

Letm1 Cas9-CKO Strategy

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Project Overview

Project Name

Letm1

Project type

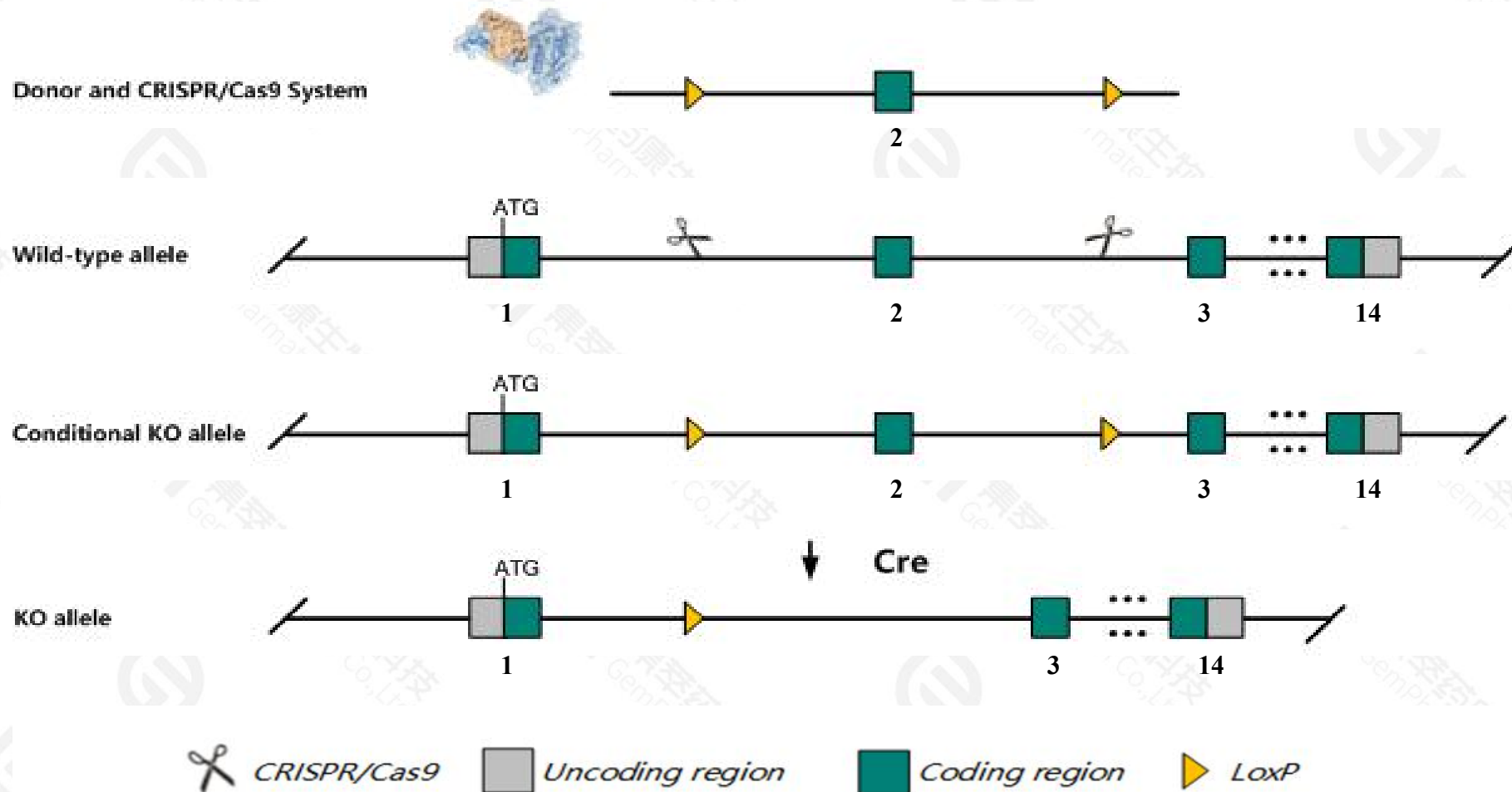
Cas9-CKO

Strain background

C57BL/6JGpt

Conditional Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Letm1* gene. The schematic diagram is as follows:



- The *Letm1* gene has 6 transcripts. According to the structure of *Letm1* gene, exon2 of *Letm1*-201(ENSMUST00000005431.5) transcript is recommended as the knockout region. The region contains 61bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Letm1* 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.

