

Efnb1 Cas9-KO Strategy

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Project Overview

Project Name

Efnb1

Project type

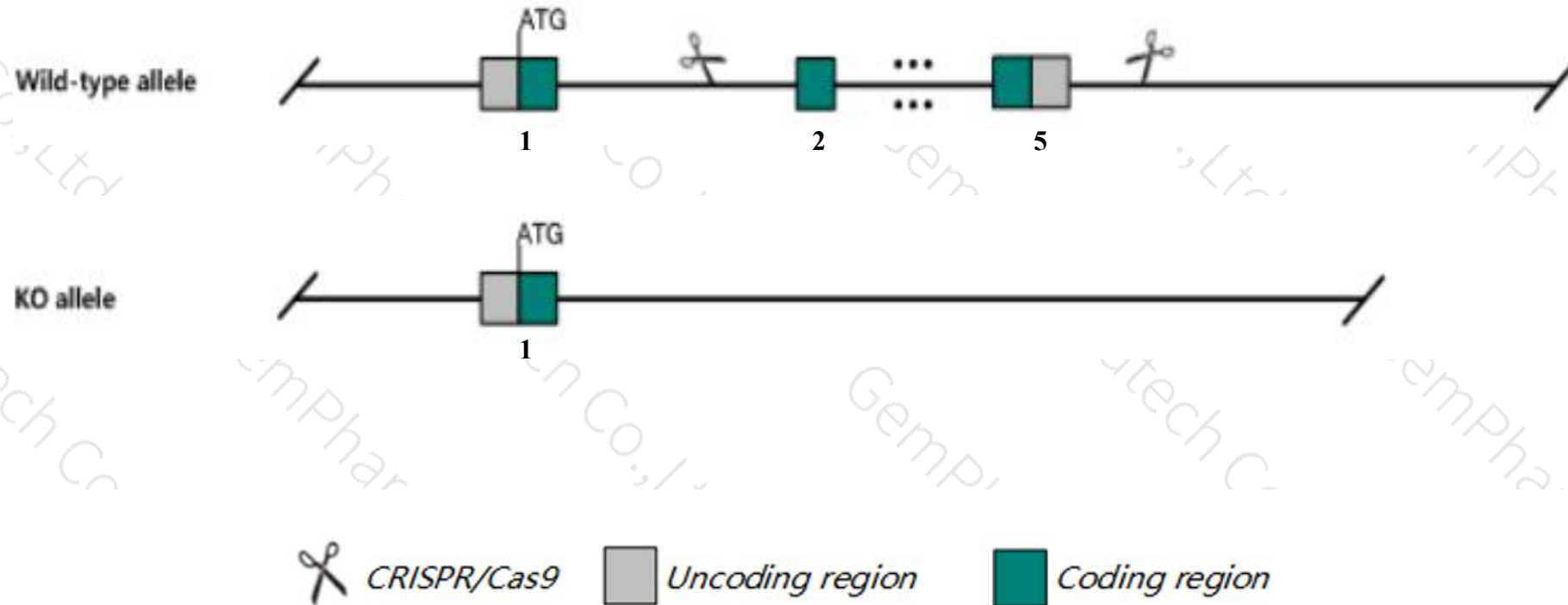
Cas9-KO

Strain background

C57BL/6JGpt

Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Efnb1* gene. The schematic diagram is as follows:



- The *Efnb1* gene has 2 transcripts. According to the structure of *Efnb1* gene, exon2-exon5 of *Efnb1-201* (ENSMUST00000052839.6) transcript is recommended as the knockout region. The region contains most of the coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Efnb1* gene. The brief process is as follows: CRISPR/Cas9 system

- According to the existing MGI data, engineered alleles of this gene produce craniofacial defects resulting in neonatal lethality.
- The *Efnb1* gene is located on the ChrX. 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)

Efnb1 ephrin B1 [Mus musculus (house mouse)]

Gene ID: 13641, updated on 13-Mar-2020

Summary

Official Symbol Efnb1 provided by [MGI](#)

Official Full Name ephrin B1 provided by [MGI](#)

Primary source [MGI:MGI:102708](#)

