

Sox21 Cas9-CKO Strategy

Designer: Zihe Cui

Reviewer: Daohua Xu

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Project Overview

Project Name

Sox21

Project type

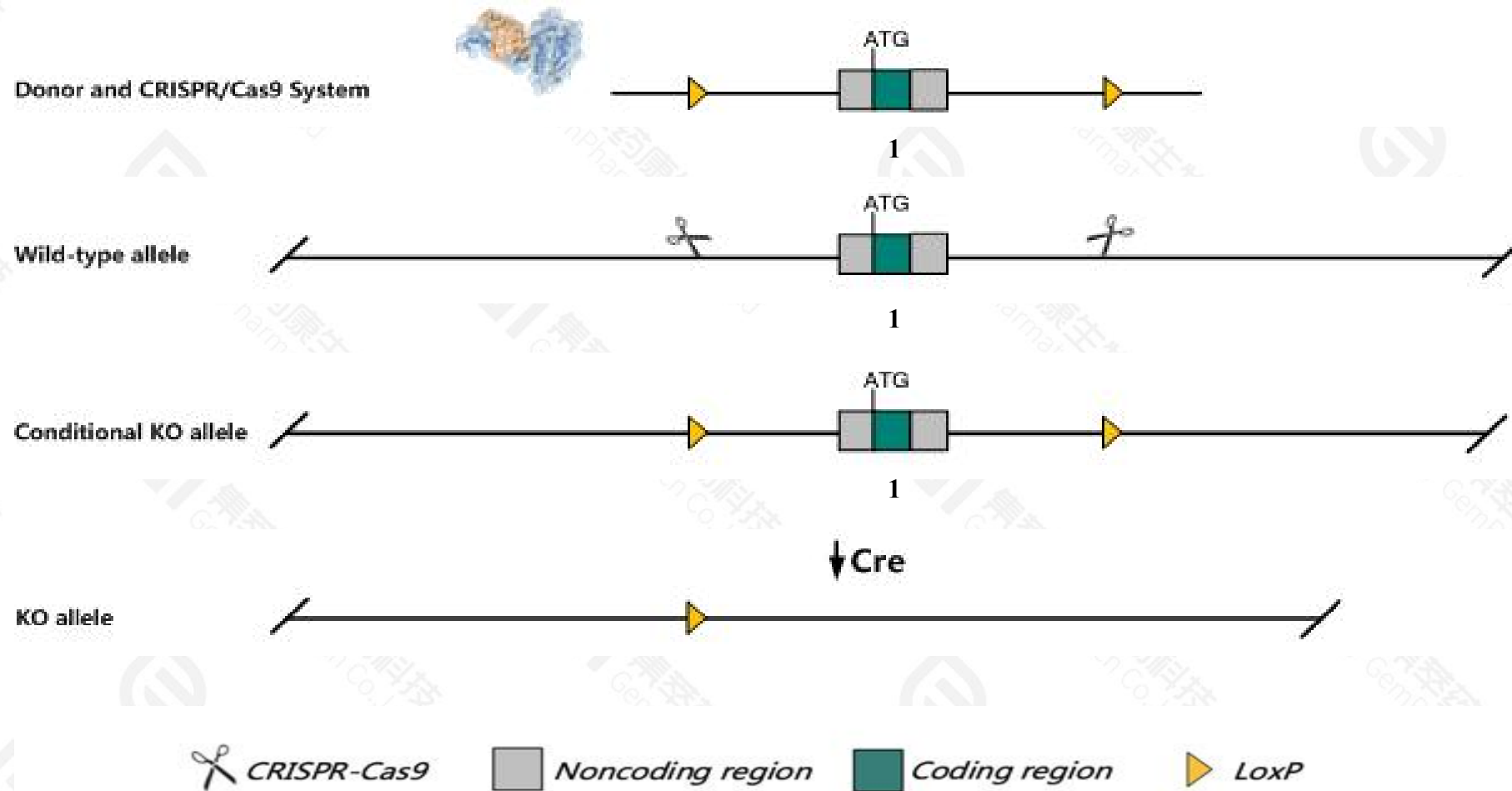
Cas9-CKO

Strain background

C57BL/6JGpt

Conditional Knockout strategy

This model will use CRISPR-Cas9 technology to edit the *Sox21* gene. The schematic diagram is as follows:



Technical routes

- The *Sox21* gene has 1 transcript. According to the structure of *Sox21* gene, exon1 of *Sox21-201*(ENSMUST00000170662.2) transcript is recommended as the knockout region. The region contains all 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 *Sox21* 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.

