

Fbln2 Cas9-CKO Strategy

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Reviewer:

Design Date: 2019-12-23

Project Overview



Project Name

Fbln2

Project type

Cas9-CKO

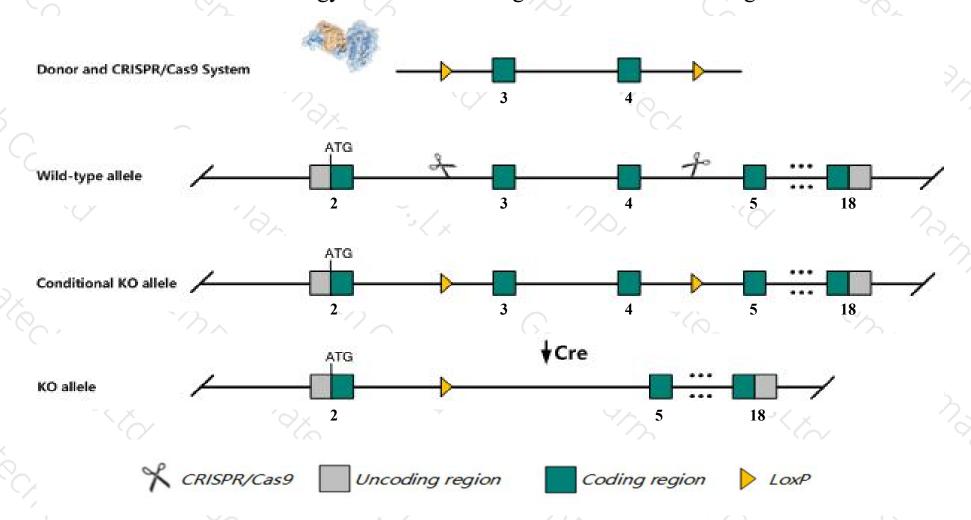
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Fbln2 gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Fbln2* gene has 9 transcripts. According to the structure of *Fbln2* gene, exon3-exon4 of *Fbln2-201*(ENSMUST00000041544.7) transcript is recommended as the knockout region. The region contains 242bp coding sequence.

 Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Fbln2* 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 alle are viable, fertile and grossly normal with no apparent defects in elastic fiber formation in the aorta and skin.
- ➤ Transcripts *Fbln2*-203/205/208 CDS are incomplete ,whether they will be affected is unknown.
- > The *Fbln2* 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)



Fbln2 fibulin 2 [Mus musculus (house mouse)]

Gene ID: 14115, updated on 12-Aug-2019

Summary

≈ ?

Official Symbol Fbln2 provided by MGI

Official Full Name fibulin 2 provided by MGI

Primary source MGI:MGI:95488

See related Ensembl: ENSMUSG00000064080

Gene type protein coding
RefSeq status VALIDATED
Organism Mus musculus

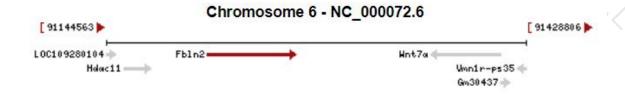
Lineage Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha;

Muroidea; Muridae; Murinae; Mus; Mus

Also known as FIBL-2; 5730577E14Rik

Expression Broad expression in limb E14.5 (RPKM 77.0), subcutaneous fat pad adult (RPKM 55.8) and 20 other tissues See more

Orthologs human all



Transcript information (Ensembl)



The gene has 9 transcripts, all transcripts are shown below:

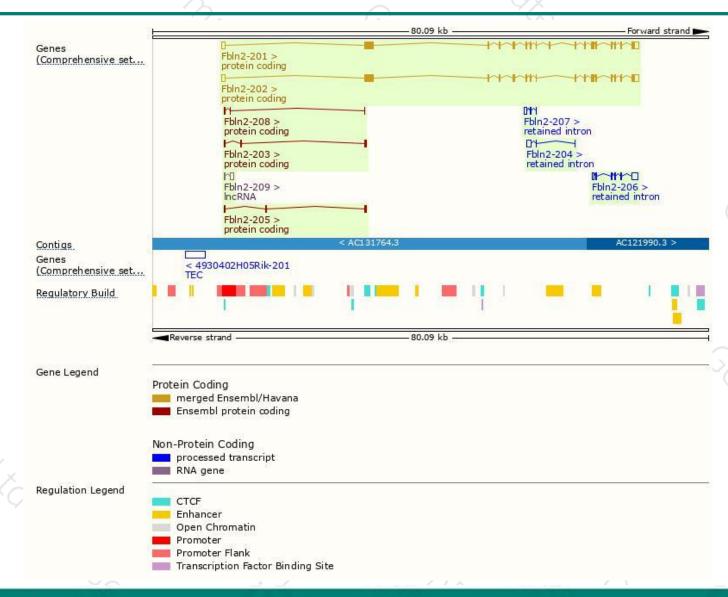
Name 🍦	Transcript ID 👙	bp 🍦	Protein	Translation ID 🝦	Biotype	CCDS 🍦	UniProt 🍦	Flags
Fbln2-201	ENSMUST00000041544.7	4786	<u>1221aa</u>	ENSMUSP00000048334.7	Protein coding	CCDS39566@	P37889 ₽	TSL:1 GENCODE basic APPRIS P1
FbIn2-202	ENSMUST00000113498.8	4644	<u>1174aa</u>	ENSMUSP00000109126.2	Protein coding	CCDS39567₽	<u>P37889</u> @ <u>Q3TGL4</u> @	TSL:1 GENCODE basic
FbIn2-203	ENSMUST00000132021.1	477	<u>69aa</u>	ENSMUSP00000116456.1	Protein coding	(+)	<u>D3Z5B0</u> ₽	CDS 3' incomplete TSL:5
FbIn2-205	ENSMUST00000134974.1	377	<u>76aa</u>	ENSMUSP00000116302.1	Protein coding	(±)	<u>D3Z5S2</u> ₽	CDS 3' incomplete TSL:2
FbIn2-208	ENSMUST00000153364.1	223	<u>10aa</u>	ENSMUSP00000120101.1	Protein coding	0±0	A0A1C7ZMZ9₽	CDS 3' incomplete TSL:5
FbIn2-206	ENSMUST00000137029.1	1612	No protein	-	Retained intron	(±)	-	TSL:1
FbIn2-204	ENSMUST00000134286.1	752	No protein	-	Retained intron	(±)	-	TSL:5
Fbln2-207	ENSMUST00000147371.7	660	No protein	-	Retained intron	(+)	-	TSL:2
Fbln2-209	ENSMUST00000203406.1	669	No protein	-	IncRNA	0. 0 0	-	TSL:1

The strategy is based on the design of *Fbln2-201* transcript, The transcription is shown below



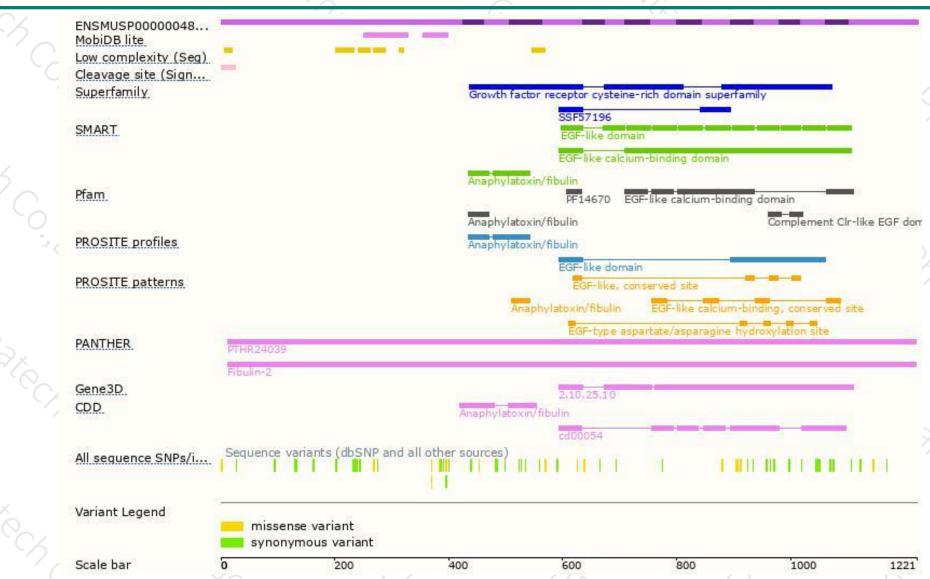
Genomic location distribution





Protein domain







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





