

# *Ccn3* Cas9-KO Strategy

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**Reviewer:**

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# Project Overview

**Project Name**

***Ccn3***

**Project type**

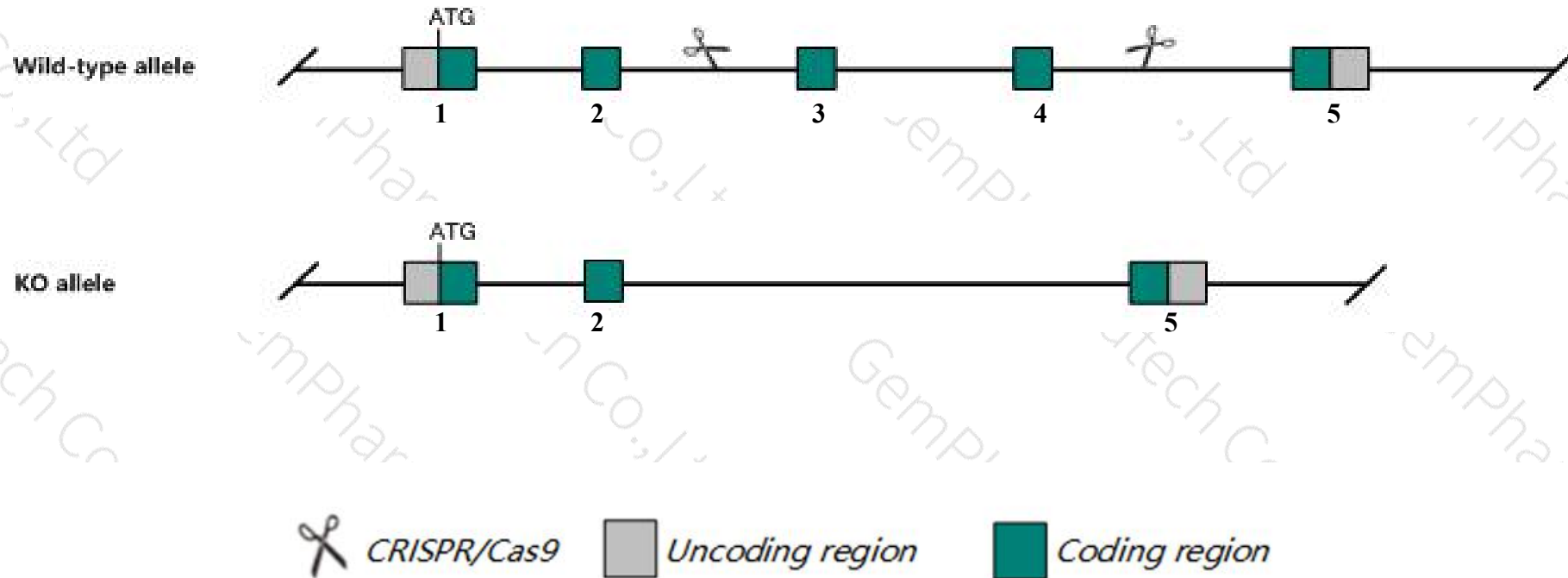
**Cas9-KO**

**Strain background**

**C57BL/6JGpt**

# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Ccn3* gene. The schematic diagram is as follows:



- The *Ccn3* gene has 1 transcript. According to the structure of *Ccn3* gene, exon3-exon4 of *Ccn3-201* (ENSMUST00000050027.8) transcript is recommended as the knockout region. The region contains 476bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Ccn3* gene. The brief process is as follows: CRISPR/Cas9 system v

- According to the existing MGI data, Heterozygotes and homozygotes for a null mutation exhibit abnormal skeletal and cardiac development, muscle atrophy and cataracts. Mice homozygous for another knock-out allele exhibit minor bone structure and physiology defects.
- The *Ccn3* 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)

