

Fgf14 Cas9-CKO Strategy

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Design Date: 2020/1/22

Project Overview



Project Name

Fgf14

Project type

Cas9-CKO

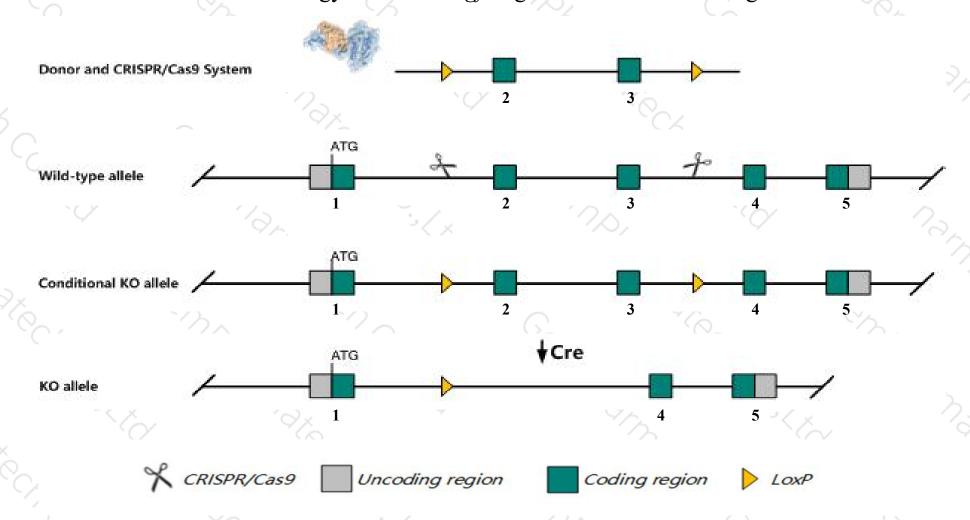
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Fgf14 gene. The schematic diagram is as follows:



Technical routes



- ➤ The Fgf14 gene has 2 transcripts. According to the structure of Fgf14 gene, exon2-exon3 of Fgf14-202 (ENSMUST00000095529.9) transcript is recommended as the knockout region. The region contains 215bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Fgf14* 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.

Notice



- ➤ According to the existing MGI data, Mice homozygous for disruptions in this gene display impaired balance and grip strength.
- The *Fgf14* 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)



△ ?

Fgf14 fibroblast growth factor 14 [Mus musculus (house mouse)]

Gene ID: 14169, updated on 10-Dec-2019

Summary

Official Full Name fibroblast growth factor 14 provided by MGI

Primary source MGI:MGI:109189

Official Symbol Fgf14 provided by MGI

See related Ensembl: ENSMUSG00000025551

Gene type protein coding
RefSeq status VALIDATED
Organism <u>Mus musculus</u>

Lineage Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae;

Murinae; Mus; Mus

Also known as Fhf4; FHF-4; mFHF-4(1B); Tg(tetO-MAPT*P301L)4510Kha

Expression Biased expression in cerebellum adult (RPKM 4.3), cortex adult (RPKM 2.4) and 7 other tissues See more

Orthologs human all

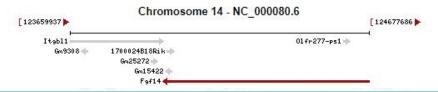
Genomic context

Location: 14 E5; 14 66.18 cM

See Fgf14 in Genome Data Viewer

Exon count: 8

| Annotation release | Status | Assembly | Chr | Location | |
|--------------------|-------------------|------------------------------|-----|--|--|
| 108 | current | GRCm38.p6 (GCF_000001635.26) | 14 | NC_000080.6 (123974441124677686, complement) | |
| Build 37.2 | previous assembly | MGSCv37 (GCF_000001635.18) | 14 | NC_000080.5 (124377513125076349, complement) | |



Transcript information (Ensembl)



