

Cwc27 Cas9-CKO Strategy

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Project Overview



Project Name

Cwc27

Project type

Cas9-CKO

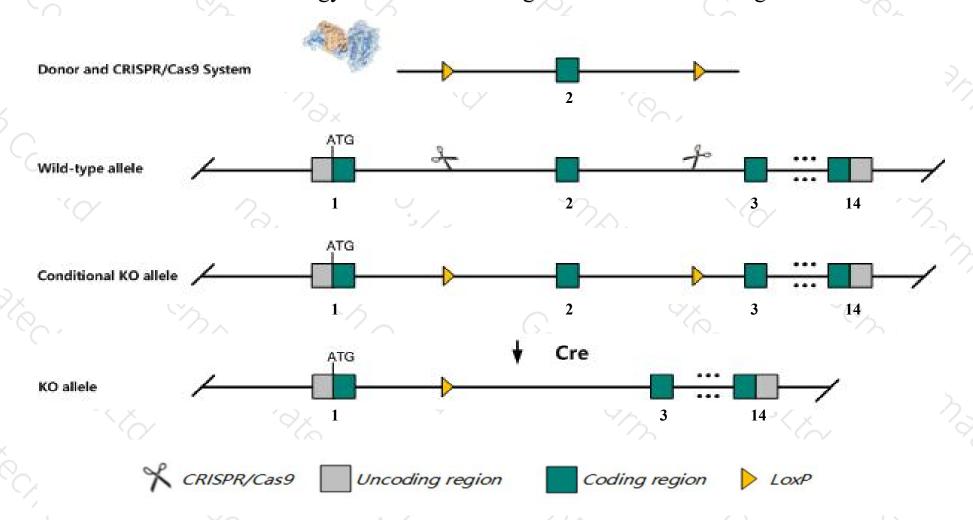
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Cwc27 gene. The schematic diagram is as follows:



Technical routes



- The Cwc27 gene has 4 transcripts. According to the structure of Cwc27 gene, exon2 of Cwc27-201 (ENSMUST00000022228.12) transcript is recommended as the knockout region. The region contains 97bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Cwc27* 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, homozygous mutant mice exhibit reduced viability. Surviors after birth show signs of growth retardation and retinal depigmentation, along with numerous neurological, immunological, and blood chemistry abnormalities.
- > The partial intron of transcript *Srek1ip1*-203 will be deleted together in this strategy.
- The floxed region is near to the N-terminal of *Srek1ip1* gene, this strategy may influence the regulatory function of the N-terminal of *Srek1ip1* gene.
- > Transcript Cwc27-202&203 may not be affected.
- The Cwc27 gene is located on the Chr13. 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)



Cwc27 CWC27 spliceosome-associated protein [Mus musculus (house mouse)]

Gene ID: 67285, updated on 13-Mar-2020

Summary

☆ ?

Official Symbol Cwc27 provided by MGI

Official Full Name CWC27 spliceosome-associated protein provided by MGI

Primary source MGI:MGI:1914535

See related Ensembl:ENSMUSG00000021715

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 3110009E13Rik, NY-CO-10, Sdccag10

Expression Broad expression in CNS E11.5 (RPKM 1.3), CNS E14 (RPKM 0.7) and 21 other tissuesSee more

Orthologs <u>human</u> all

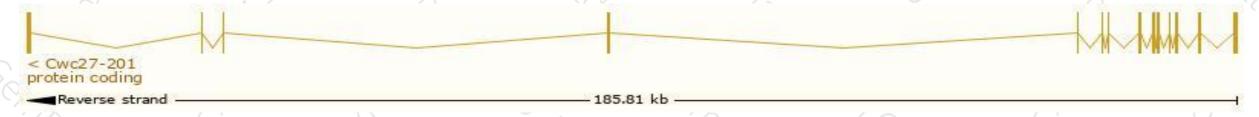
Transcript information (Ensembl)



The gene has 4 transcripts, all transcripts are shown below:

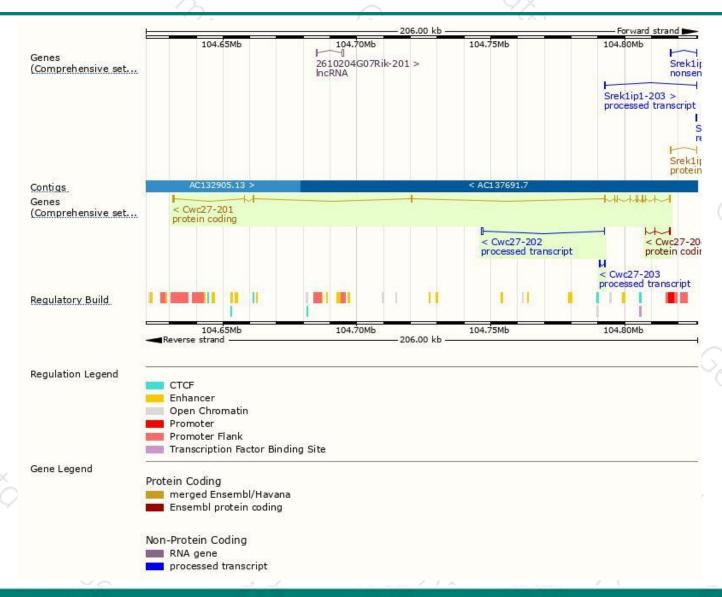
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Cwc27-201	ENSMUST00000022228.12	2071	469aa	Protein coding	CCDS26752	Q3TKY6	TSL:1 GENCODE basic APPRIS is a system to annotate alternatively spliced transcripts based on a range of computational methods to identify the most functionally important transcript(s) of a gene. APPRIS P
Cwc27-204	ENSMUST00000154165.1	353	83aa	Protein coding	0.0	D3Z2U3	CDS 3' incomplete TSL:3
Cwc27-202	ENSMUST00000141495.1	588	No protein	Processed transcript	8-8	-	TSL:3
Cwc27-203	ENSMUST00000147514.1	451	No protein	Processed transcript	12		TSL:3

The strategy is based on the design of Cwc27-201 transcript, The transcription is shown below



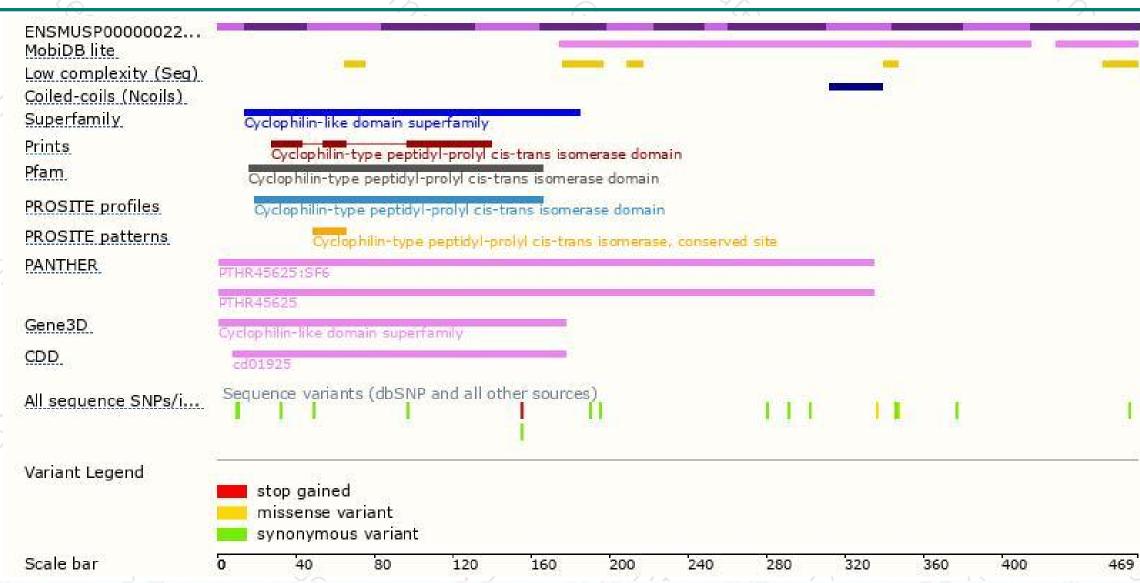
Genomic location distribution





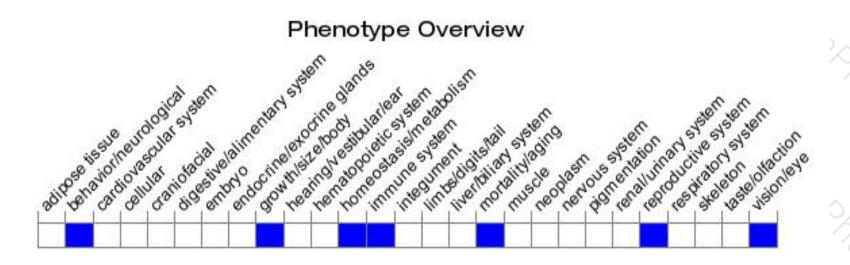
Protein domain





Mouse phenotype description(MGI)





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

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





