

Tfec Cas9-CKO Strategy

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Design Date:

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Project Overview

Project Name

Tfec

Project type

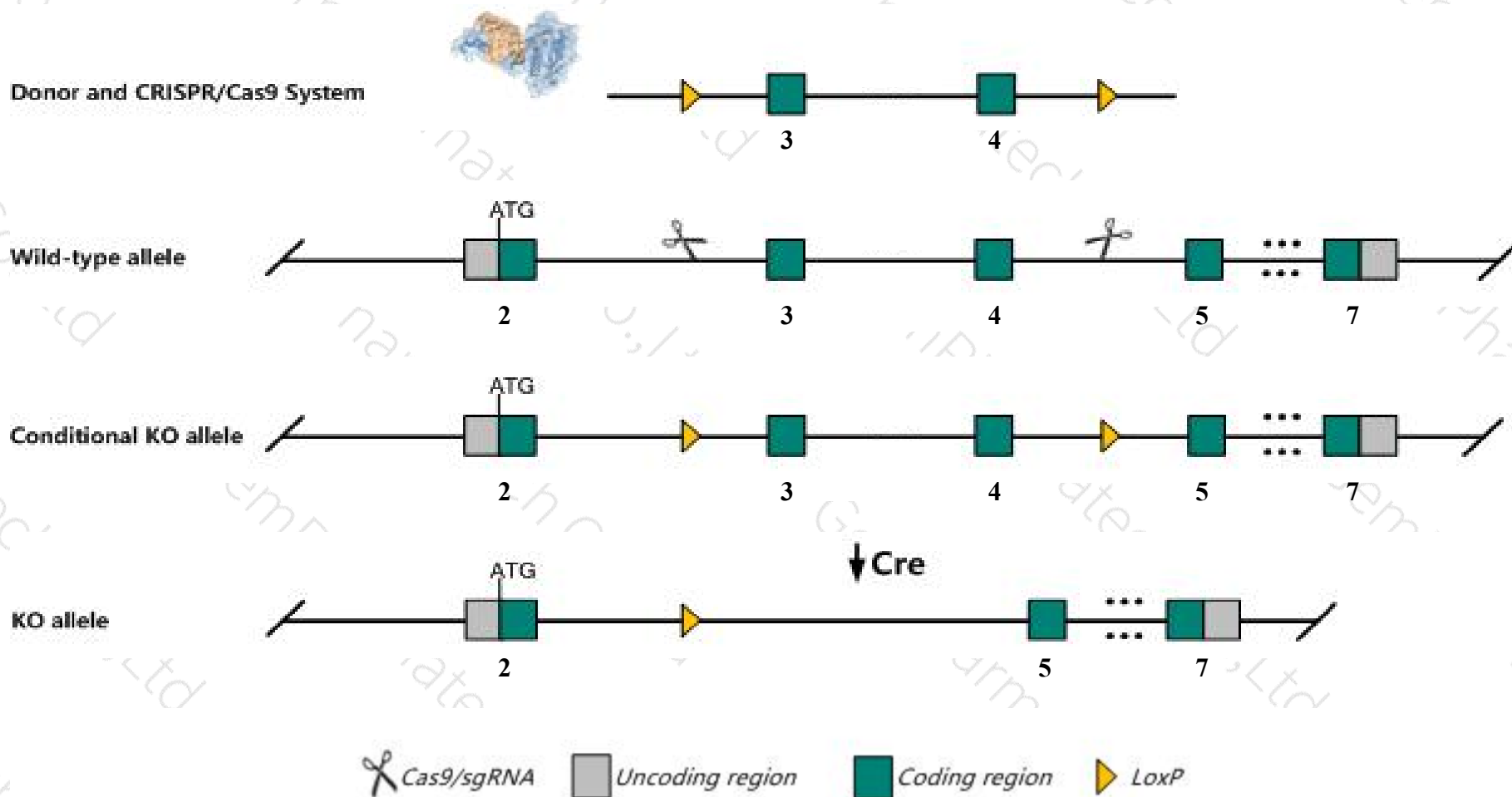
Cas9-CKO

Strain background

C57BL/6JGpt

Conditional Knockout strategy

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



- The *Tfec* gene has 5 transcripts. According to the structure of *Tfec* gene, exon3-exon4 of *Tfec-201* (ENSMUST00000031533.10) transcript is recommended as the knockout region. The region contains 172bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Tfec* 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 a knock-out allele are viable and fertile, normally pigmented, have normal eyes and mast cells, and show no evidence of osteopetrosis.
- The *Tfec* gene is located on the Chr6. 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)

Tfec transcription factor EC [*Mus musculus* (house mouse)]

