

Fgf17 Cas9-KO Strategy

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Overview

Target Gene Name

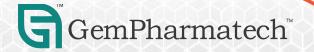
• Fgf17

Project Type

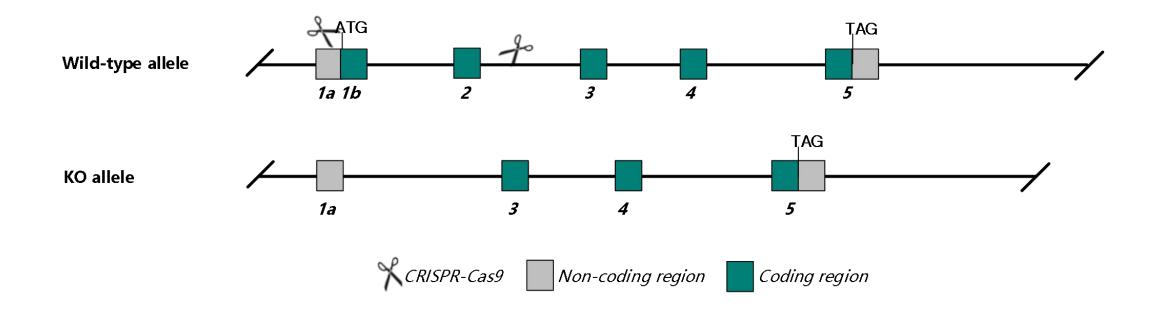
• Cas9-KO

Genetic Background

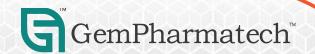
• C57BL/6JGpt



Strain Strategy



Schematic representation of CRISPR-Cas9 engineering used to edit the Fgf17 gene.



Technical Information

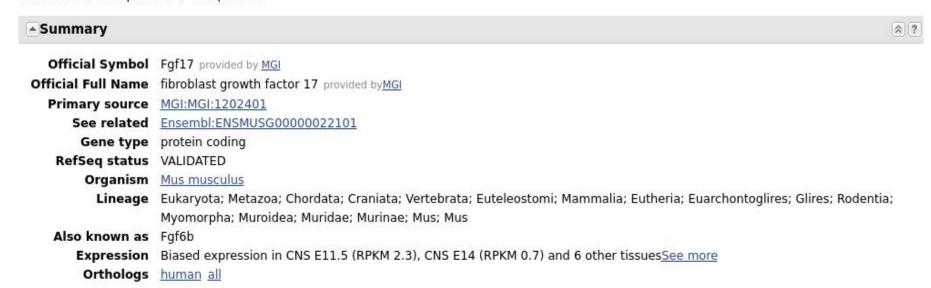
- The *Fgf17* gene has 2 transcripts. According to the structure of *Fgf17* gene, encoding region of exon 1 and exon2 of *Fgf17*-201 (ENSMUST00000022697.7) transcript is recommended as the knockout region. The region contains start codon ATG. Knocking out the region will result in disruption of protein function.
- In this project we use CRISPR-Cas9 technology to modify *Fgf17* 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

Fgf17 fibroblast growth factor 17 [Mus musculus (house mouse)]

Gene ID: 14171, updated on 12-Apr-2023



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

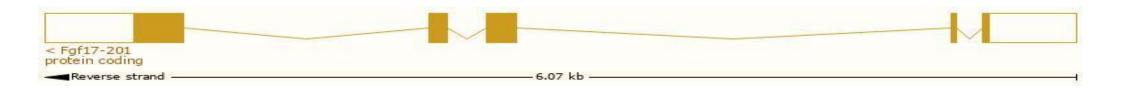


Transcript Information

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

Transcript ID 🗼	Name 🍦	bp 🛊	Protein 🍦	Biotype 👙	CCDS	UniProt Match	Flags			
ENSMUST00000022697.7	Fgf17-201	1689	216aa	Protein coding	CCDS27260 ₽	P63075 & Q0VF19 ₽	Ensembl Canonical	GENCODE basic	APPRIS P2	TSL:1
ENSMUST00000227123.2	Fgf17-202	1336	205aa	Protein coding		B7ZMS7 ₽	GENCODE basic APPRIS ALT1			

