

Exoc5 Cas9-CKO Strategy

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Project Overview



Project Name

Exoc5

Project type

Cas9-CKO

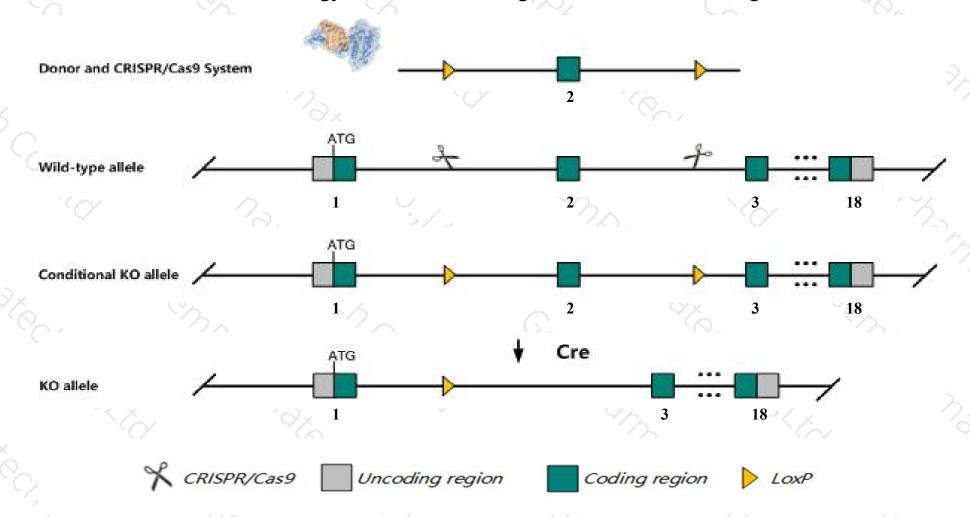
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Exoc5* gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Exoc5* gene has 7 transcripts. According to the structure of *Exoc5* gene, exon2 of *Exoc5*207(ENSMUST00000162175.8) transcript is recommended as the knockout region. The region contains 95bp coding sequence.

 Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Exoc5* 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 conditional allele activated in all cells die prior to E8.5. Mice homozygous for a conditional allele activated in kidney cells exhibit ureteropelvic junction obstructions leading to neontal death.
- > The *Exoc5* gene is located on the Chr14. 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)



Exoc5 exocyst complex component 5 [Mus musculus (house mouse)]

Gene ID: 105504, updated on 13-Mar-2020

Summary

☆ ?

Official Symbol Exoc5 provided by MGI

Official Full Name exocyst complex component 5 provided by MGI

Primary source MGI:MGI:2145645

See related Ensembl:ENSMUSG00000061244

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 Al447711, Al448003, Gm76, PRO1912, SEC10, Sec10l1

Expression Ubiquitous expression in CNS E11.5 (RPKM 11.7), CNS E14 (RPKM 11.3) and 28 other tissuesSee more

Orthologs <u>human</u> all

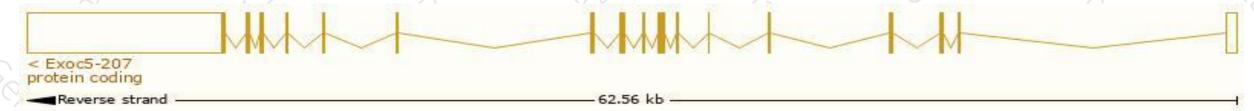
Transcript information (Ensembl)



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

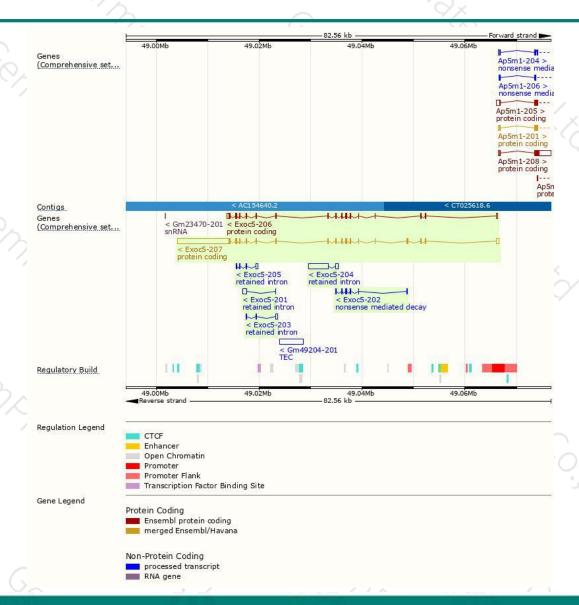
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Exoc5-207	ENSMUST00000162175.8	12699	708aa	Protein coding	CCDS49474	Q3TPX4	TSL:1 GENCODE basic APPRIS P1
Exoc5-206	ENSMUST00000161504.7	2582	643aa	Protein coding		E9PZ92	TSL:5 GENCODE basic
Exoc5-202	ENSMUST00000160386.2	694	49aa	Nonsense mediated decay	-	E0CYX2	CDS 5' incomplete TSL:3
Exoc5-204	ENSMUST00000160723.1	4429	No protein	Retained intron	70	-	TSL:1
Exoc5-201	ENSMUST00000159651.1	880	No protein	Retained intron	-		TSL:3
Exoc5-205	ENSMUST00000160833.1	749	No protein	Retained intron		-	TSL:5
Exoc5-203	ENSMUST00000160453.1	559	No protein	Retained intron	-		TSL:3
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The strategy is based on the design of *Exoc5-207* transcript, the transcription is shown below:



Genomic location distribution





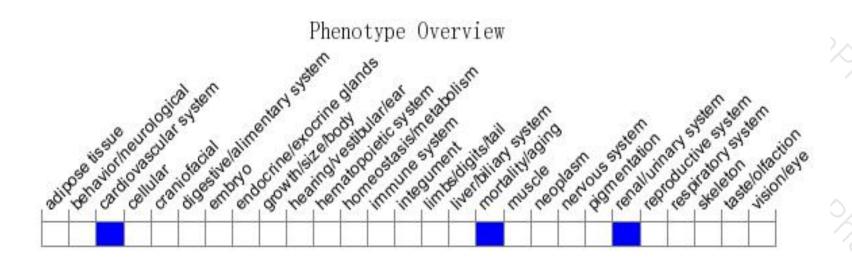
Protein domain





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 a conditional allele activated in all cells die prior to E8.5. Mice homozygous for a conditional allele activated in kidney cells exhibit ureteropelvic junction obstructions leading to neontal death.



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