See related [Ensembl:ENSMUSG00000031217](#)

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 Cek5-L, EFL-3, Elk-L, Epl2, Eplg2, LERK-2, Lerk2, Stra1

Summary This gene encodes a membrane protein that acts as a ligand for Eph family receptors. Signalling occurs bidirectionally in both the cell containing the receptor and the cell expressing this protein. Activity of this protein is important in neuronal axon growth and other developmental processes. [provided by RefSeq, May 2015]

Expression Ubiquitous expression in lung adult (RPKM 47.3), colon adult (RPKM 44.5) and 26 other tissues [See more](#)

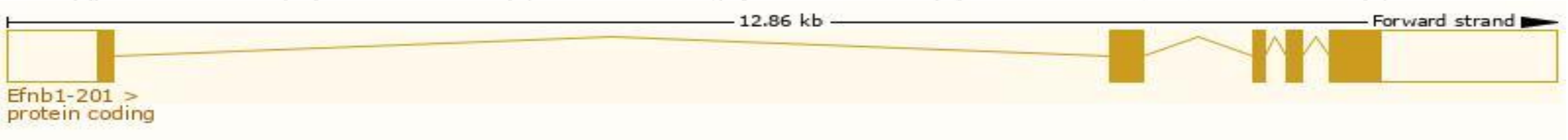
Orthologs [human](#) [all](#)

Transcript information (Ensembl)

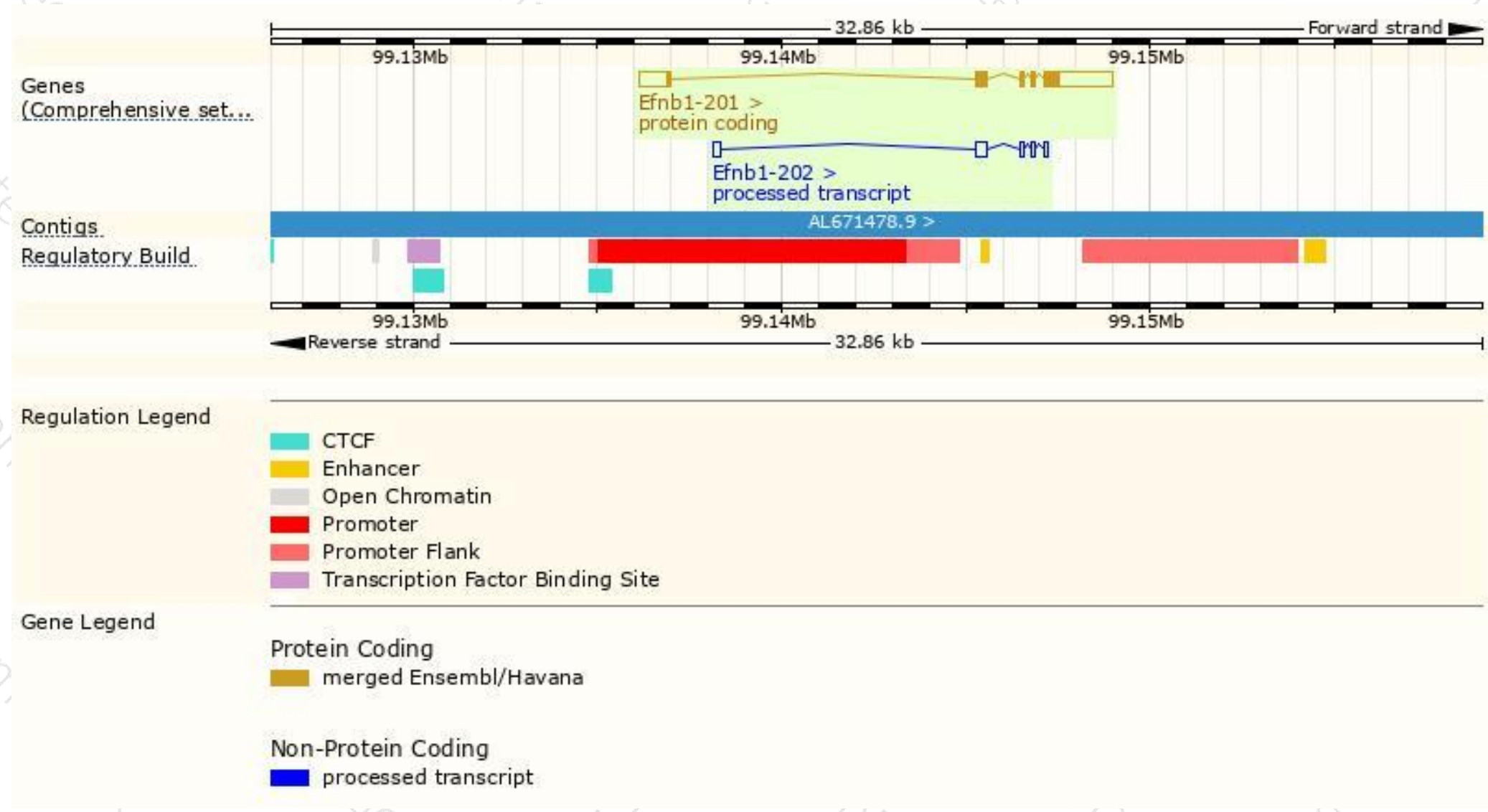
The gene has 2 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Efnb1-201	ENSMUST00000052839.6	3258	345aa	Protein coding	CCDS30298	P52795 Q544L9	TSL:1 GENCODE basic APPRIS is a system to annotate alternatively spliced transcripts based on a range of computational methods to identify the most functionally important transcript(s) of a gene. APPRIS P1
Efnb1-202	ENSMUST000000151750.1	807	No protein	Processed transcript	-	-	TSL:3

