

Nfatc4 Cas9-KO Strategy

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Project Overview



Project Name Nfatc4

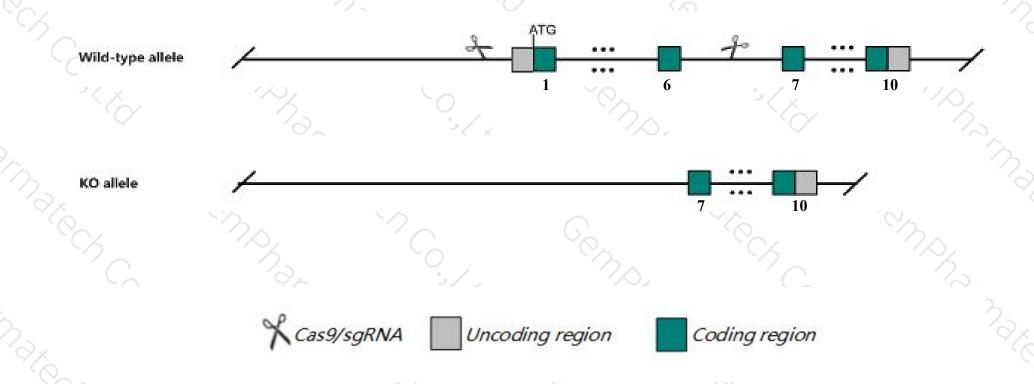
Project type Cas9-KO

Strain background C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Nfatc4* gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Nfatc4* gene has 11 transcripts. According to the structure of *Nfatc4* gene, exon1-exon6 of *Nfatc4-201* (ENSMUST00000024179.5) transcript is recommended as the knockout region. The region contains start codon ATG. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Nfatc4* gene. The brief process is as follows: CRISPR/Cas9 system

Notice



- ➤ According to the existing MGI data, Mice homozygous for a knock-out allele are viable and overtly normal and exhibit normal embryonic heart morphology as well as normal pathophysiologic cardiac hypertrophy in response to angiotensin II infusion or aortic banding.
- > The *Nfatc4* 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 the gene knockout on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.

Gene information (NCBI)



Nfatc4 nuclear factor of activated T cells, cytoplasmic, calcineurin dependent 4 [Mus musculus (house mouse)]

Gene ID: 73181, updated on 12-Aug-2019

Summary



Official Symbol Nfatc4 provided by MGI

Official Full Name nuclear factor of activated T cells, cytoplasmic, calcineurin dependent 4 provided by MGI

Primary source MGI:MGI:1920431

See related Ensembl:ENSMUSG00000023411

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 Nfat3; AW107667; AW546455; 3110041H08Rik

Expression Biased expression in limb E14.5 (RPKM 41.5), CNS E11.5 (RPKM 14.6) and 14 other tissues See more

Orthologs <u>human</u> all

Transcript information (Ensembl)



The gene has 11 transcripts, all transcripts are shown below:

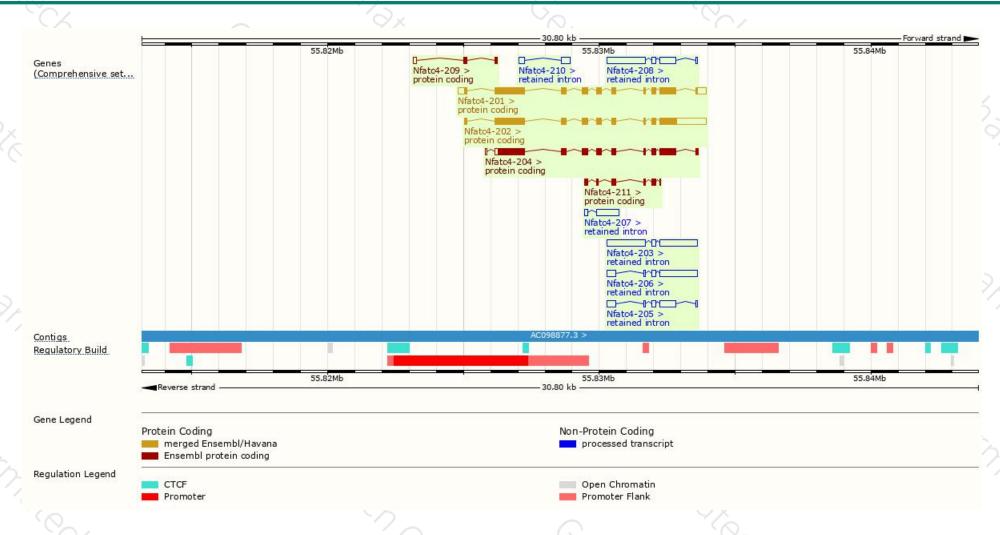
Name 🍦	Transcript ID 👙	bp 🛊	Protein 🍦	Biotype	CCDS 🍦	UniProt 🍦	Flags
Nfatc4-202	ENSMUST00000172271.8	3775	890aa	Protein coding	CCDS49500@	Q8K120 ₪	TSL:1 GENCODE basic APPRIS ALT2
Nfatc4-201	ENSMUST00000024179.5	3267	<u>901aa</u>	Protein coding	CCDS27132 ₽	Q8K120 ₪	TSL:1 GENCODE basic APPRIS P3
Nfatc4-204	ENSMUST00000226357.1	2680	831aa	Protein coding	-	A0A2I3BRR0 ₽	GENCODE basic APPRIS ALT2
Nfatc4-211	ENSMUST00000228308.1	509	<u>170aa</u>	Protein coding	-	A0A2I3BRX2函	CDS 5' and 3' incomplete
Nfatc4-209	ENSMUST00000226979.1	325	<u>58aa</u>	Protein coding	-	A0A2I3BS51₽	CDS 3' incomplete
Nfatc4-203	ENSMUST00000226293.1	2969	No protein	Retained intron	-	-	-
Nfatc4-208	ENSMUST00000226869.1	2217	No protein	Retained intron	-	-	-
Nfatc4-206	ENSMUST00000226716.1	1913	No protein	Retained intron	-	-	-
Nfatc4-205	ENSMUST00000226536.1	1161	No protein	Retained intron	-	-	-
Nfatc4-207	ENSMUST00000226834.1	916	No protein	Retained intron	-	-	-
Nfatc4-210	ENSMUST00000227746.1	540	No protein	Retained intron	-	-	*

The strategy is based on the design of Nfatc4-201 transcript, The transcription is shown below



Genomic location distribution





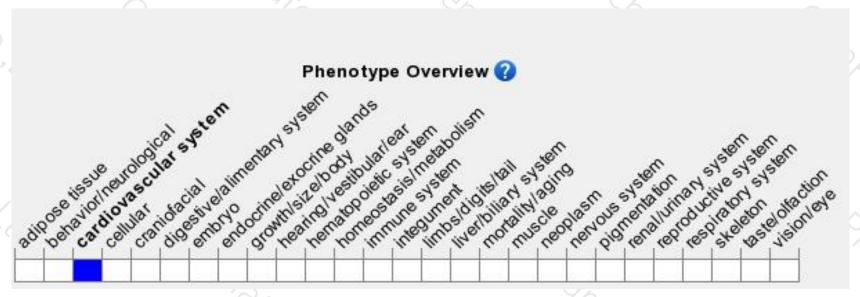
Protein domain





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 overtly normal and exhibit normal embryonic heart morphology as well as normal pathophysiologic cardiac hypertrophy in response to angiotensin II infusion or aortic banding.



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