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

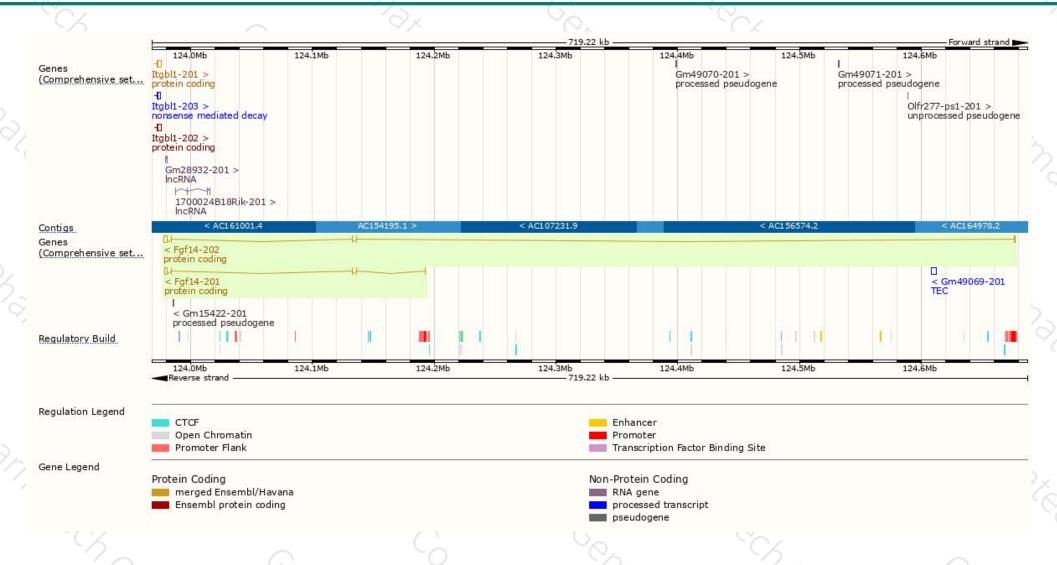
| Name | Transcript ID | bp 🌲 | Protein 🍦 | Biotype | CCDS | UniProt 🍦 | Flags | | |
|-----------|----------------------|------|--------------|----------------|-------------|-----------------|---------------------|------------|-----------|
| Fgf14-202 | ENSMUST00000095529.9 | 3608 | <u>252aa</u> | Protein coding | CCDS27355 ₽ | <u>O89096</u> ₽ | TSL:1 GENCODE basic | | |
| Fgf14-201 | ENSMUST00000026631.5 | 3180 | 247aa | Protein coding | CCDS27356 ₺ | A0A0R4J063₺ | TSL:1 GEN | CODE basic | APPRIS P1 |

The strategy is based on the design of Fgf14-202 transcript, The transcription is shown below



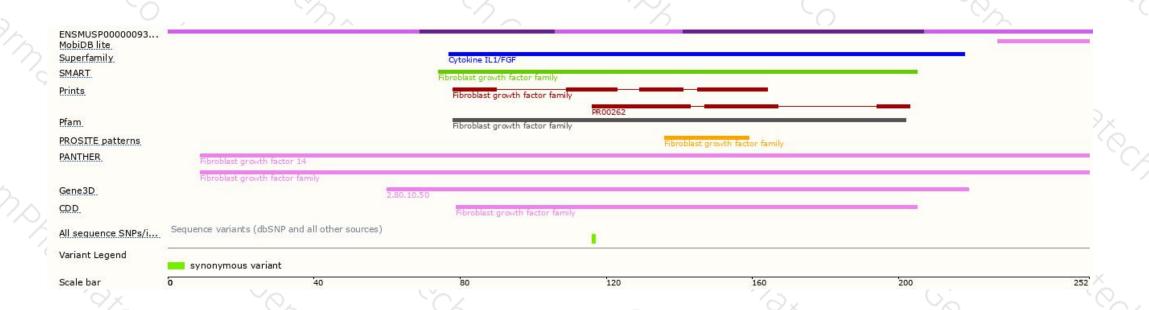
Genomic location distribution





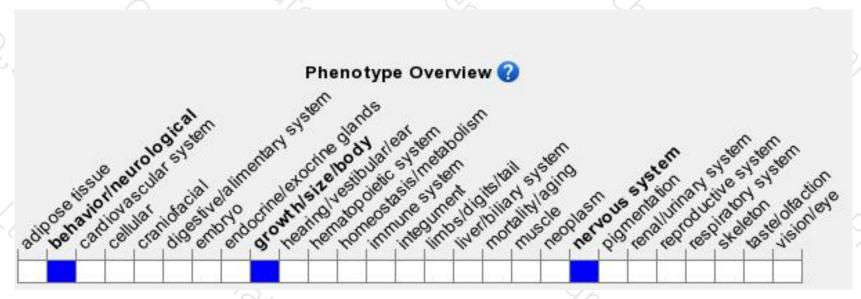
Protein domain





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 disruptions in this gene display impaired balance and grip strength.



If you have any questions, you are welcome to inquire. Tel: 400-9660890