The strategy is based on the design of *Efnb1-201* transcript,the transcription is shown below:



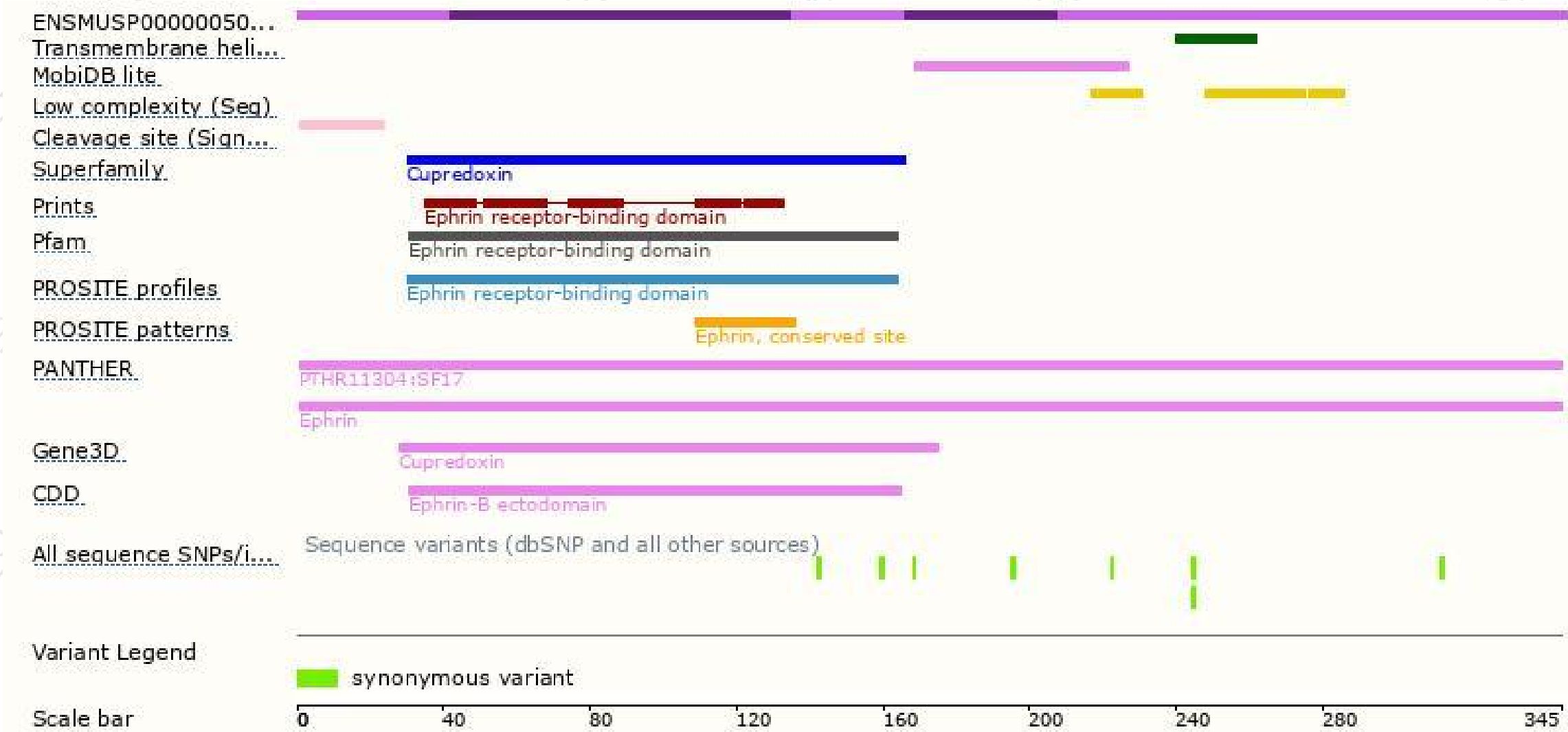
Genomic location distribution



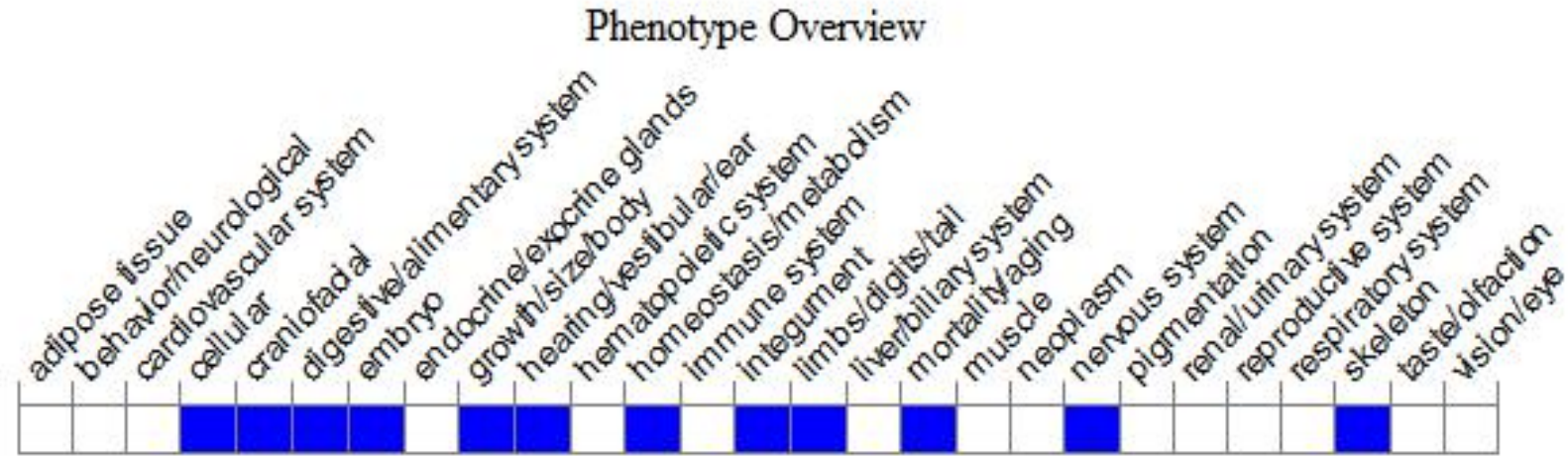
Protein domain



集萃药康
GemPharmatech



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, engineered alleles of this gene produce craniofacial defects resulting in neonatal lethality.

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

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