

Clcc1 Cas9-CKO Strategy

Designer:

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Design Date:

2019-7-24

Project Overview



Project Name

Clcc1

Project type

Cas9-CKO

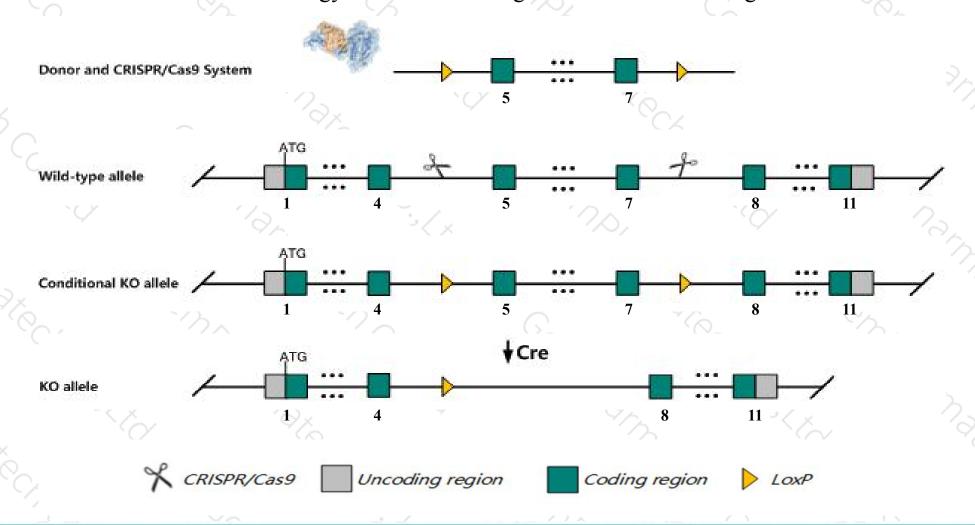
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Clcc1* gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Clcc1* gene has 8 transcripts. According to the structure of *Clcc1* gene, exon5-exon7 of *Clcc1-203*(ENSMUST00000106613.1) transcript is recommended as the knockout region. The region contains 457bp coding sequence.

 Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Clcc1* 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 spontaneous mutation show strain-dependent cerebellar granule cell death and peripheral motor axon degeneration. The peripheral neuropathy, neurogenic muscular atrophy and mild truncal ataxia observed on the C57BL/6JGpt background are not found on the C3H/HeSnJ background.
- > The *Clcc1* gene is located on the Chr3. 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)



Clcc1 chloride channel CLIC-like 1 [Mus musculus (house mouse)]

Gene ID: 229725, updated on 31-Jan-2019

Summary

↑ ?

Official Symbol Clcc1 provided by MGI

Official Full Name chloride channel CLIC-like 1 provided by MGI

Primary source MGI:MGI:2385186

See related Ensembl: ENSMUSG00000027884

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 McIc

Expression Ubiquitous expression in bladder adult (RPKM 9.9), limb E14.5 (RPKM 9.7) and 28 other tissuesSee more

Orthologs human all

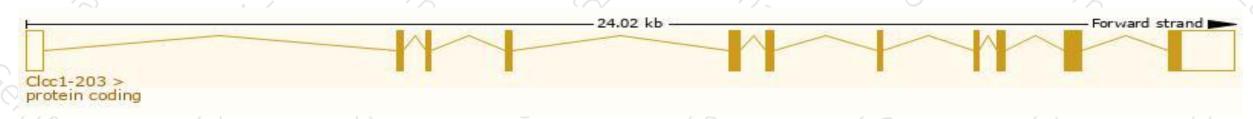
Transcript information (Ensembl)