- According to the existing MGI data, mice homozygous for a null mutation display cyclic alopecia, epidermal hyperplasia, enlarged sebaceous glands, and hair shaft and cuticle abnormalities.
- The KO region overlaps with *Gm4675* gene. Knockout the region may affect the function of *Gm4675* gene.
- The *Sox21* gene is located on the Chr14. 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)

Sox21 SRY (sex determining region Y)-box 21 [Mus musculus (house mouse)]

Gene ID: 223227, updated on 24-Apr-2022

Summary



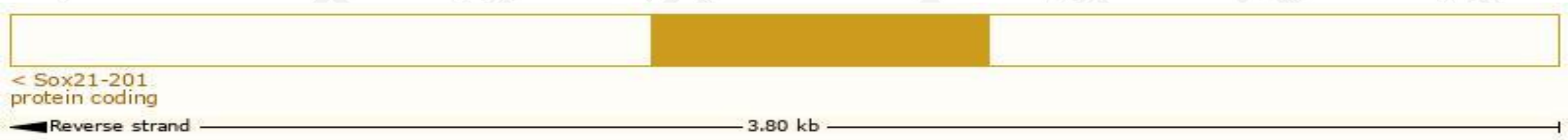
Official Symbol	Sox21 provided by MGI
Official Full Name	SRY (sex determining region Y)-box 21 provided by MGI
Primary source	MGI:MGI:2654070
See related	Ensembl:ENSMUSG00000061517
Gene type	protein coding
RefSeq status	VALIDATED
Organism	Mus musculus
Lineage	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
Also known as	Sox25
Orthologs	human all

Transcript information (Ensembl)

