

Mafb Cas9-KO Strategy

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Overview

Target Gene Name

• Mafb

Project Type

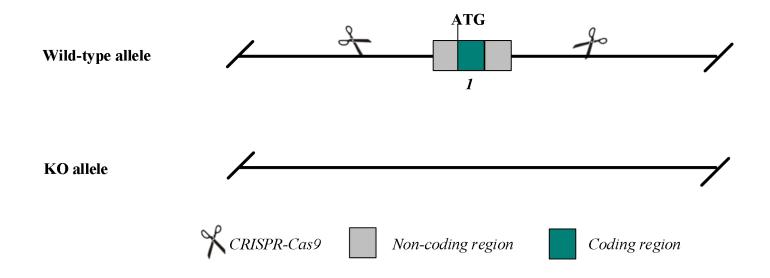
• Cas9-KO

Genetic Background

• C57BL/6JGpt



Strain Strategy

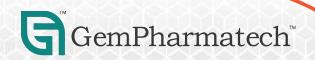


Schematic representation of CRISPR-Cas9 engineering used to edit the Mafb gene.



Technical Information

- The *Mafb* gene has 1 transcript. According to the structure of *Mafb* gene, exon1 of *Mafb*-201 (ENSMUST00000099126.5) transcript is recommended as the knockout region. The region contains 116bp of coding sequences. Knocking out the region will result in disruption of protein function.
- In this project we use CRISPR-Cas9 technology to modify *Mafb* gene. The brief process is as follows: gRNAs were transcribed in vitro. Cas9 and gRNAs 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 ontarget amplicon sequencing. A stable F1-generation mouse strain was obtained by mating positive F0-generation mice with C57BL/6JGpt mice and confirmation of the desired mutant allele was carried out by PCR and on-target amplicon sequencing.



Gene Information



Source: https://www.ncbi.nlm.nih.gov/

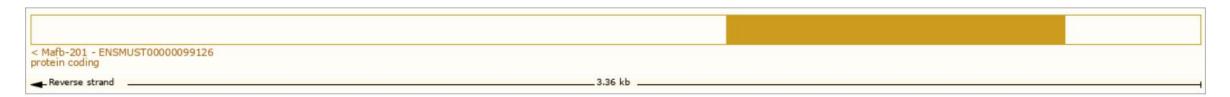


Transcript Information

The gene has 1 transcript, all transcripts are shown below:

Transcript ID	Name 🍦	bp 🛊	Protein	Biotype	CCDS	UniProt Match	Flags			
ENSMUST00000099126.5	Mafb-201	3363	<u>323aa</u>	Protein coding	CCDS16994 €	P54841@	Ensembl Canonical	GENCODE basic	APPRIS P1	TSL:NA

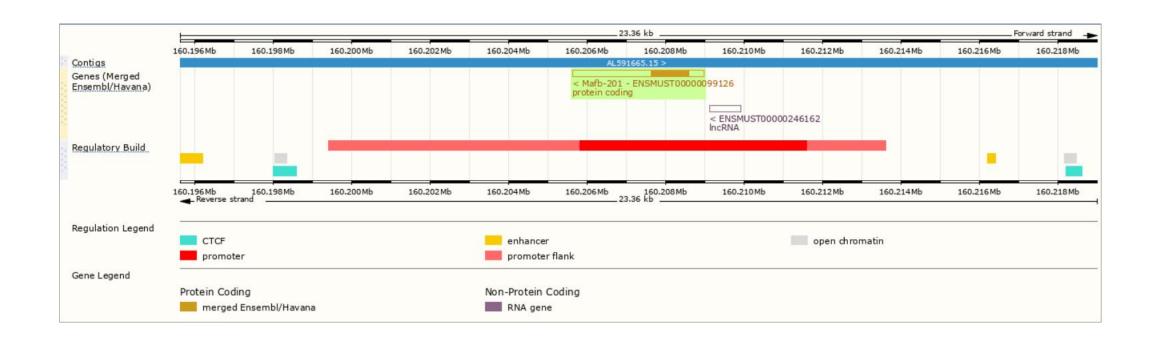
The strategy is based on the design of *Mafb*-201 transcript, the transcription is shown below:

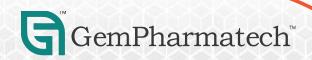


Source: https://www.ensembl.org



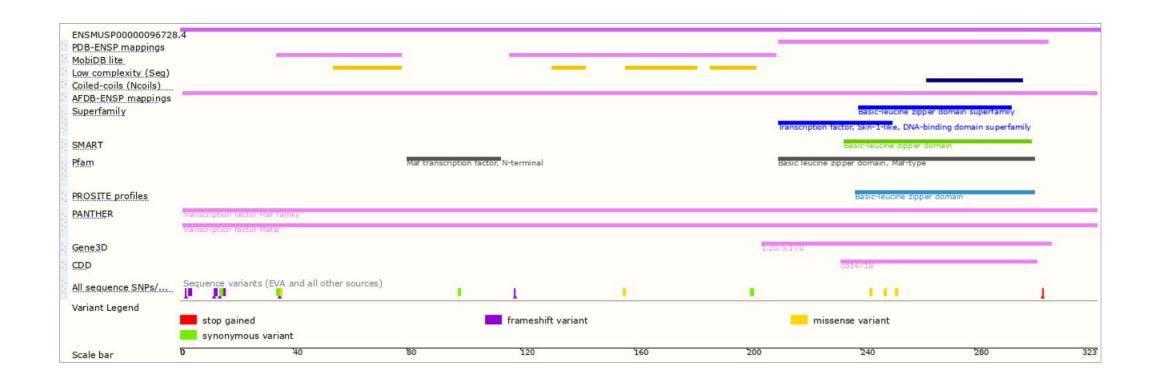
Genomic Information

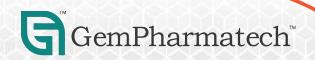




Source: : https://www.ensembl.org

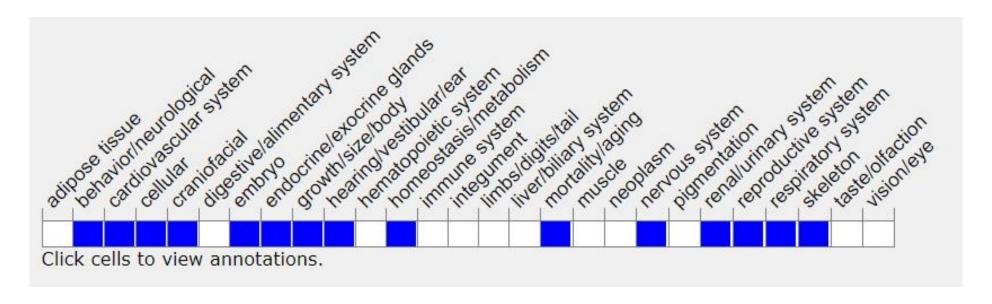
Protein Information





Source: : https://www.ensembl.org

Mouse Phenotype Information (MGI)



• Mutant homozygotes exhibit segmentation defects in the caudal hindbrain, loss of facial motor neurons, impaired inner ear development, arrested maturation of kidney podocytes, reduced fertility, and, in some cases, lethality at birth from apnea. Homozygous KO mice die shortly after birth.



Source: https://www.informatics.jax.org

Important Information

- According to the existing MGI data, Mutant homozygotes exhibit segmentation defects in the caudal hindbrain, loss of facial motor neurons, impaired inner ear development, arrested maturation of kidney podocytes, reduced fertility, and, in some cases, lethality at birth from apnea. Homozygous KO mice die shortly after birth.
- The effect of ENSMUST00000246162.1 gene is unknown.
- *Mafb* is located on Chr4. If the knockout mice are crossed with other mouse strains to obtain double homozygous mutant offspring, please avoid the situation that the second gene is on the same chromosome.
- This Strategy is designed based on genetic information in existing databases. Due to the complexity of biological processes, all risks of the mutation on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.

