, updated on 9-Apr-2019

### Summary

**Official Symbol** Stat3 provided by [MGI](#)

**Official Full Name** signal transducer and activator of transcription 3 provided by [MGI](#)

**Primary source** [MGI:MGI:103038](#)

**See related** [Ensembl:ENSMUSG00000004040](#)

**Gene type** protein coding

**RefSeq status** REVIEWED

**Organism** [Mus musculus](#)

**Lineage** Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus

**Also known as** 1110034C02Rik, AW109958, Aprf

**Summary** The protein encoded by this gene is a member of the STAT protein family. In response to cytokines and growth factors, STAT family members are phosphorylated by the receptor associated kinases, and then form homo- or heterodimers that translocate to the cell nucleus where they act as transcription activators. This protein is activated through phosphorylation in response to various cytokines and growth factors including IFNs, EGF, IL5, IL6, HGF, LIF and BMP2. This protein mediates the expression of a variety of genes in response to cell stimuli, and thus plays a key role in many cellular processes such as cell growth and apoptosis. The small GTPase Rac1 has been shown to bind and regulate the activity of this protein. PIAS3 protein is a specific inhibitor of this protein. Alternative splicing results in multiple transcript variants encoding distinct isoforms. [provided by RefSeq, Sep 2015]

**Expression** Ubiquitous expression in lung adult (RPKM 48.1), mammary gland adult (RPKM 46.4) and 28 other tissues [See more](#)

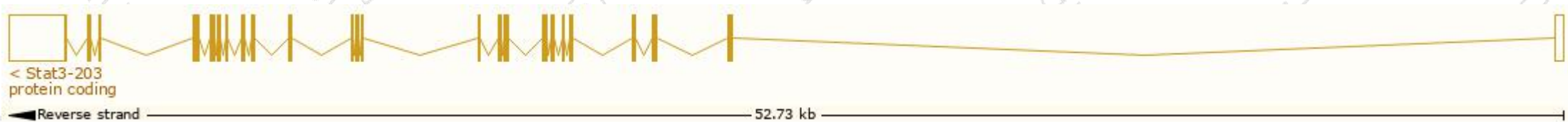
**Orthologs** [human](#) [all](#)

