



# **Postn Cas9-CKO Strategy**

**Designer:**

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**Design Date:**

**2019-7-23**

# Project Overview

**Project Name**

*Postn*

**Project type**

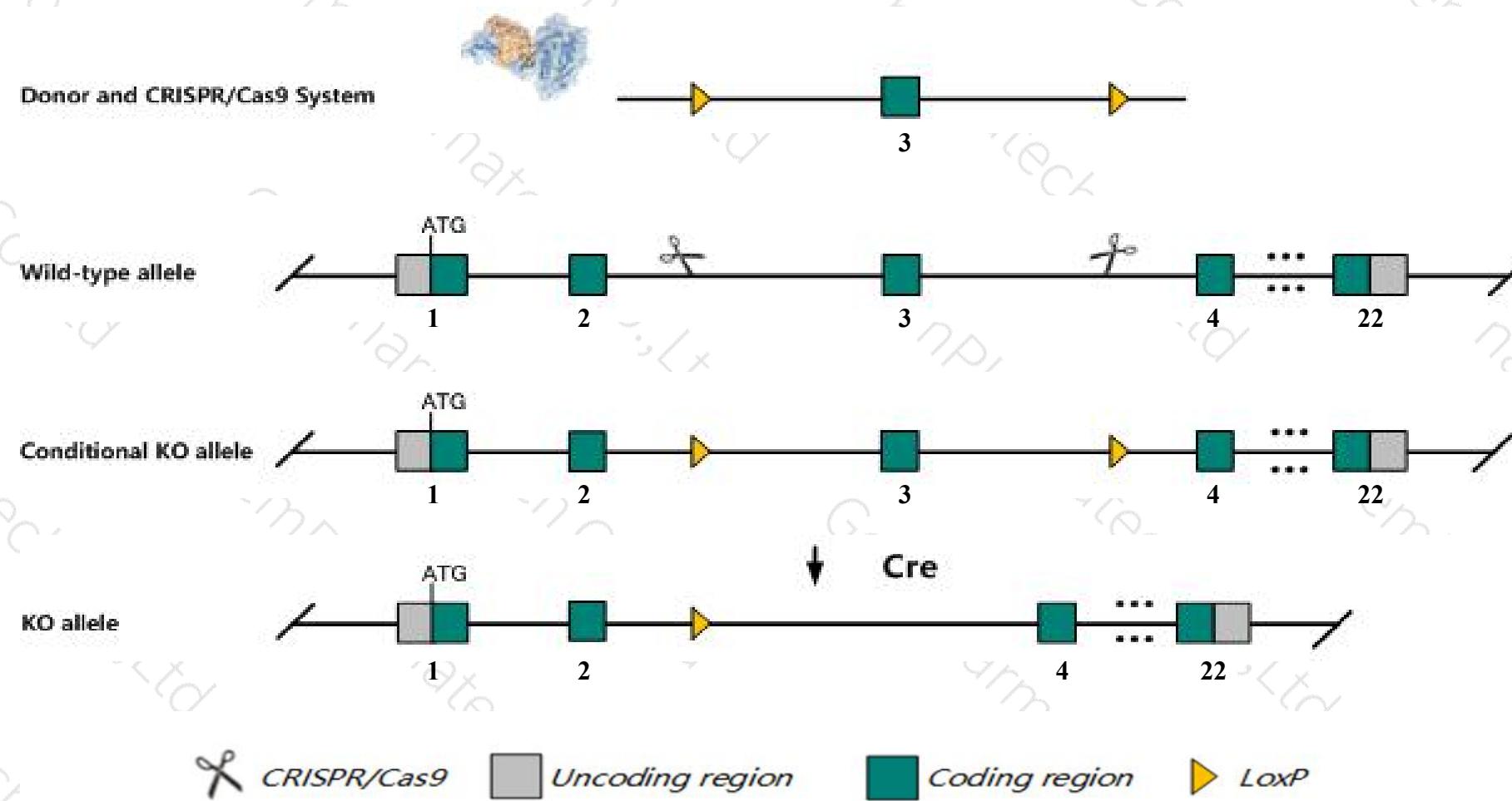
**Cas9-CKO**

**Strain background**

**C57BL/6JGpt**

# Conditional Knockout strategy

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



# Technical routes

- The *Postn* gene has 9 transcripts. According to the structure of *Postn* gene, exon3 of *Postn-203* (ENSMUST00000107985.9) transcript is recommended as the knockout region. The region contains 65bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Postn* 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.