- According to the existing MGI data, homozygous deletion of this gene causes embryonic lethality prior to E6.5 while ~50% of heterozygotes die before E13.5. Surviving heterozygous mice show altered glucose metabolism, impaired control of brain ATP levels, and increased susceptibility to kainic acid-induced seizures.
- The *Letm1* gene is located on the Chr5. 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)

Letm1 leucine zipper-EF-hand containing transmembrane protein 1 [Mus musculus (house mouse)]

Gene ID: 56384, updated on 13-Mar-2020

Summary

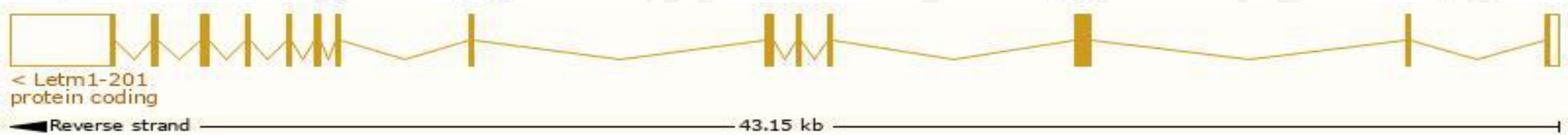
Official Symbol	Letm1 provided by MGI
Official Full Name	leucine zipper-EF-hand containing transmembrane protein 1 provided by MGI
Primary source	MGI:MGI:1932557
See related	Ensembl:ENSMUSG00000005299
Gene type	protein coding
RefSeq status	VALIDATED
Organism	Mus musculus
Lineage	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
Expression	Ubiquitous expression in duodenum adult (RPKM 18.7), colon adult (RPKM 16.5) and 28 other tissues See more
Orthologs	human all

Transcript information (Ensembl)

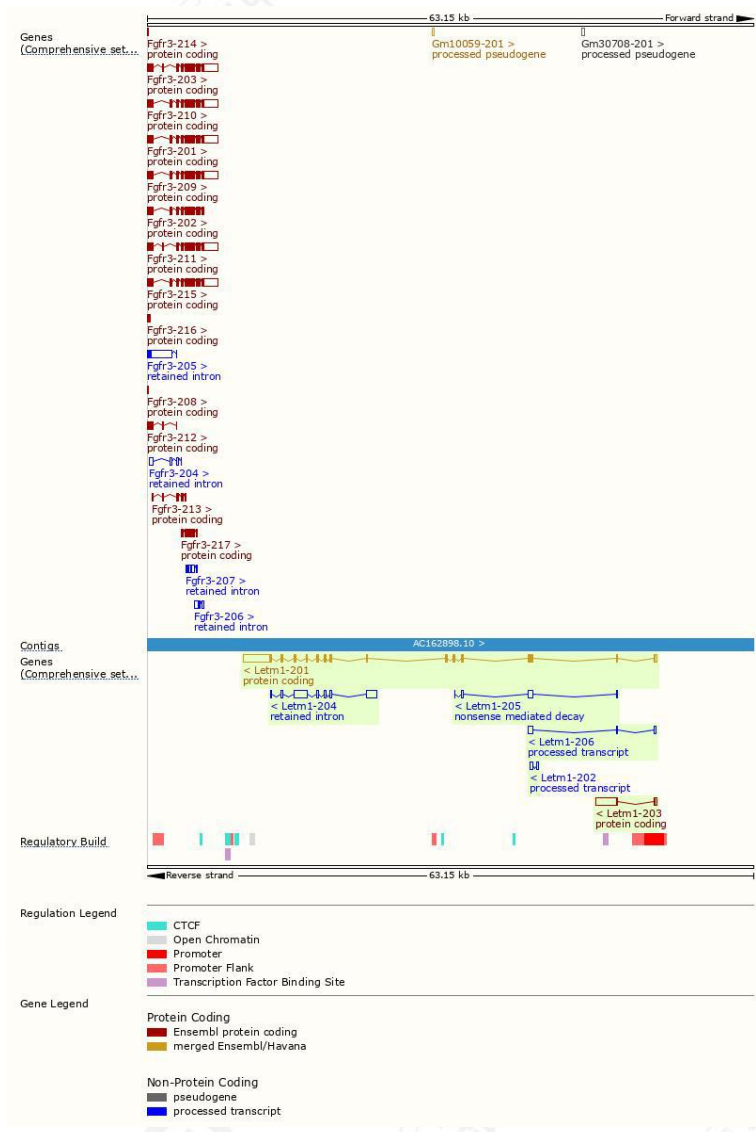
The gene has 6 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Letm1-201	ENSMUST00000005431.5	5272	738aa	Protein coding	CCDS19207	Q9Z2I0	TSL:1 GENCODE basic APPRIS P1
Letm1-203	ENSMUST00000148451.1	2478	47aa	Protein coding	-	A0A0J9YTQ9	TSL:2 GENCODE basic
Letm1-205	ENSMUST00000200827.1	642	13aa	Nonsense mediated decay	-	A0A0J9YV52	CDS 5' incomplete TSL:5
Letm1-206	ENSMUST00000201981.1	747	No protein	Processed transcript	-	-	TSL:3
Letm1-202	ENSMUST00000144071.1	488	No protein	Processed transcript	-	-	TSL:2
Letm1-204	ENSMUST00000149886.1	3056	No protein	Retained intron	-	-	TSL:1