## Ccn3 cellular communication network factor 3 [Mus musculus (house mouse)]

Gene ID: 18133, updated on 5-Mar-2019

### Summary



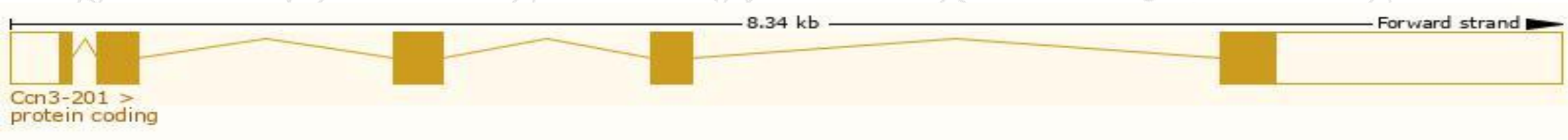
<b>Official Symbol</b>	Ccn3 provided by <a href="#">MGI</a>
<b>Official Full Name</b>	cellular communication network factor 3 provided by <a href="#">MGI</a>
<b>Primary source</b>	<a href="#">MGI:MGI:109185</a>
<b>See related</b>	<a href="#">Ensembl:ENSMUSG000000037362</a>
<b>Gene type</b>	protein coding
<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>	C130088N23Rik, Nov
<b>Expression</b>	Biased expression in colon adult (RPKM 83.7), frontal lobe adult (RPKM 20.9) and 4 other tissues <a href="#">See more</a>
<b>Orthologs</b>	<a href="#">human</a> <a href="#">all</a>

# Transcript information (Ensembl)

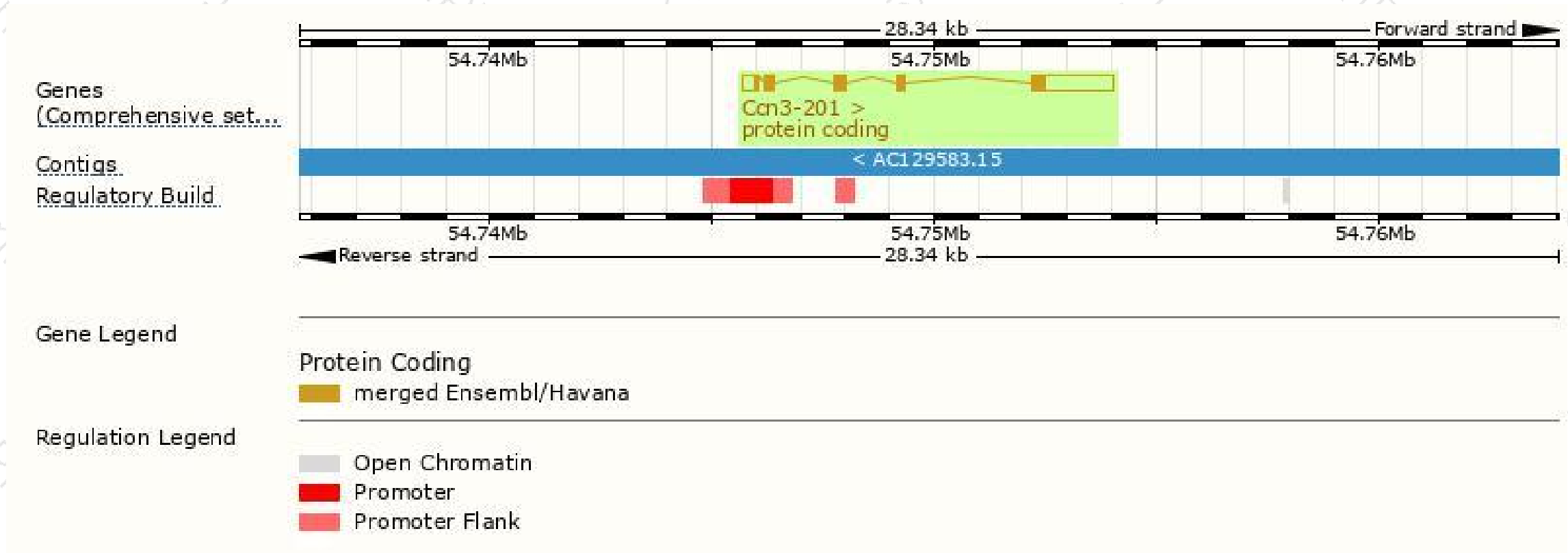
The gene has 1 transcript, and the transcript is shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Ccn3-201	<a href="#">ENSMUST00000050027.8</a>	2869	<a href="#">354aa</a>	Protein coding	<a href="#">CCDS27471</a>	<a href="#">Q64299</a>	TSL:1 GENCODE basic APPRIS P1