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# Notice

- According to the existing MGI data, Homozygous null mice display abnormalities of the enamel, periodontal ligament, ameloblasts, and incisors. For one allele changing the hardness of the food alters the severity of the abnormalities.
- Transcript *Postn-206* may not be affected.
- The *Postn* gene is located on the Chr3. 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)

## Postn periostin, osteoblast specific factor [Mus musculus (house mouse)]

Gene ID: 50706, updated on 16-Feb-2019

### Summary



<b>Official Symbol</b>	Postn provided by <a href="#">MGI</a>
<b>Official Full Name</b>	periostin, osteoblast specific factor provided by <a href="#">MGI</a>
<b>Primary source</b>	<a href="#">MGI:MGI:1926321</a>
<b>See related</b>	<a href="#">Ensembl:ENSMUSG00000027750</a>
<b>Gene type</b>	protein coding
<b>RefSeq status</b>	REVIEWED
<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>	A630052E07Rik, AI747096, OSF-2, Osf2, PLF, PN
<b>Summary</b>	This gene encodes a secreted extracellular matrix protein that functions in tissue development and regeneration, including wound healing and ventricular remodeling following myocardial infarction. The encoded protein binds to integrins to support adhesion and migration of epithelial cells. This protein plays a role in cancer stem cell maintenance and metastasis. Mice lacking this gene exhibit cardiac valve disease, and skeletal and dental defects. Alternative splicing results in multiple transcript variants encoding different isoforms. [provided by RefSeq, Sep 2015]
<b>Expression</b>	Biased expression in limb E14.5 (RPKM 151.8), placenta adult (RPKM 112.7) and 12 other tissues <a href="#">See more</a>
<b>Orthologs</b>	<a href="#">human</a> <a href="#">all</a>

# Transcript information (Ensembl)

The gene has 9 transcripts, all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Postn-203	<a href="#">ENSMUST00000107985.9</a>	3205	<a href="#">810aa</a>	Protein coding	<a href="#">CCDS57211</a>	<a href="#">Q62009</a>	TSL:1 GENCODE basic APPRIS ALT2
Postn-201	<a href="#">ENSMUST00000073012.12</a>	3189	<a href="#">811aa</a>	Protein coding	<a href="#">CCDS17351</a>	<a href="#">Q62009</a>	TSL:1 GENCODE basic APPRIS P3
Postn-204	<a href="#">ENSMUST00000117373.7</a>	3121	<a href="#">783aa</a>	Protein coding	<a href="#">CCDS57212</a>	<a href="#">Q62009</a>	TSL:1 GENCODE basic APPRIS ALT2
Postn-202	<a href="#">ENSMUST00000081564.12</a>	2670	<a href="#">838aa</a>	Protein coding	-	<a href="#">Q62009</a>	TSL:5 GENCODE basic APPRIS ALT2
Postn-206	<a href="#">ENSMUST00000143258.1</a>	655	<a href="#">102aa</a>	Protein coding	-	<a href="#">F7C9H0</a>	CDS 5' incomplete TSL:2
Postn-207	<a href="#">ENSMUST00000145036.2</a>	5223	No protein	Retained intron	-	-	TSL:1
Postn-205	<a href="#">ENSMUST00000127452.1</a>	900	No protein	Retained intron	-	-	TSL:2
Postn-209	<a href="#">ENSMUST00000154157.7</a>	891	No protein	Retained intron	-	-	TSL:3
Postn-208	<a href="#">ENSMUST00000150868.1</a>	624	No protein	Retained intron	-	-	TSL:2