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

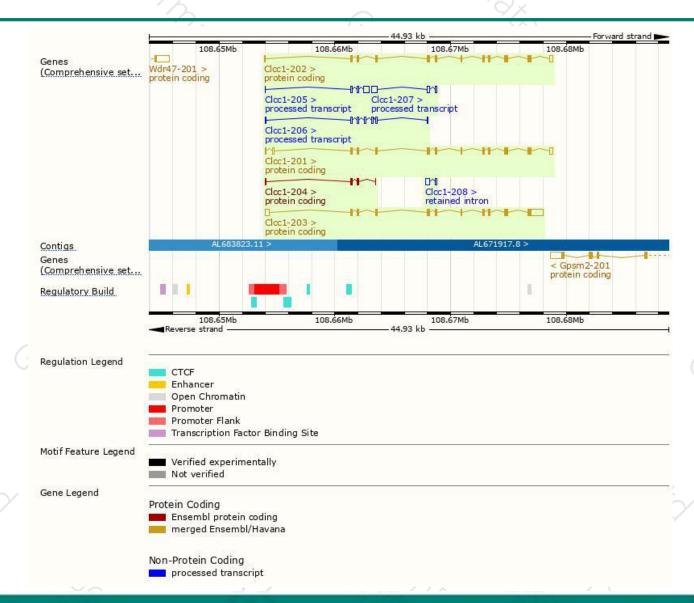
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Clcc1-203	ENSMUST00000106613.1	3064	544aa	Protein coding	CCDS51047	A2AEM2	TSL:1 GENCODE basic APPRIS ALT2
Clcc1-201	ENSMUST00000029483.14	2224	<u>539aa</u>	Protein coding	CCDS17766	Q99LI2	TSL:5 GENCODE basic APPRIS P3
Clcc1-202	ENSMUST00000106609.7	2043	<u>539aa</u>	Protein coding	CCDS17766	Q99LI2	TSL:1 GENCODE basic APPRIS P3
Clcc1-204	ENSMUST00000124384.7	358	<u>84aa</u>	Protein coding	1528	Z4YMG1	CDS 3' incomplete TSL:3
CIcc1-206	ENSMUST00000130352.3	895	No protein	Processed transcript	1.5	5/2	TSL:3
Clcc1-207	ENSMUST00000139016.2	795	No protein	Processed transcript	-	+0	TSL:3
Clcc1-205	ENSMUST00000125274.7	762	No protein	Processed transcript	827	20	TSL:3
Clcc1-208	ENSMUST00000156811.2	469	No protein	Retained intron	-	20	TSL:3

The strategy is based on the design of Clcc1-203 transcript, The transcription is shown below



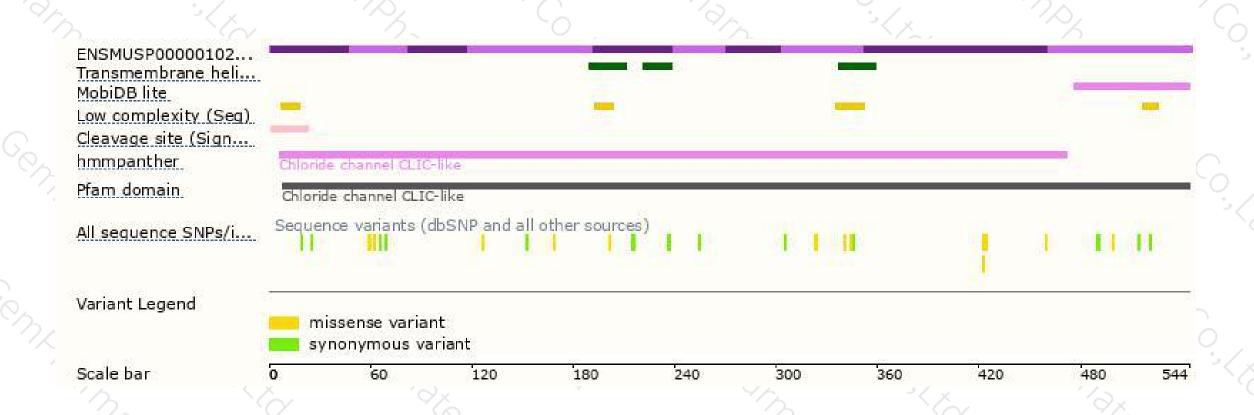
Genomic location distribution





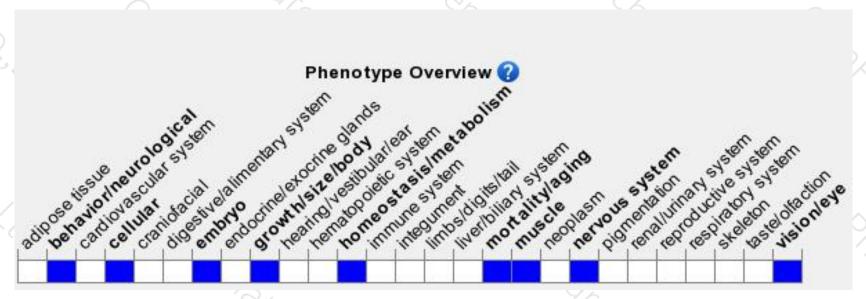
Protein domain





Mouse phenotype description(MGI)





Phenotypes affected by the gene are marked in blue.Data quoted from MGI database(http://www.informatics.jax.org/).

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