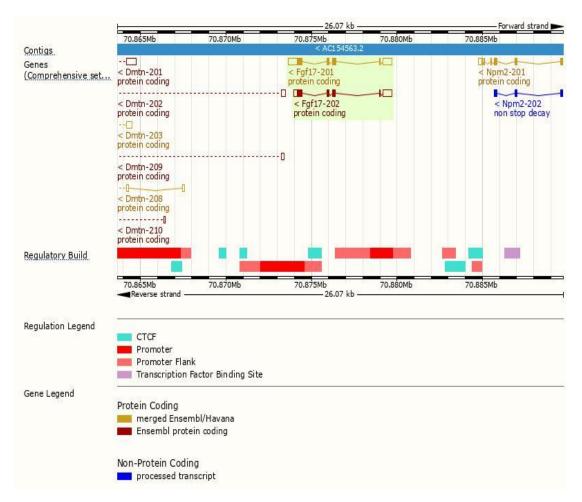
The strategy is based on the design of Fgf17-201 transcript, the transcription is shown below:



Source: https://www.ensembl.org



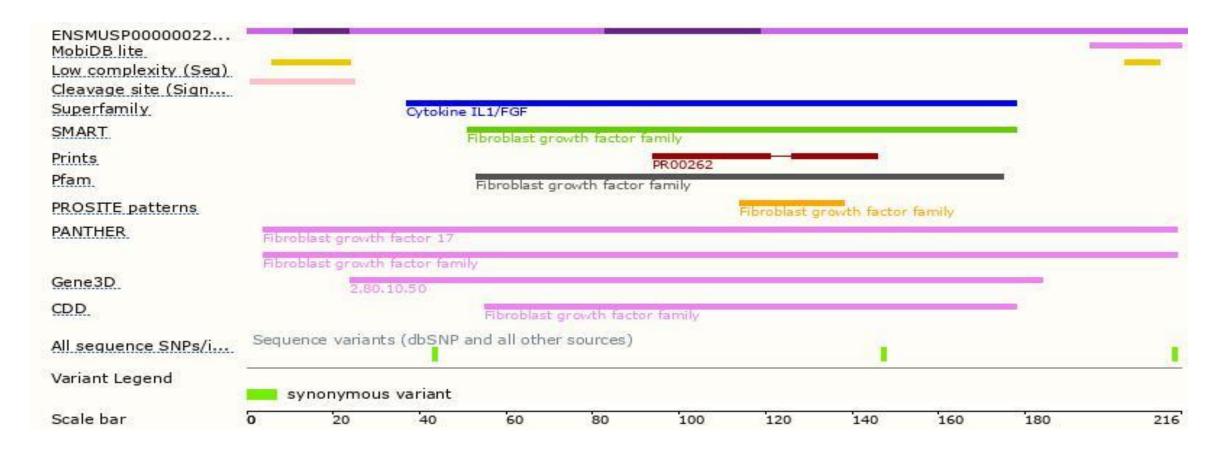
Genomic Information





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

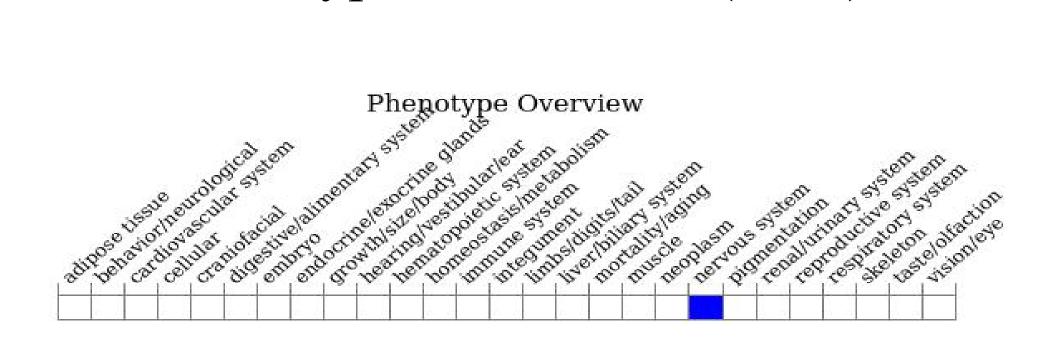
Protein Information





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

Mouse Phenotype Information (MGI)



• Mice homozygous for disruptions in this gene are grossly normal at birth and apparently healthy at birth. However, there are tissue losses in the inferior colliculus and the anterior vermis of the brain.



Important Information

- According to the existing MGI data, mice homozygous for disruptions in this gene are grossly normal at birth and apparently healthy at birth. However, there are tissue losses in the inferior colliculus and the anterior vermis of the brain.
- After deleting the exon containing the starting codon ATG in this strategy, there is a risk of restarting translation.
- Fgf17 is located on Chr14. 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 risk of loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.