Gene ID: 21426, updated on 13-Mar-2020

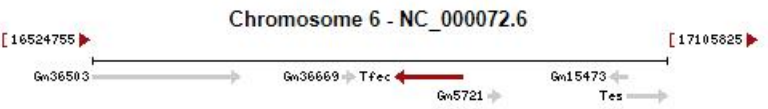
Summary

Official Symbol Tfec provided by [MGI](#)
Official Full Name transcription factor EC provided by [MGI](#)
Primary source [MGI:MGI:1333760](#)
See related [Ensembl:ENSMUSG00000029553](#)
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 Tcfec; bHLHe34; BB107417
Expression Biased expression in placenta adult (RPKM 1.9), liver E18 (RPKM 1.4) and 12 other tissues [See more](#)
Orthologs [human](#) [all](#)

Genomic context

Location: 6; 6 A2 [See Tfec in Genome Data Viewer](#)
Exon count: 9

Annotation release	Status	Assembly	Chr	Location
108	current	GRCm38.p6 (GCF_000001635.26)	6	NC_000072.6 (16830927..16898464, complement)
Build 37.2	previous assembly	MGSCv37 (GCF_000001635.18)	6	NC_000072.5 (16783381..16848441, complement)

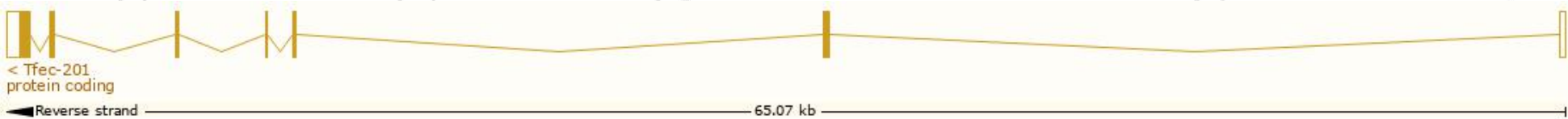


Transcript information (Ensembl)

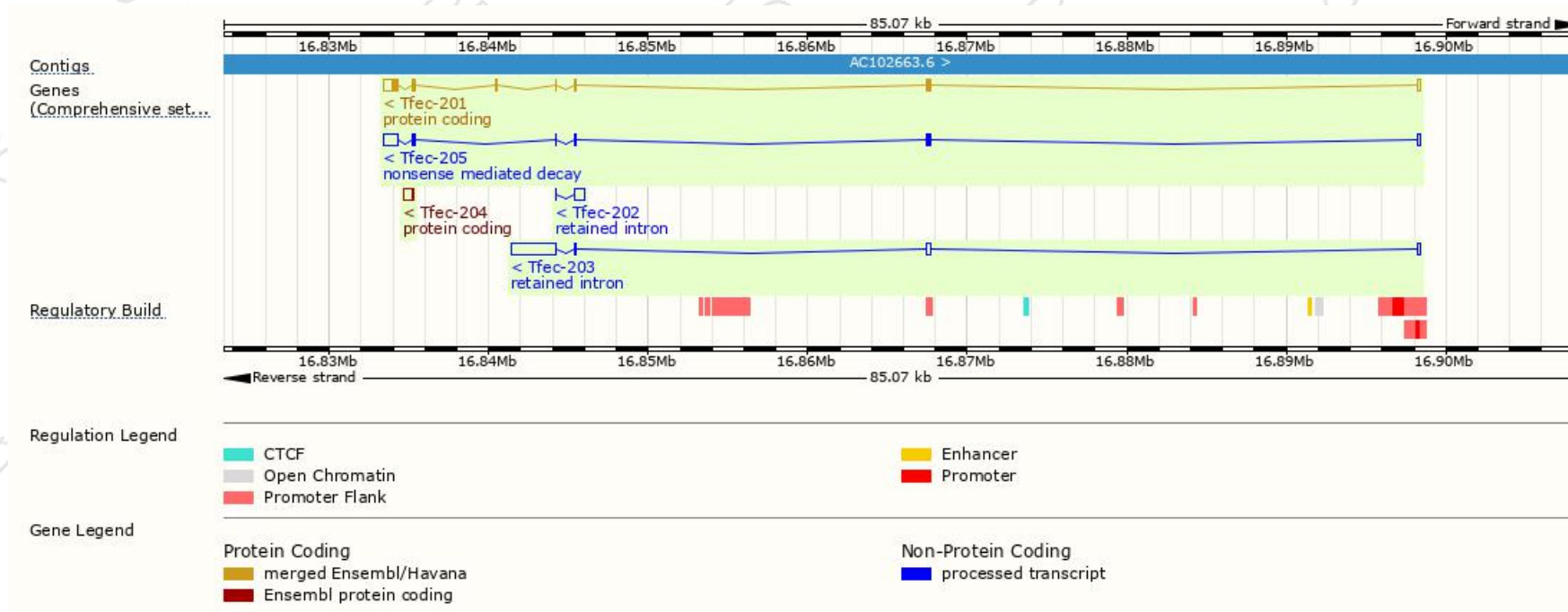
The gene has 5 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Tfec-201	ENSMUST000000031533.10	1787	317aa	Protein coding	CCDS19920	Q9WTW4	TSL:1 GENCODE basic APPRIS P1
Tfec-204	ENSMUST000000201406.1	572	28aa	Protein coding	-	A0A0J9YV19	CDS 5' incomplete TSL:NA
Tfec-205	ENSMUST000000202997.1	1654	127aa	Nonsense mediated decay	-	A0A0J9YTW5	TSL:1
Tfec-203	ENSMUST000000201104.1	3384	No protein	Retained intron	-	-	TSL:1
Tfec-202	ENSMUST000000200984.1	688	No protein	Retained intron	-	-	TSL:5