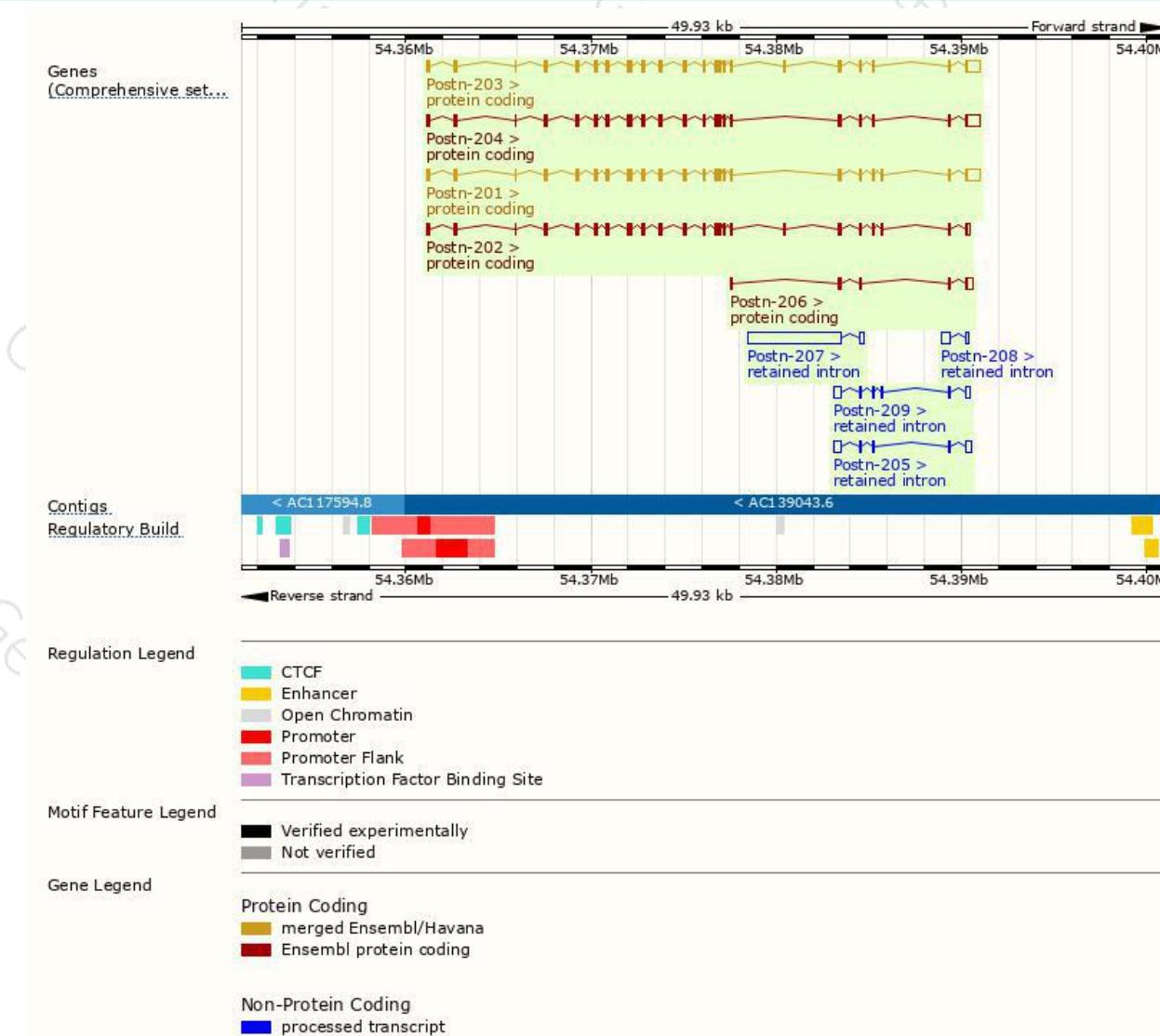
The strategy is based on the design of Postn-203 transcript, The transcription is shown below



