Cd300lf Cas9-CKO Strategy

Designer:

Date:

Zengyang 2019-7-30

Project Overview



Project Name

Cd300lf

Project type

Cas9-CKO

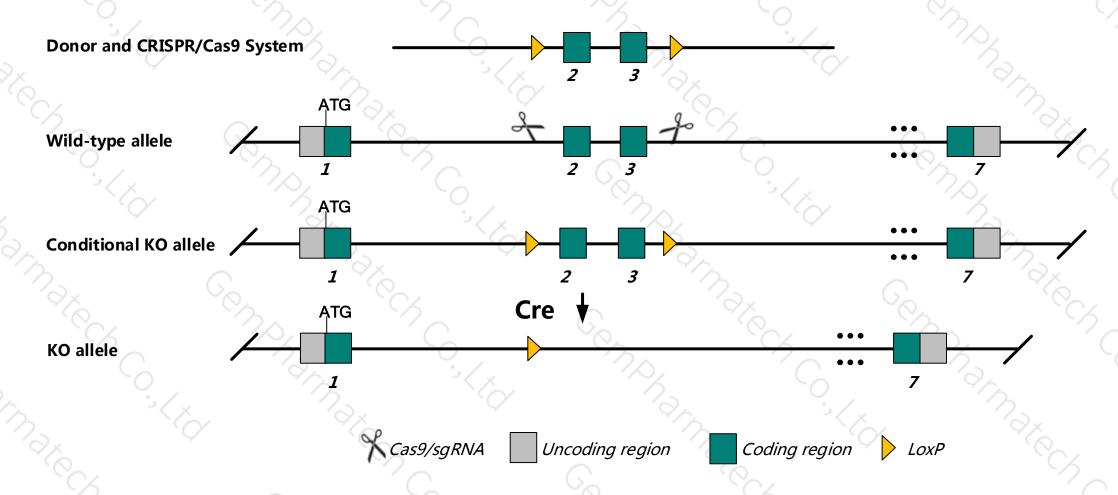
Strain background

C57BL/6J

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Cd300lf* gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Cd300lf* gene has 7 transcripts. According to the structure of *Cd300lf* gene, exon2-3 of *Cd300lf-202* (ENSMUST00000106561.7) transcript is recommended as the knockout region. The region contains 472bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Cd300lf* gene. The brief process is as follows: sgRNA was transcribed in vitro, donor vector was constructed.Cas9, sgRNA and Donor were microinjected into the fertilized eggs of C57BL/6J 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/6J mice.
- The flox mice was knocked out after mating with mice expressing Cre recombinase, resulting in the loss of function of the target gene in specific tissues or cell types.

Notice



According to the existing MGI data, mice homozygous for a knock-out allele exhibit increased severity of experimental autoimmune encephalomyelitis with increased demyelination.

➤ The KO region contains functional region of the *Rab37* gene. Knockout the region may affect the function of *Rab37* gene.

➤ The *Cd300lf* gene is located on the Chr11. 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.

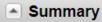
This strategy is designed based on genetic information in existing databases. Due to the complexity of biological processes, all risk of the gene knockout on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.

Gene information (NCBI)



Cd300lf CD300 molecule like family member F [Mus musculus (house mouse)]

Gene ID: 246746, updated on 4-Jun-2019





Official Symbol Cd300lf provided by MGI

Official Full Name CD300 molecule like family member F provided by MGI

Primary source MGI:MGI:2442359

See related Ensembl: ENSMUSG00000047798

Gene type protein coding
RefSeq status VALIDATED
Organism <u>Mus musculus</u>

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

Murinae; Mus; Mus

Also known as CLM1; CLIM1; CLM-1; Digr2; IREM1; LMIR3; Pigr3; IgSF13; F730004D16Rik

Expression Biased expression in spleen adult (RPKM 7.7), liver E18 (RPKM 5.1) and 12 other tissues See more

Orthologs human all

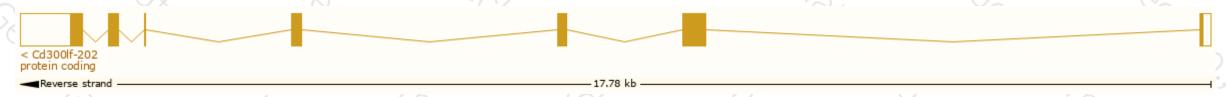
Transcript information (Ensembl)



The gene has 7 transcripts, and all transcripts are shown below: :

