

Fxn Cas9-CKO Strategy

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Project Overview



Project Name Fxn

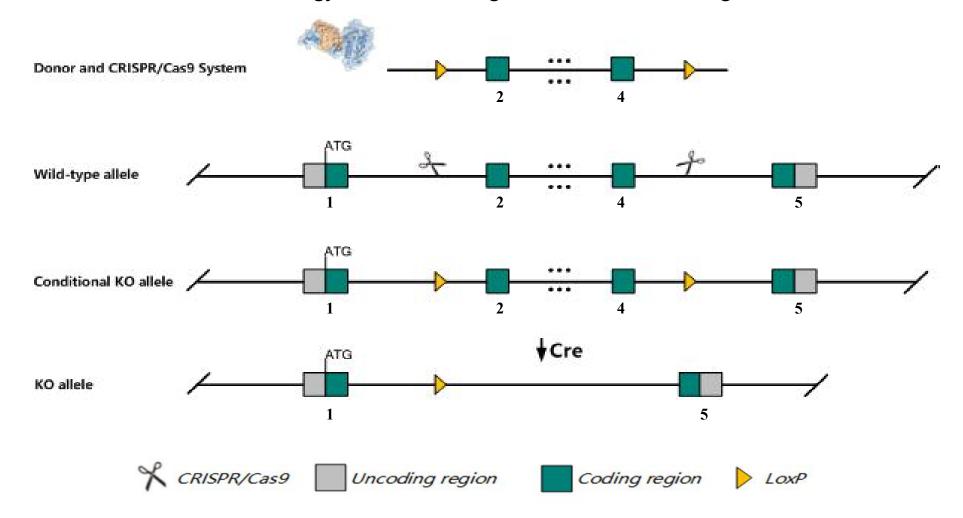
Project type Cas9-CKO

Strain background C57BL/6JGpt

Conditional Knockout strategy



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



Technical routes



The Fxn gene has 3 transcripts. According to the structure of Fxn gene, exon2-exon4 of Fxn-201

(ENSMUST00000081333.10) transcript is recommended as the knockout region. The region contains 314bp coding sequence. Knock out the region will result in disruption of protein function.

In this project we use CRISPR/Cas9 technology to modify *Fxn* 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, Homozygotes for a targeted null mutation exhibit early post-implantation lethality, in the absence of intramitochondrial iron accumulation. Conditional knockouts, specific to striated muscle and neuron/striated muscle, show cardiac hypertrophy and large sensory neuron dysfunction, respectively.

The *Fxn* gene is located on the Chr19. 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



Fxn frataxin [Mus musculus (house mouse)]

Gene ID: 14297, updated on 31-Jan-2019

Summary

☆ ?

Official Symbol Fxn provided by MGI

Official Full Name frataxin provided by MGI

Primary source MGI:MGI:1096879

See related Ensembl:ENSMUSG00000059363

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 FA, FARR, Frda, X25

Expression Ubiquitous expression in adrenal adult (RPKM 20.6), liver E14.5 (RPKM 15.3) and 28 other tissuesSee more

Orthologs <u>human all</u>

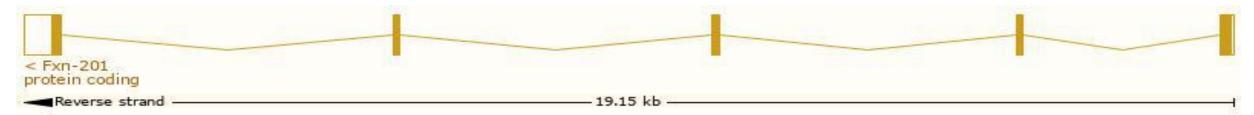
Transcript information Ensembl



The gene has 3 transcripts, all transcripts are shown below:

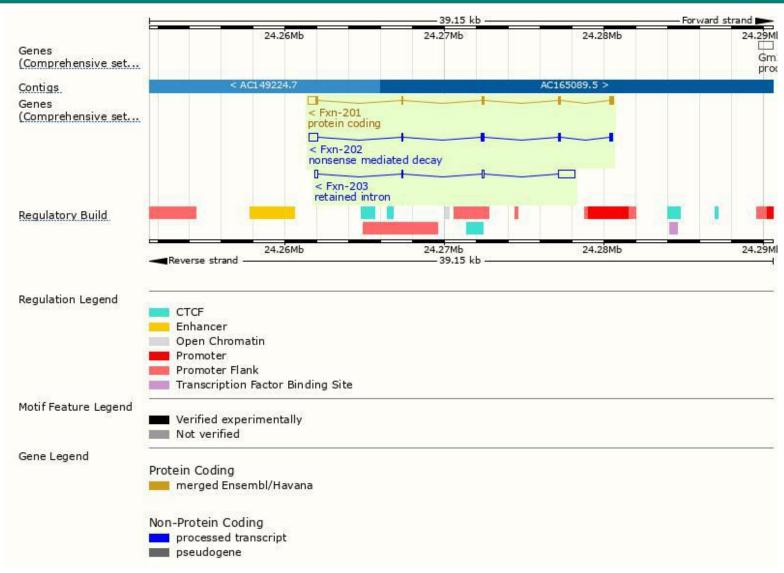
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Fxn-201	ENSMUST00000081333.10	1114	<u>207aa</u>	Protein coding	CCDS29711	O35943 Q3TV21	TSL:1 GENCODE basic APPRIS P1
Fxn-202	ENSMUST00000123684.1	1072	<u>145aa</u>	Nonsense mediated decay		E9Q2P9	TSL:1
Fxn-203	ENSMUST00000132688.1	1402	No protein	Retained intron	-	-	TSL:1

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



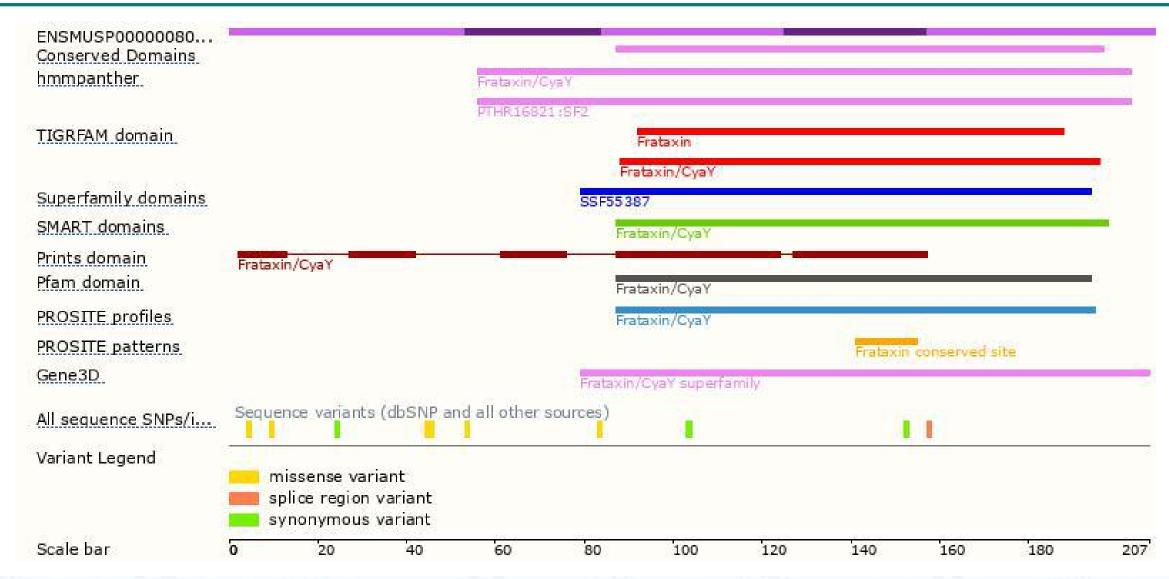
Genomic location distribution





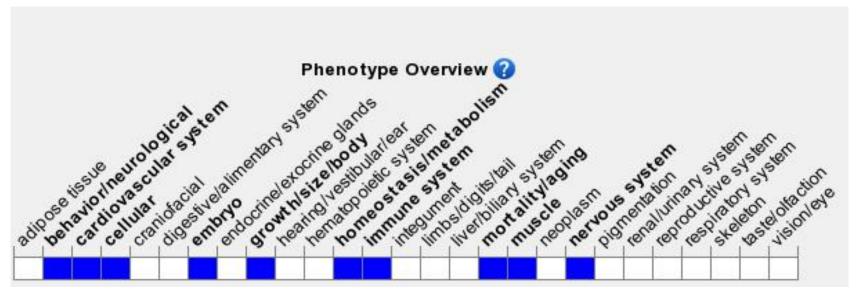
Protein domain





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, Homozygotes for a targeted null mutation exhibit early post-implantation lethality, in the absence of intramitochondrial iron accumulation. Conditional knockouts, specific to striated muscle and neuron/striated muscle, show cardiac hypertrophy and large sensory neuron dysfunction, respectively.



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