The strategy is based on the design of *Ccn3-201* transcript, The transcription is shown below

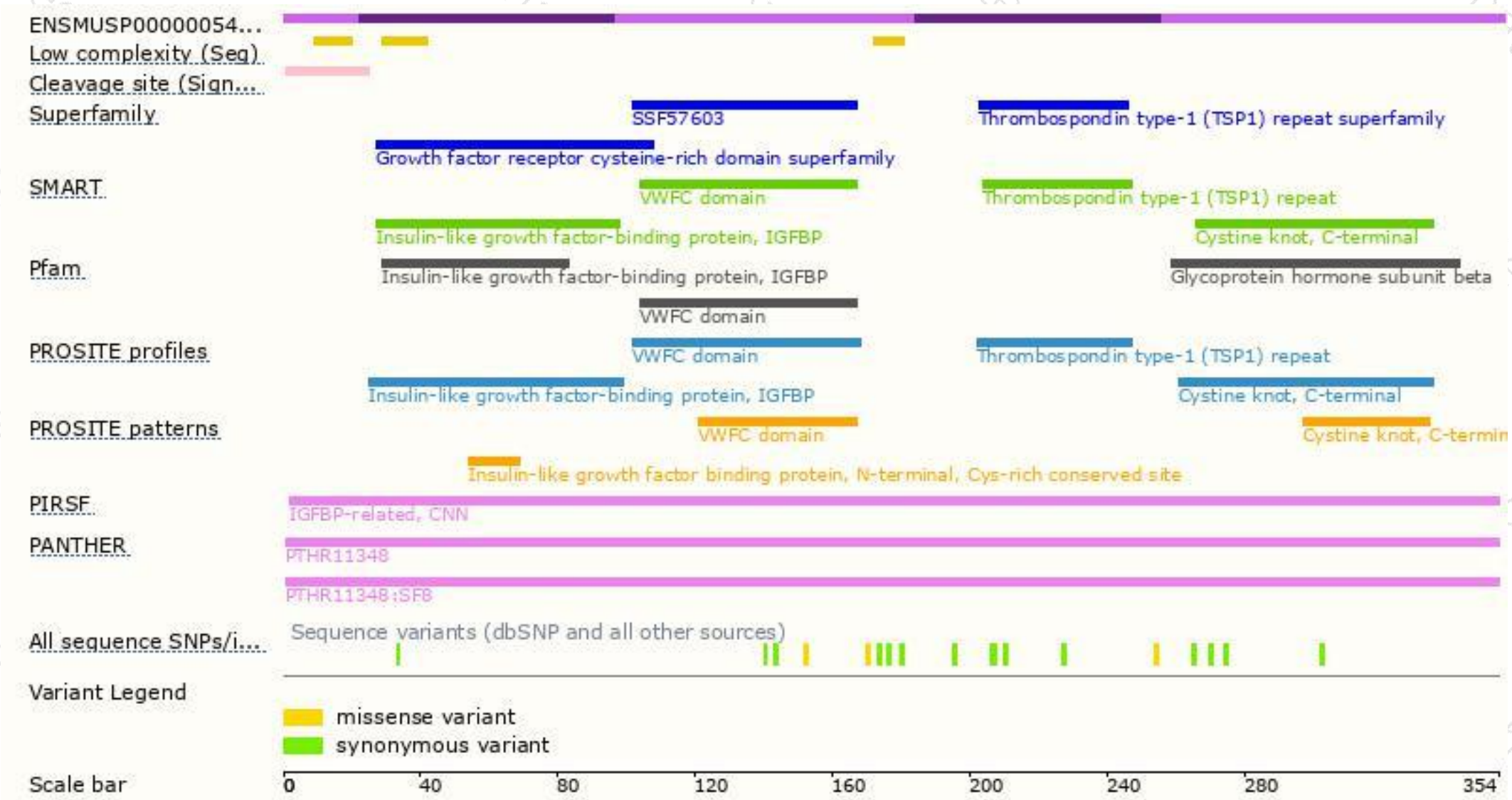


# Genomic location distribution

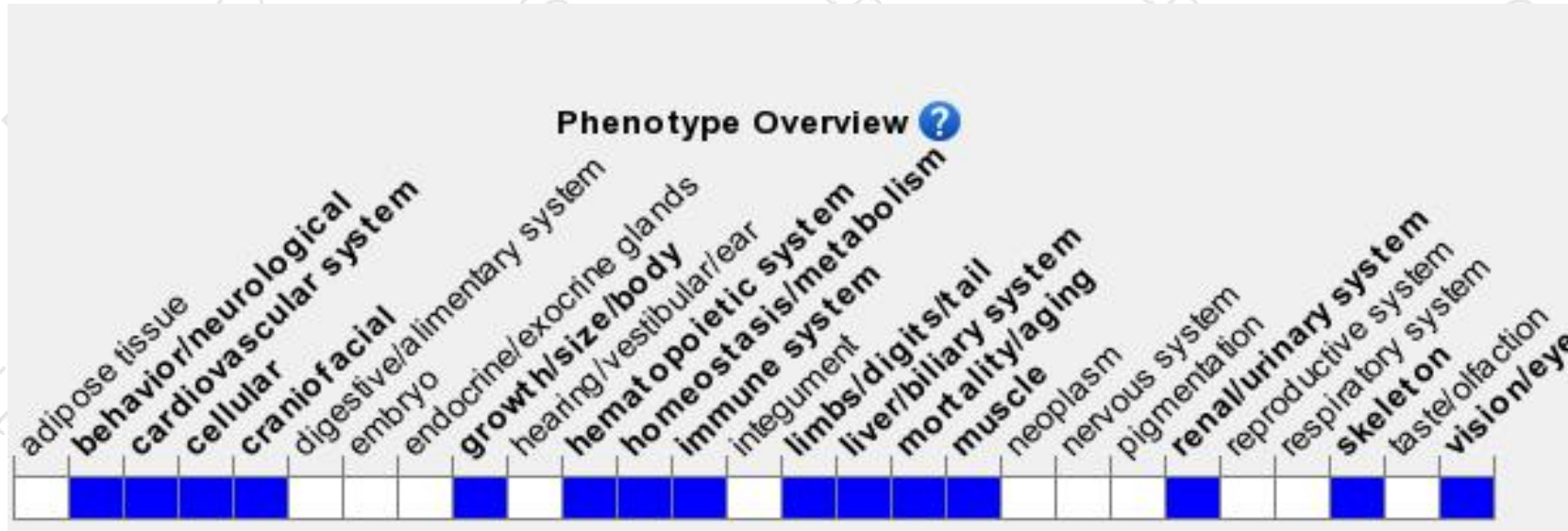




# Protein domain



# 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, Heterozygotes and homozygotes for a null mutation exhibit abnormal skeletal and cardiac development, muscle atrophy and cataracts. Mice homozygous for another knock-out allele exhibit minor bone structure and physiology defects.

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

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