

Tcf19 Cas9-KO Strategy

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Overview

Target Gene Name

• *Tcf19*

Project Type

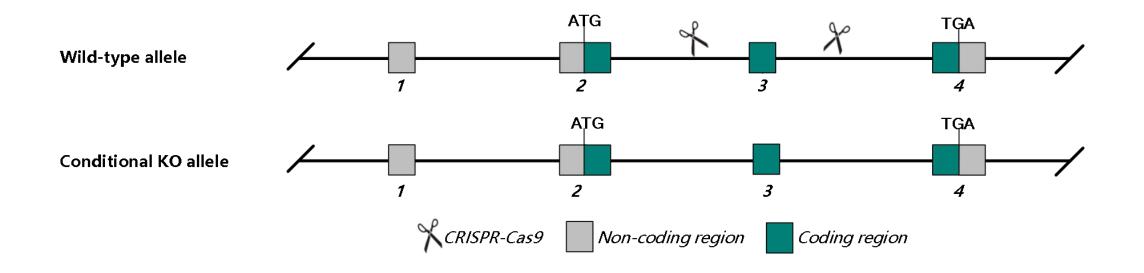
• Cas9-KO

Genetic Background

• C57BL/6JGpt



Strain Strategy



Schematic representation of CRISPR-Cas9 engineering used to edit the Tcf19 gene.

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Technical Information

- The *Tcf19* gene has 4 transcripts. According to the structure of *Tcf19* gene, exon 3 of *Tcf19*-202 (ENSMUST00000160885.2) is recommended as the knockout region. The region contains 547 bp of coding sequences. Knocking out the region will result in disruption of gene function.
- In this project we use CRISPR-Cas9 technology to modify *Tcf19* 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 on-target 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

Full Report -

Tcf19 transcription factor 19 [Mus musculus (house mouse)]

Gene ID: 106795, updated on 26-Sep-2022

Summary

Official Symbol Tcf19 provided by MGI Official Full Name transcription factor 19 provided by MGI Primary source MGI:MGI:103180 See related Ensembl:ENSMUSG00000050410 AllianceGenome:MGI:103180 Gene type protein coding RefSeg 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 5730403J10Rik Summary Acts upstream of or within G1/S transition of mitotic cell cycle; regulation of gene expression; and type B pancreatic cell proliferation. Located in nucleus. Is expressed in forebrain; genitourinary system; and limb. Orthologous to human TCF19 (transcription factor 19). [provided by Alliance of Genome Resources, Apr 2022] Expression Broad expression in limb E14.5 (RPKM 21.9), thymus adult (RPKM 20.3) and 25 other tissues See more Orthologs human all NEW Try the new Gene table Try the new Transcript table Genomic context ☆ ? Location: 17: 17 B1 See Tcf19 in Genome Data Viewer Exon count: 4 Source: https://www.ncbi.nlm.nih.gov/gene/106795



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Transcript Information

The gene has 4 transcripts, all transcripts are shown below:

Transcript ID 🔹	Name 🖕	bp 🖕	Protein 🖕	Biotype 💧	CCDS 🖕	UniProt Match	Flags
ENSMUST00000162683.2	Tcf19-204	873	<u>147aa</u>	Protein coding		<u>E0CZ64</u> &	TSL:1 CDS 3' incomplete
ENSMUST00000161012.8	Tcf19-203	1665	<u>263aa</u>	Protein coding	<u>CCDS50089</u> 교	<u>G3XA31</u> ₽	GENCODE basic APPRIS P1 TSL:1
ENSMUST00000160885.2	Tcf19-202	1732	<u>263aa</u>	Protein coding	<u>CCDS50089</u> &	<u>G3XA31</u> ₽	Ensembl Canonical GENCODE basic APPRIS P1 TSL:1
ENSMUST00000159009.2	Tcf19-201	982	<u>201aa</u>	Protein coding		<u>E0CY69</u> 교	TSL:2 CDS 3' incomplete

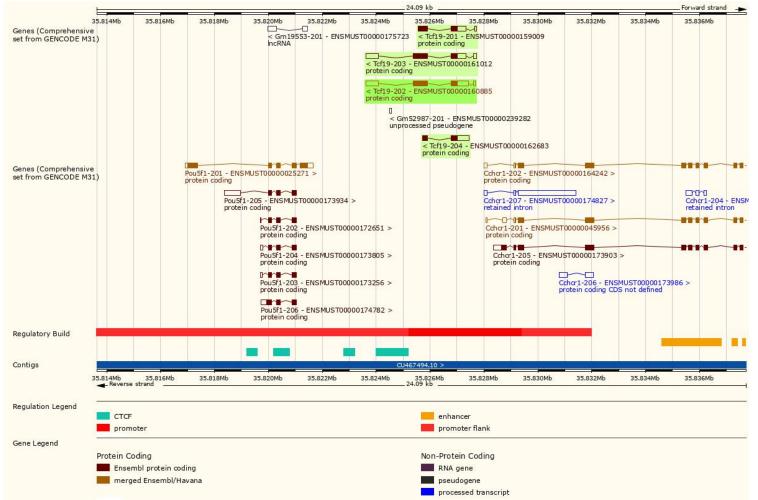
The strategy is based on the design of *Tcf19*-202 transcript, the transcription is shown below:



Source: http://asia.ensembl.org/



Genomic Information



Source:http://asia.ensembl.org/

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Important Information

- The knockout region is about 1.8 kb away from the 5th end of the *Cchcr1* gene, which may affect the regulation of these gene.
- The knockout region is about 500 bp away from the 5th end of the *Gm52987* gene, which may affect the regulation of this gene.
- The knockout region is about 3.5 kb away from the 5th end of the *Gm19553* gene, which may affect the regulation of this gene.
- A part of amino acid sequence will still remain at the N-terminal of the *Tcf19* gene.
- *Tcf19* is located on Chr 17. 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.

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