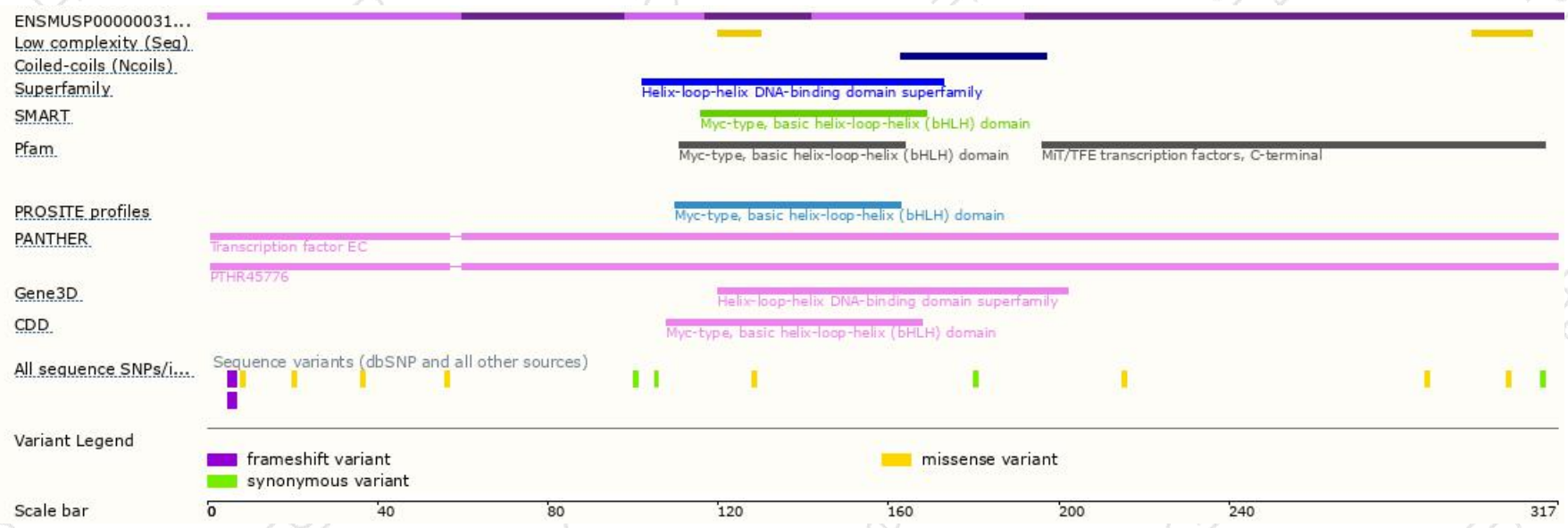
The strategy is based on the design of *Tfec-201* transcript, the transcription is shown below



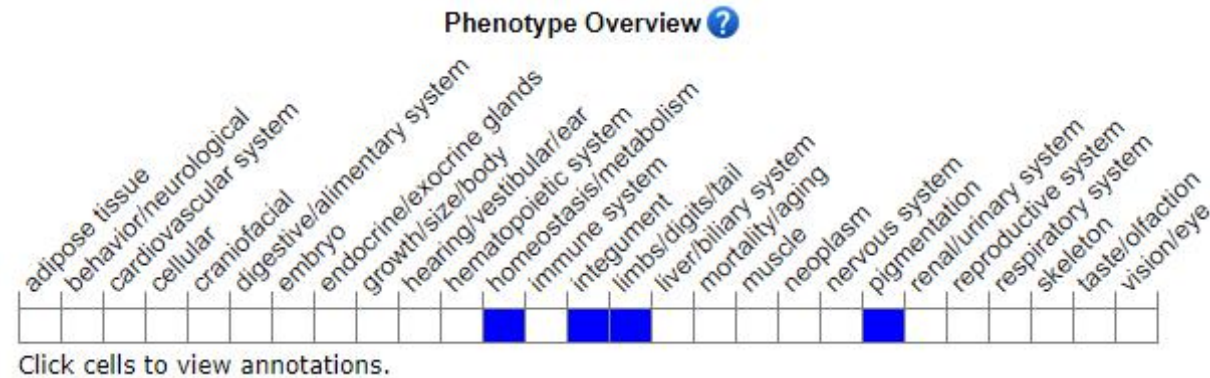
Genomic location distribution



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 knock-out allele are viable and fertile, normally pigmented, have normal eyes and mast cells, and show no evidence of osteopetrosis..

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

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