# Transcript information (Ensembl)

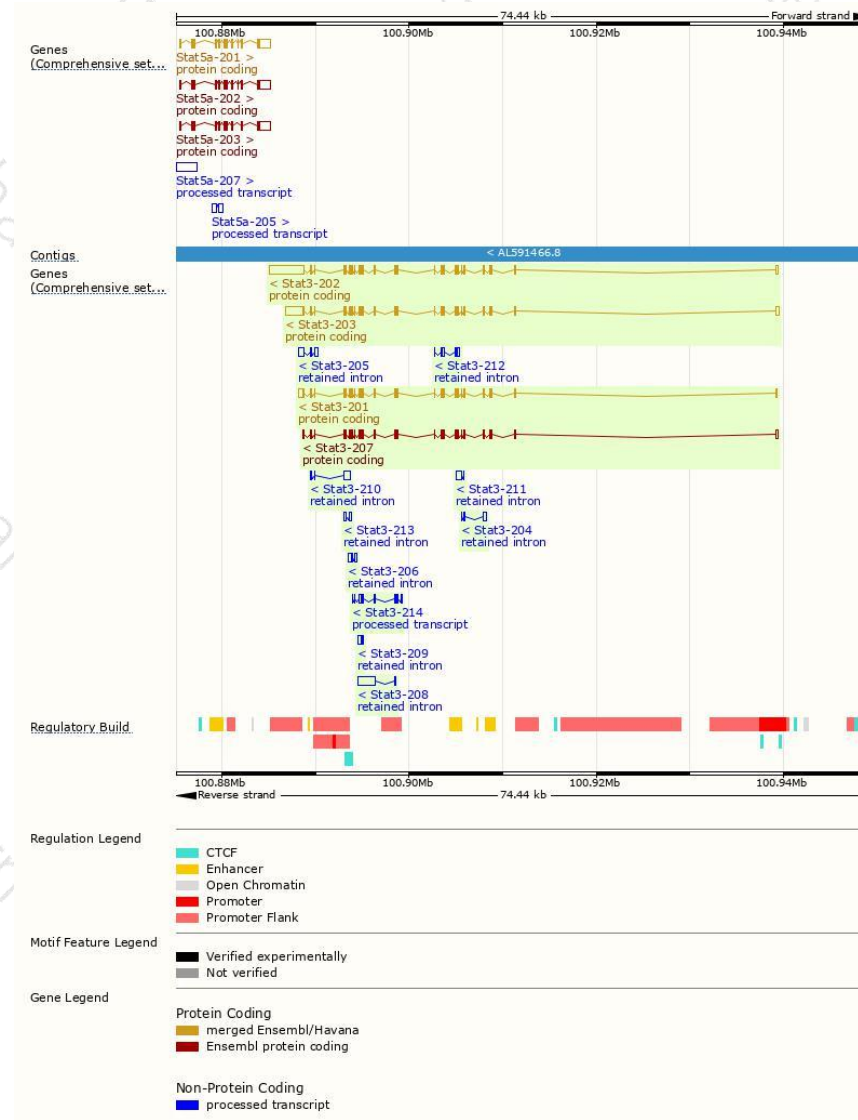
The gene has 14 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Stat3-202	<a href="#">ENSMUST00000103114.7</a>	6042	<a href="#">722aa</a>	Protein coding	<a href="#">CCDS25441</a>	<a href="#">P42227_Q6GU23</a>	TSL:1 GENCODE basic APPRIS ALT1
Stat3-203	<a href="#">ENSMUST00000127638.7</a>	4516	<a href="#">770aa</a>	Protein coding	<a href="#">CCDS25440</a>	<a href="#">P42227</a>	TSL:1 GENCODE basic APPRIS P4
Stat3-201	<a href="#">ENSMUST00000092671.11</a>	2927	<a href="#">769aa</a>	Protein coding	<a href="#">CCDS48934</a>	<a href="#">P42227</a>	TSL:1 GENCODE basic APPRIS ALT1
Stat3-207	<a href="#">ENSMUST00000138438.1</a>	2506	<a href="#">744aa</a>	Protein coding	-	<a href="#">B7ZC18</a>	TSL:5 GENCODE basic
Stat3-214	<a href="#">ENSMUST00000156645.1</a>	651	No protein	Processed transcript	-	-	TSL:5
Stat3-208	<a href="#">ENSMUST00000146971.1</a>	1995	No protein	Retained intron	-	-	TSL:1
Stat3-205	<a href="#">ENSMUST00000137093.7</a>	1079	No protein	Retained intron	-	-	TSL:2
Stat3-210	<a href="#">ENSMUST00000152601.1</a>	826	No protein	Retained intron	-	-	TSL:1
Stat3-212	<a href="#">ENSMUST00000154170.1</a>	672	No protein	Retained intron	-	-	TSL:3
Stat3-206	<a href="#">ENSMUST00000137367.1</a>	646	No protein	Retained intron	-	-	TSL:2
Stat3-211	<a href="#">ENSMUST00000152623.1</a>	609	No protein	Retained intron	-	-	TSL:3
Stat3-209	<a href="#">ENSMUST00000151544.1</a>	408	No protein	Retained intron	-	-	TSL:3
Stat3-213	<a href="#">ENSMUST00000155972.1</a>	396	No protein	Retained intron	-	-	TSL:3
Stat3-204	<a href="#">ENSMUST00000130686.1</a>	349	No protein	Retained intron	-	-	TSL:3

The strategy is based on the design of *Stat3-203* transcript,The transcription is shown below

