

Ltbp1 Cas9-CKO Strategy

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Project Overview



Project Name Ltbp1

Project type Cas9-CKO

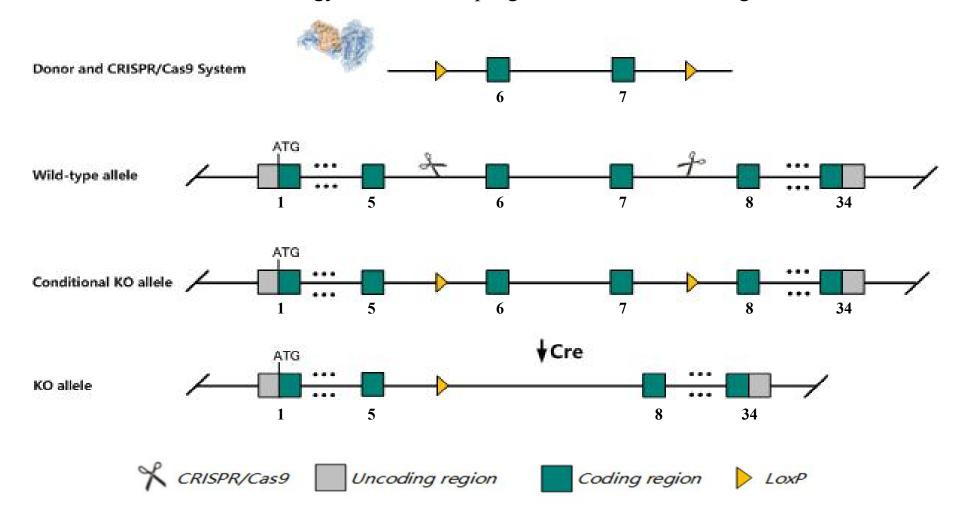
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Ltbp1* gene. The schematic diagram is as follows:



Technical routes



The *Ltbp1* gene has 10 transcripts. According to the structure of *Ltbp1* gene, exon6-exon7 of *Ltbp1-201* (ENSMUST0000001927.11) transcript is recommended as the knockout region. The region contains 500bp coding sequence. Knock out the region will result in disruption of protein function.

In this project we use CRISPR/Cas9 technology to modify *Ltbp1* 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 null allele exhibit embryonic and neonatal lethality associated with defects in the aortic arch and outflow tract.

Transcript *Ltbp1-209* may not be affected.

The *Ltbp1* gene is located on the Chr17. 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



Ltbp1 latent transforming growth factor beta binding protein 1 [Mus musculus (house mouse)]

Gene ID: 268977, updated on 3-Feb-2019

Summary



Official Symbol Ltbp1 provided by MGI

Official Full Name latent transforming growth factor beta binding protein 1 provided by MGI

Primary source MGI:MGI:109151

See related Ensembl: ENSMUSG00000001870

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 9430031G15Rik, 9830146M04, Ltbp-1, Ltbp1L, TGF-beta1-BP-1, Tgfb, b2b1000Clo

Expression Broad expression in ovary adult (RPKM 29.6), limb E14.5 (RPKM 18.7) and 16 other tissuesSee more