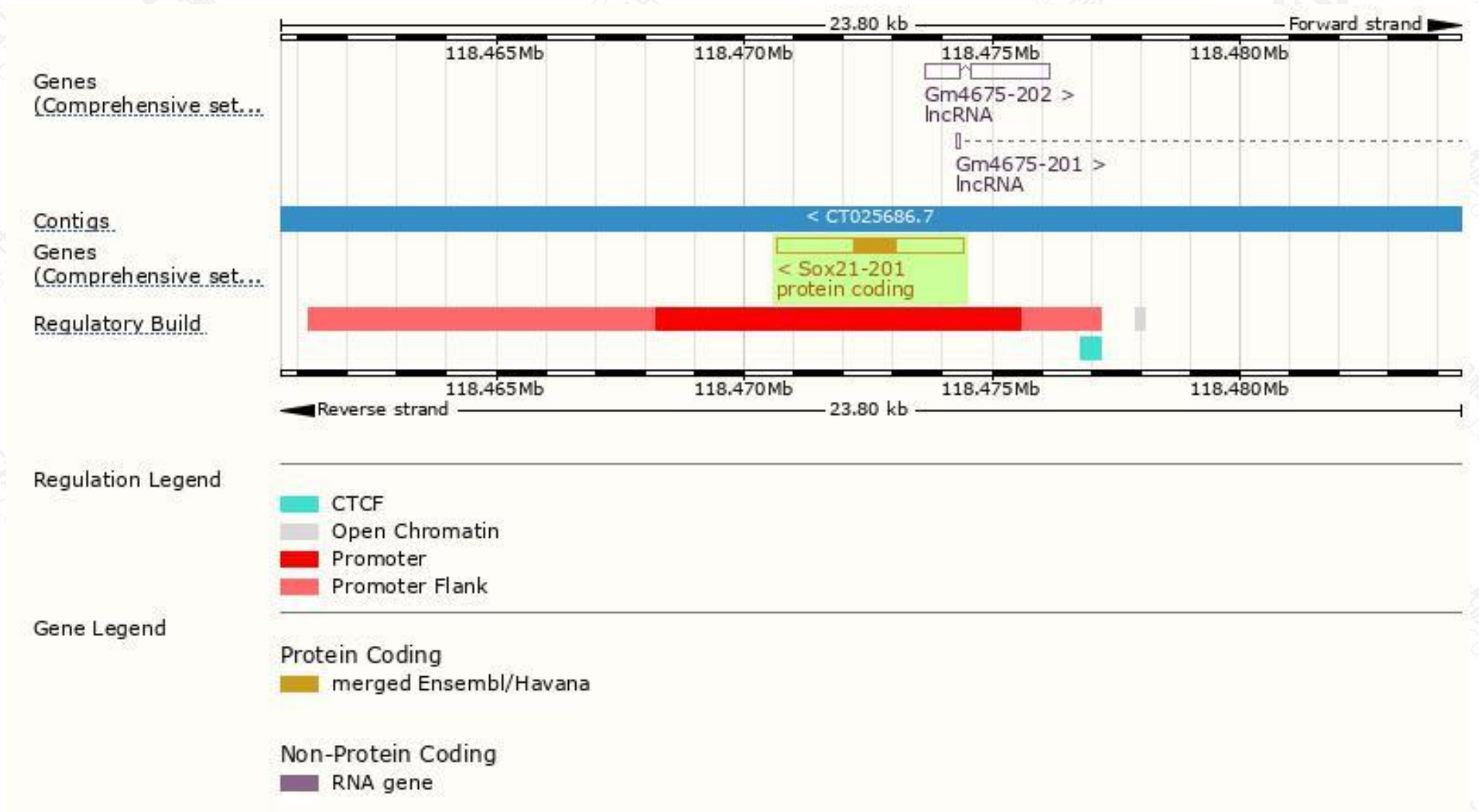
The gene has 1 transcript,and the transcript is shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Sox21-201	ENSMUST00000170662.2	3799	276aa	Protein coding	CCDS27334		TSL:NA , GENCODE basic , APPRIS P1 ,

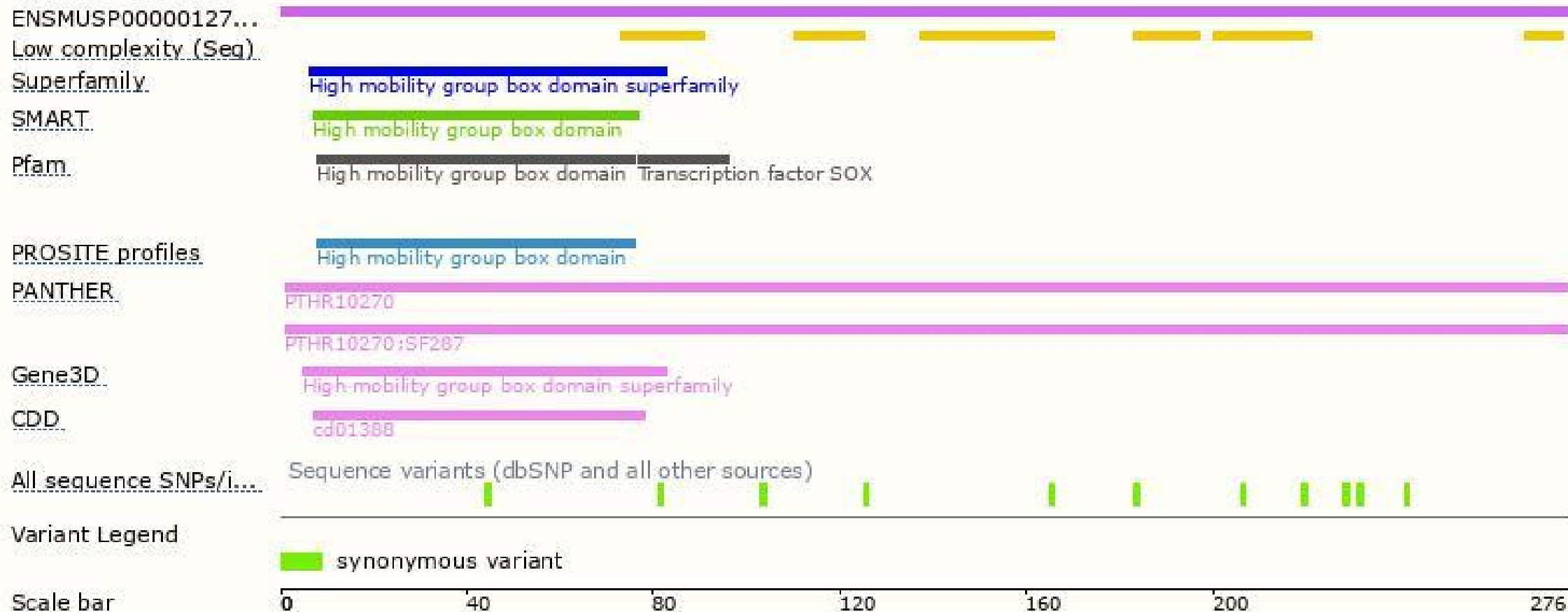
The strategy is based on the design of *Sox21-201* transcript,the transcription is shown below:



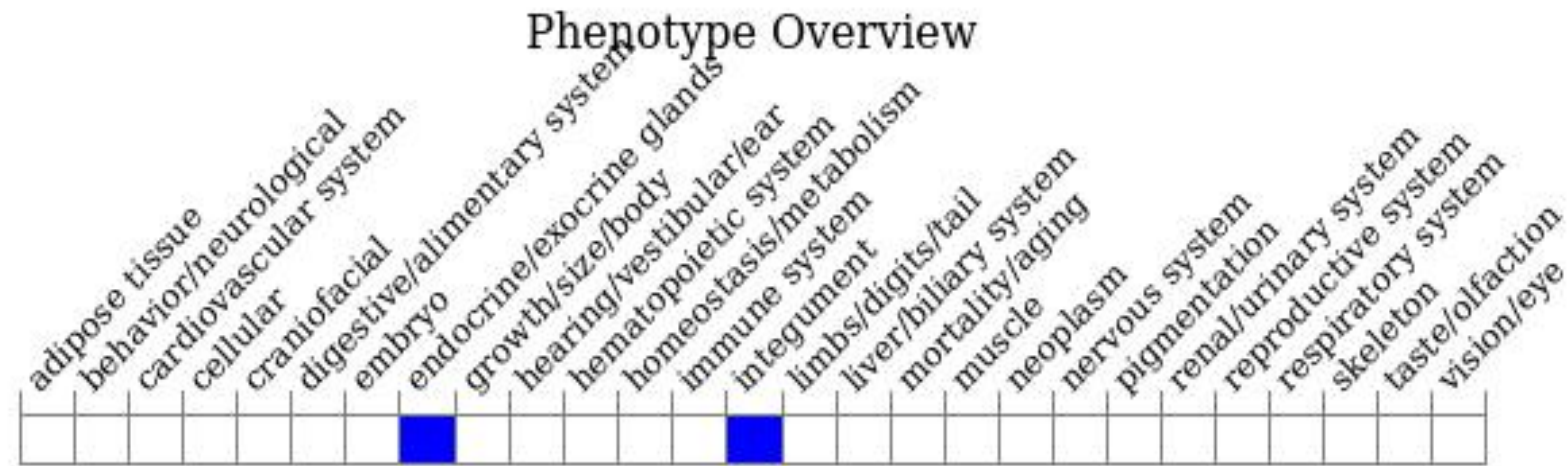
Genomic location distribution



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, mice homozygous for a null mutation display cyclic alopecia, epidermal hyperplasia, enlarged sebaceous glands, and hair shaft and cuticle abnormalities.

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

Tel: 400-9660890