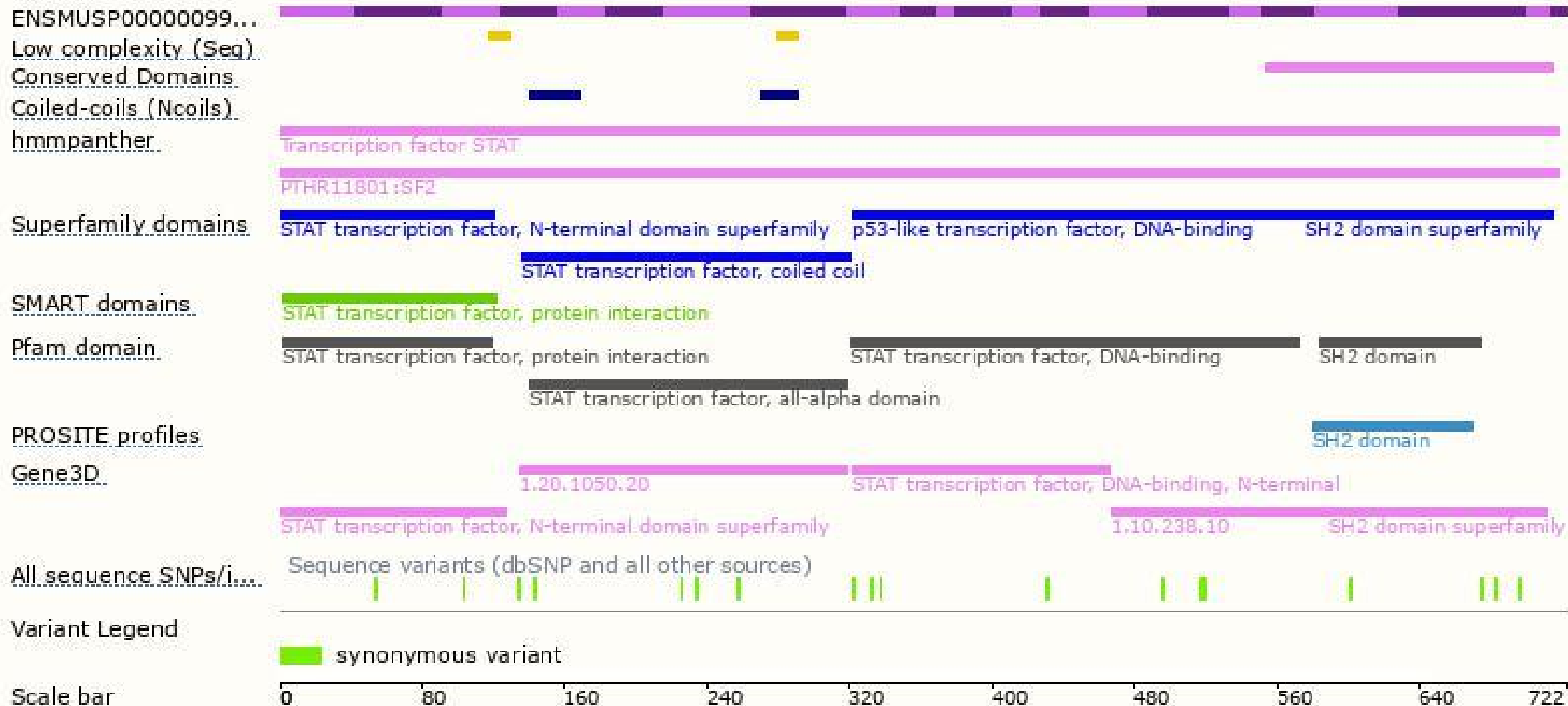


# Genomic location distribution

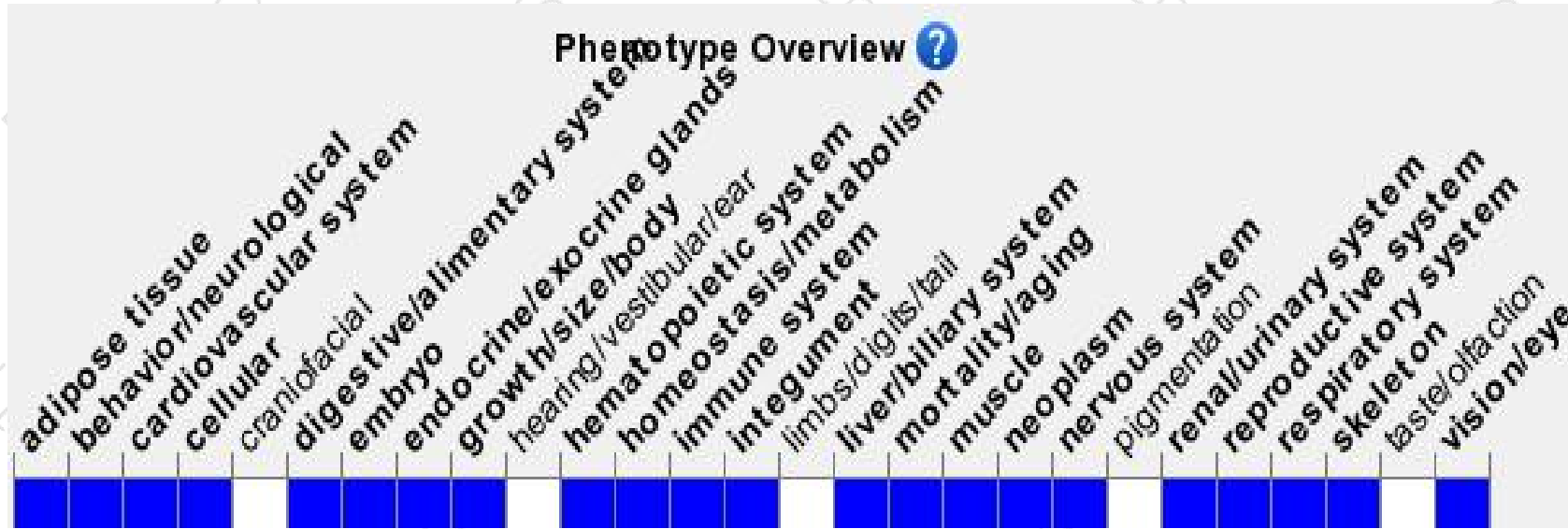




# Protein domain



# 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, Homozygotes for targeted null mutations die at embryonic day 6.5-7.5. Conditional, tissue specific mutants are variably viable and show diverse defects including obesity, diabetes, thermal dysregulation and infertility.

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

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