Orthologs <u>human all</u>

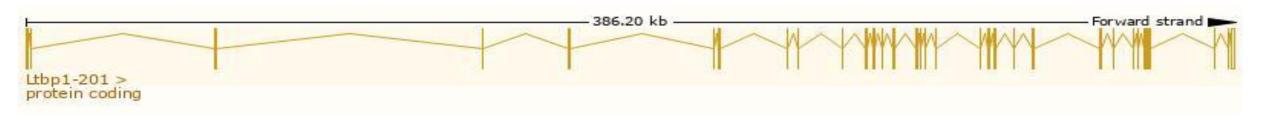
Transcript information Ensembl



The gene has 10 transcripts, all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Ltbp1-201	ENSMUST00000001927.11	6671	<u>1712aa</u>	Protein coding	CCDS28973	Q8CG19	TSL:5 GENCODE basic
Ltbp1-203	ENSMUST00000112516.7	5140	<u>1394aa</u>	Protein coding	CCDS37694	Q8CG19	TSL:5 GENCODE basic
Ltbp1-202	ENSMUST00000112514.1	4870	<u>1341aa</u>	Protein coding	CCDS84336	Q8CG19	TSL:5 GENCODE basic APPRIS P1
Ltbp1-210	ENSMUST00000234490.1	6235	<u>1713aa</u>	Protein coding	102	-	GENCODE basic
Ltbp1-205	ENSMUST00000135447.7	3114	980aa	Protein coding		B1B1E2	CDS 3' incomplete TSL:1
Ltbp1-209	ENSMUST00000234327.1	246	82aa	Protein coding	19-		5' and 3' truncations in transcript evidence prevent annotation of the start and the end of the CDS. CDS 5' and 3' incomplete
Ltbp1-206	ENSMUST00000135750.1	3277	No protein	Retained intron	¥4	2	TSL:1
Ltbp1-204	ENSMUST00000127458.1	1905	No protein	Retained intron	12	-	TSL:1
Ltbp1-208	ENSMUST00000234008.1	823	No protein	IncRNA			
Ltbp1-207	ENSMUST00000146839.1	519	No protein	IncRNA			TSL:2

The strategy is based on the design of *Ltbp1-201* transcript, The transcription is shown below

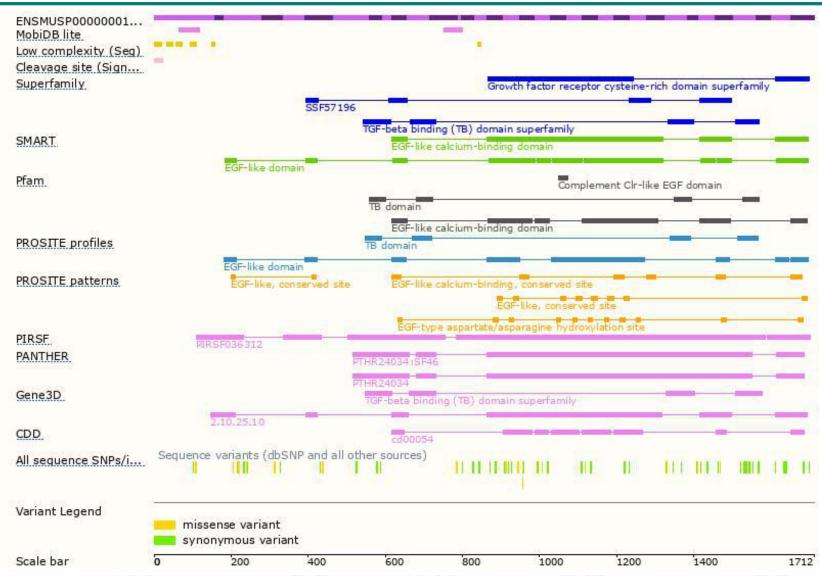


Genomic location distribution



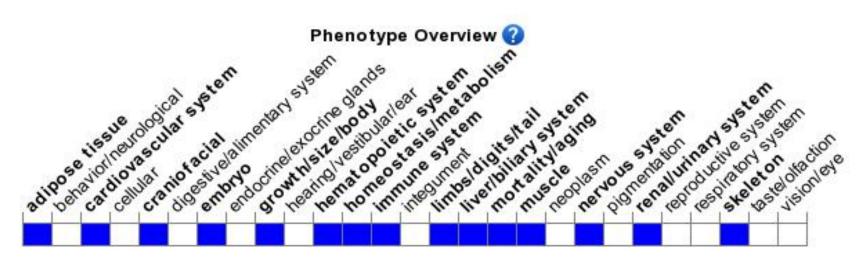
Protein domain





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 null allele exhibit embryonic and neonatal lethality associated with defects in the aortic arch and outflow tract.



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