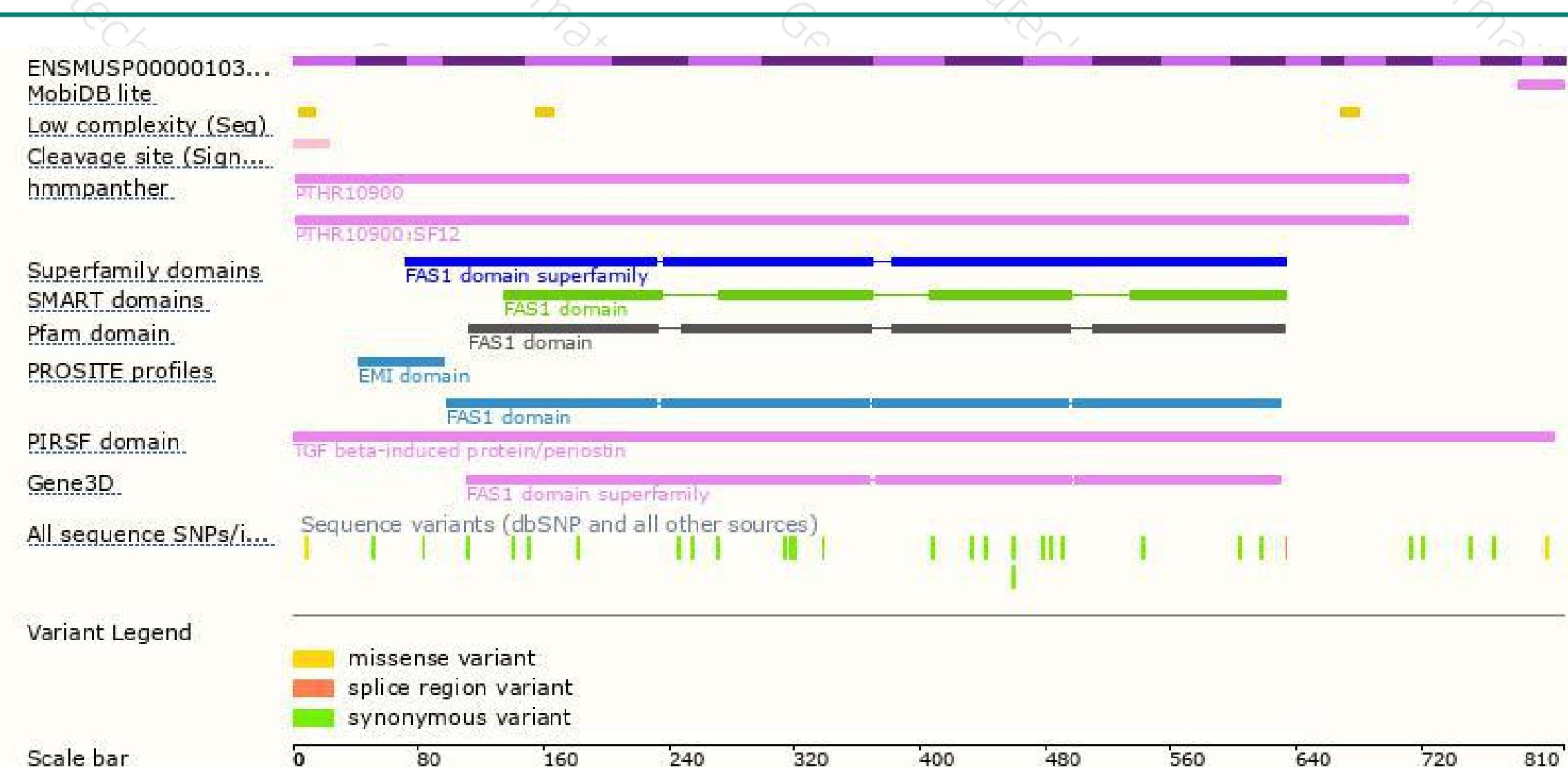


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# Genomic location distribution



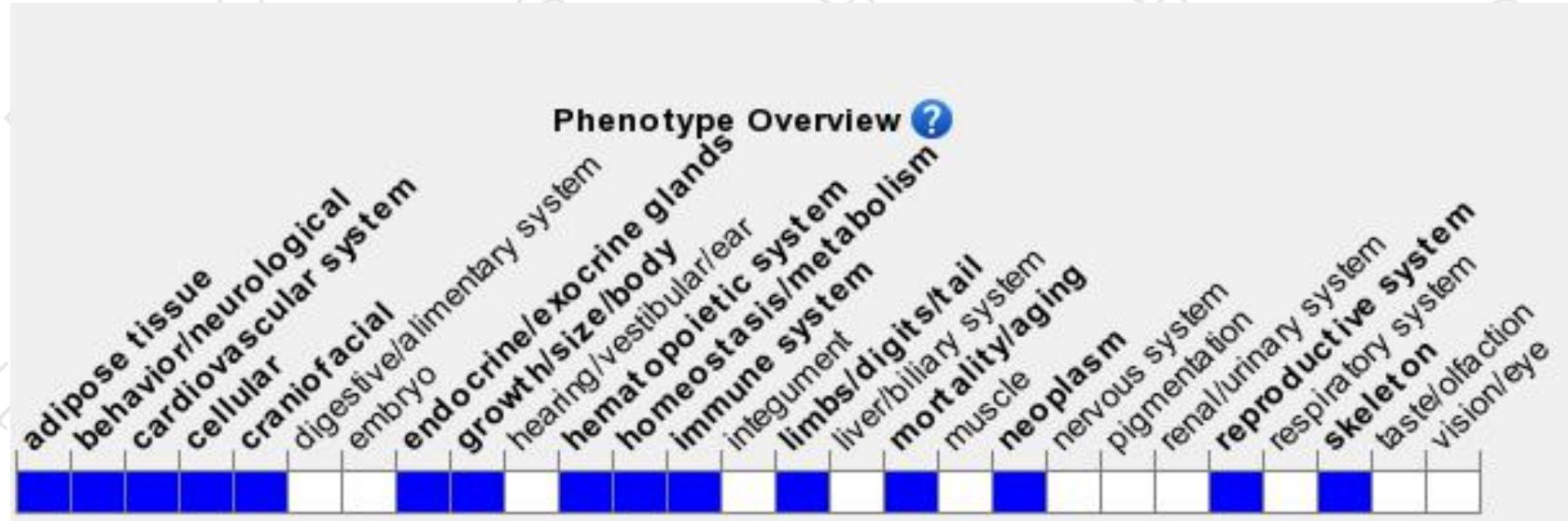
# Protein domain





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# 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, Homozygous null mice display abnormalities of the enamel, periodontal ligament, ameloblasts, and incisors. For one allele changing the hardness of the food alters the severity of the abnormalities.



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

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