

Ldb1 Cas9-CKO Strategy

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Design Date: 2019-10-23

Project Overview



Project Name

Ldb1

Project type

Cas9-CKO

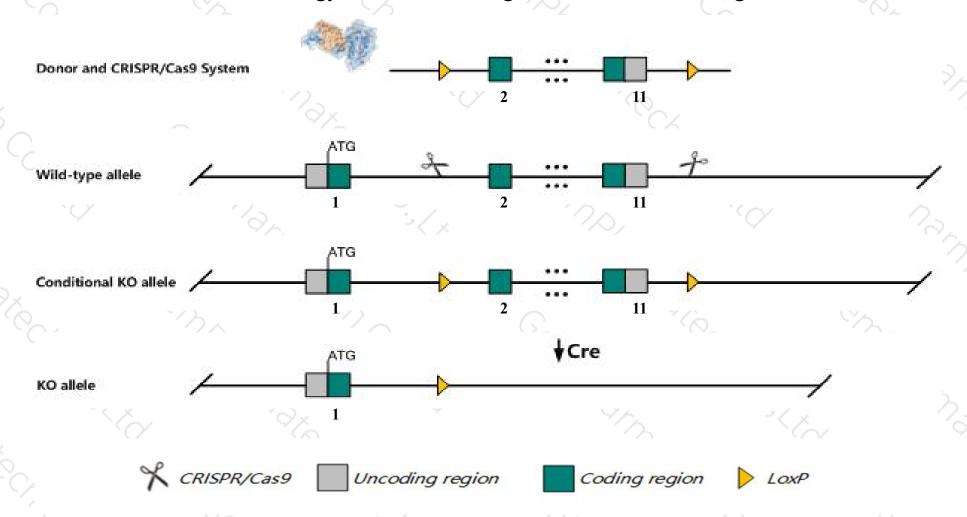
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Ldb1* gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Ldb1* gene has 8 transcripts. According to the structure of *Ldb1* gene, exon2-exon11 of *Ldb1-207*(ENSMUST00000156585.8) transcript is recommended as the knockout region. The region contains 1211bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Ldb1* 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 disruptions in this gene die as embryos at E9.5-E10 with impaired primitive erythropoiesis and vascular development.
- > The *Ldb1* 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)



Ldb1 LIM domain binding 1 [*Mus musculus* (house mouse)]

Gene ID: 16825, updated on 14-Oct-2019

Summary

☆ ?

Official Symbol Ldb1 provided by MGI

Official Full Name LIM domain binding 1 provided by MGI

Primary source MGI:MGI:894762

See related Ensembl:ENSMUSG00000025223

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 NLI; CLIM2

Expression Ubiquitous expression in thymus adult (RPKM 131.0), CNS E14 (RPKM 82.2) and 27 other tissues See more

Orthologs <u>human</u> all

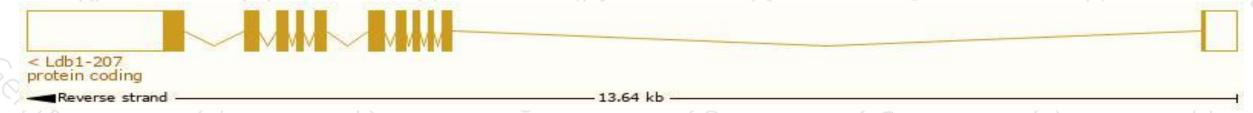
Transcript information (Ensembl)



The gene has 8 transcripts, all transcripts are shown below:

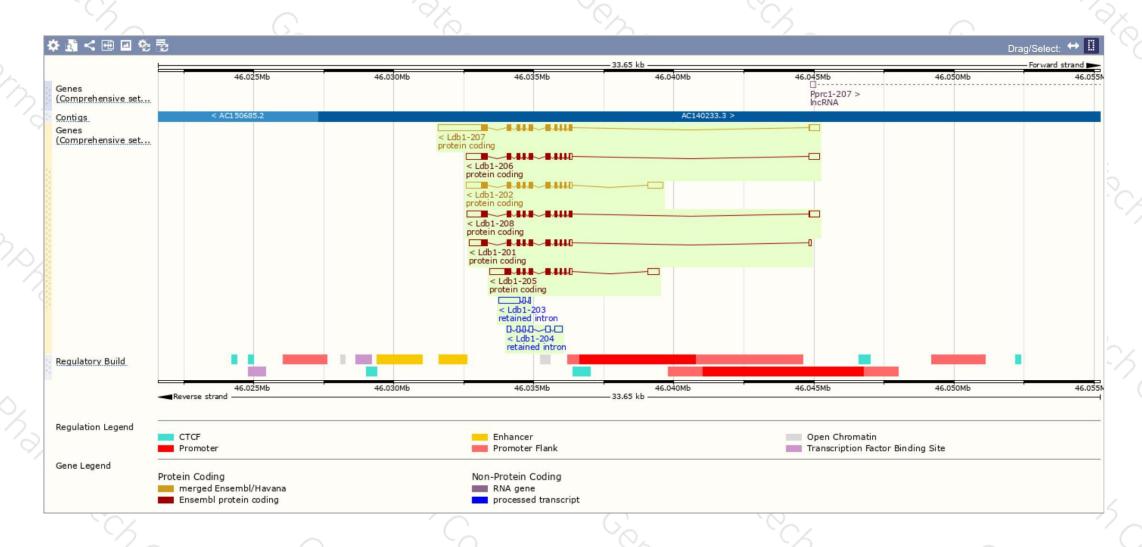
Show/hide columns (1 hidden)								Filter	1 ×
Name 🝦	Transcript ID ▼	bp 🍦	Protein 🍦	Biotype	CCDS 🍦	UniProt 🌲	Flags -		
Ldb1-208	ENSMUST00000185355.6	2088	411aa	Protein coding	<u>CCDS50455</u> &	P70662 &	TSL:1	GENCODE basic	APPRIS ALT1
Ldb1-207	ENSMUST00000156585.8	3133	<u>411aa</u>	Protein coding	<u>CCDS50455</u> &	P70662 &	TSL:1	GENCODE basic	APPRIS ALT1
Ldb1-206	ENSMUST00000152946.7	2108	373aa	Protein coding	-	D3Z1C5 _日	TSL:5	GENCODE basic	APPRIS ALT1
Ldb1-205	ENSMUST00000137771.1	2014	319aa	Protein coding	-	<u>P70662</u> 굡	TSL:5 GENCODE basic		
Ldb1-204	ENSMUST00000136203.1	900	No protein	Retained intron	8=	-	TSL:2		
Ldb1-203	ENSMUST00000126320.7	907	No protein	Retained intron	15	150	TSL:2		
Ldb1-202	ENSMUST00000056931.13	2295	<u>375aa</u>	Protein coding	CCDS29870 ₺	P70662 &	TSL:1	GENCODE basic	APPRIS P3
Ldb1-201	ENSMUST00000026252.13	1718	<u>375aa</u>	Protein coding	CCDS29870 &	P70662 母	TSL:1	GENCODE basic	APPRIS P3

The strategy is based on the design of *Ldb1-207* transcript, The transcription is shown below



Genomic location distribution





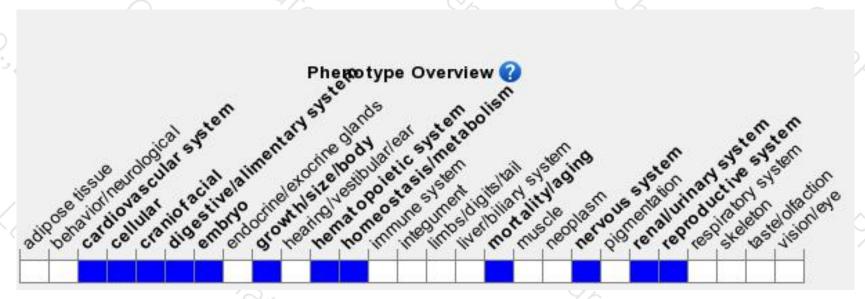
Protein domain





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 disruptions in this gene die as embryos at E9.5-E10 with impaired primitive erythropoiesis and vascular development.



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