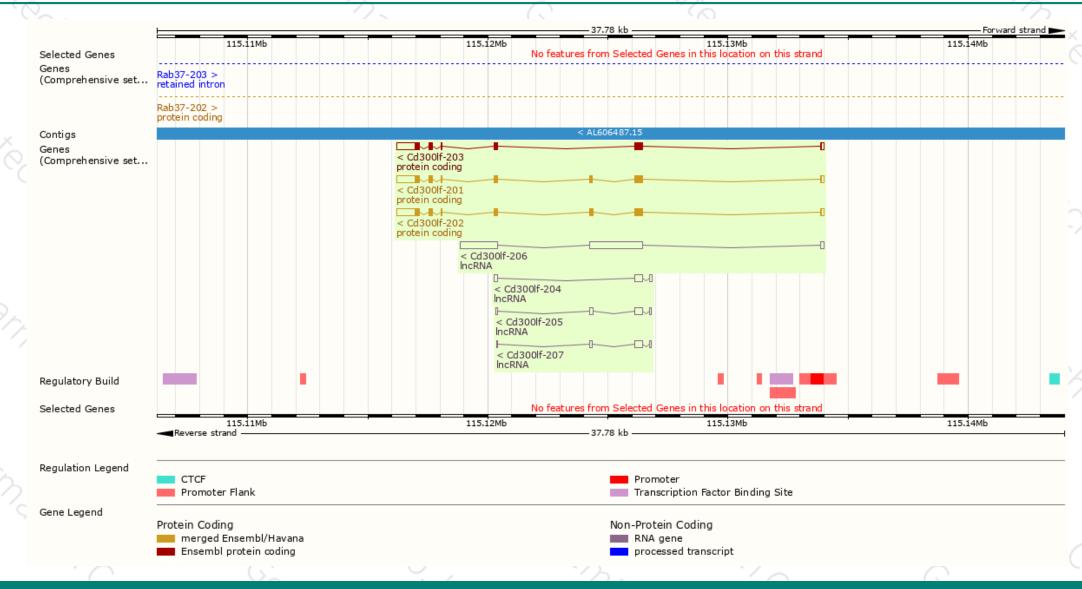
	Name ≜	Transcript ID	bp 🛦	Protein A	Biotype 🌢	CCDS A	UniProt ▼	Flags
-		· · ·			· · · · · ·			
	Cd300lf-201	ENSMUST00000051264.13	1864	<u>330aa</u>	Protein coding	CCDS25619&	<u>Q6SJQ7</u> &	TSL:1 GENCODE basic APPRIS P3
	Cd300lf-203	ENSMUST00000106562.2	1762	<u>296aa</u>	Protein coding	CCDS83925₺	<u>A2A6Z2</u> ₽	TSL:1 GENCODE basic
	Cd300lf-202	ENSMUST00000106561.7	1885	<u>337aa</u>	Protein coding	CCDS48980 ₺	<u>A0A1C9ZQ14</u> @ <u>Q6SJQ7</u> @	TSL:1 GENCODE basic APPRIS ALT2
	Cd300lf-206	ENSMUST00000146254.1	3929	No protein	IncRNA ■	-	-	TSL:1
	Cd300lf-204	ENSMUST00000124083.7	615	No protein	IncRNA	-	-	TSL:3
	Cd300lf-205	ENSMUST00000127927.7	608	No protein	IncRNA	-	-	TSL:3
	Cd300lf-207	ENSMUST00000149335.1	569	No protein	IncRNA	-	-	TSL:3

The strategy is based on the design of *Cd300lf-202* transcript, The transcription is shown below



Genomic location (Ensembl)





Protein domain (Ensembl)

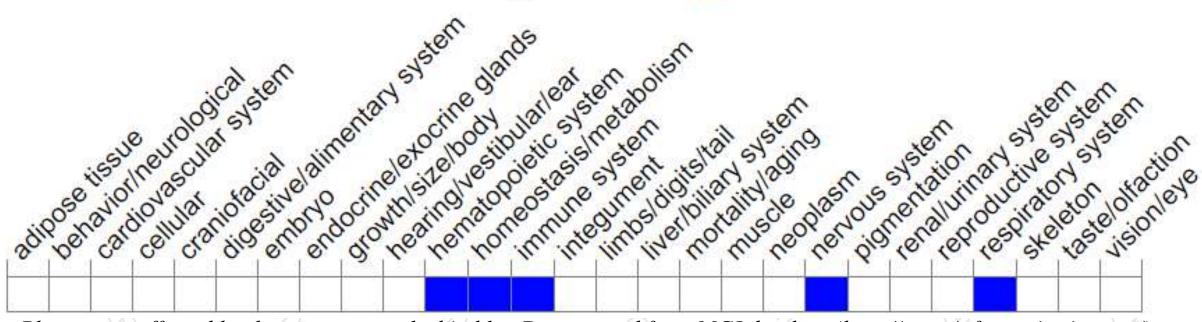




Mouse phenotype description(MGI)







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 knock-out allele exhibit increased severity of experimental autoimmune encephalomyelitis with increased demyelination.

If you have any questions, you are welcome to inquire. Tel: 025-5864 1534