The strategy is based on the design of *Letm1-201* transcript,the transcription is shown below:



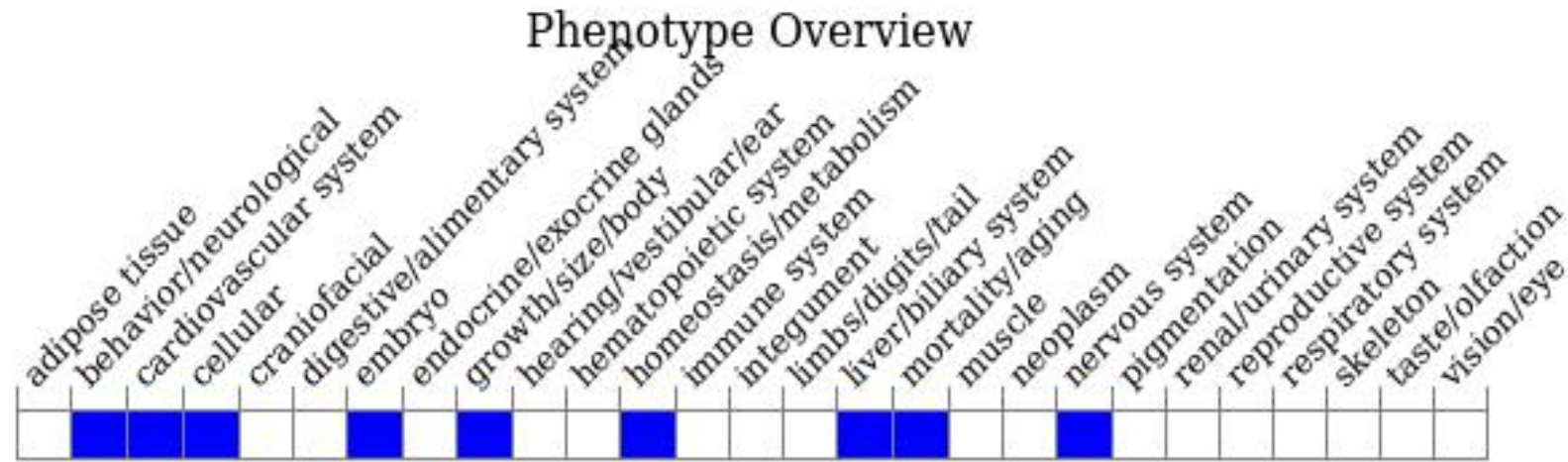
Genomic location distribution



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, homozygous deletion of this gene causes embryonic lethality prior to E6.5 while ~50% of heterozygotes die before E13.5. Surviving heterozygous mice show altered glucose metabolism, impaired control of brain ATP levels, and increased susceptibility to kainic acid-induced seizures.

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