

Kcp Cas9-CKO Strategy

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Project Overview



Project Name Kcp

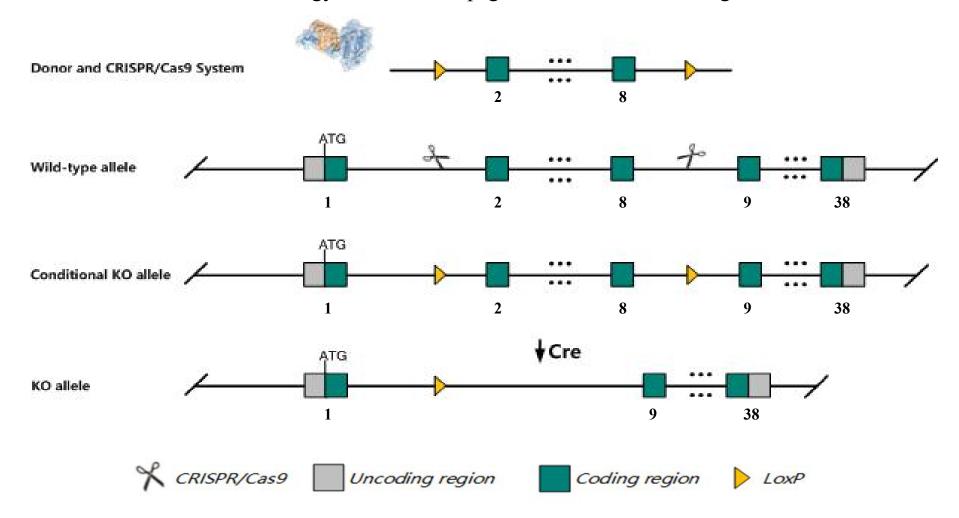
Project type Cas9-CKO

Strain background C57BL/6JGpt

Conditional Knockout strategy



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



Technical routes



The *Kcp* gene has 10 transcripts. According to the structure of *Kcp* gene, exon2-exon8 of *Kcp-203* (ENSMUST00000101614.9) transcript is recommended as the knockout region. The region contains 773bp coding sequence. Knock out the region will result in disruption of protein function.

In this project we use CRISPR/Cas9 technology to modify *Kcp* 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 null mice display increased sensitivity to renal injury.

Kcp-205 will not be affected.

The *Kcp* gene is located on the Chr6. 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



Kcp kielin/chordin-like protein [Mus musculus (house mouse)]

Gene ID: 333088, updated on 13-Mar-2020

Summary

☆ ?

Official Symbol Kcp provided by MGI

Official Full Name kielin/chordin-like protein provided by MGI

Primary source MGI:MGI:2141640

See related Ensembl:ENSMUSG00000059022

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 AW060220, CRIM-2, Crim2, Gm793, KCP-1

Expression Broad expression in limb E14.5 (RPKM 11.6), ovary adult (RPKM 5.1) and 16 other tissuesSee more

Orthologs <u>human all</u>

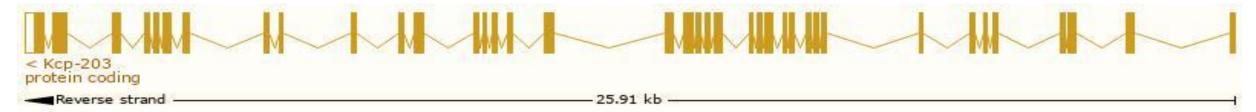
Transcript information Ensembl



The gene has 10 transcripts, all transcripts are shown below:

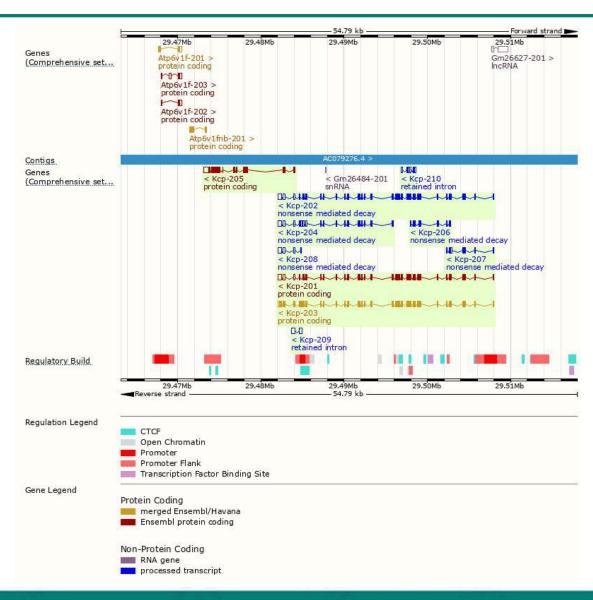
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Kcp-203	ENSMUST00000101614.9	4885	<u>1550aa</u>	Protein coding	CCDS57415	Q3U492	TSL:1 GENCODE basic APPRIS P1
Kcp-201	ENSMUST00000078112.12	4775	<u>1255aa</u>	Protein coding	353	Q3U492	TSL:5 GENCODE basic
Кср-205	ENSMUST00000159482.8	2742	728aa	Protein coding	127	13 5 0	CDS 5' incomplete TSL:1
Kcp-202	ENSMUST00000091391.10	4758	<u>1254aa</u>	Nonsense mediated decay	758	E9QNB3	TSL:2
Kcp-204	ENSMUST00000159479.7	2676	<u>359aa</u>	Nonsense mediated decay	-	F6UH68	CDS 5' incomplete TSL:1
Kcp-208	ENSMUST00000161276.7	971	<u>19aa</u>	Nonsense mediated decay	100	F7DED4	CDS 5' incomplete TSL:1
Kcp-207	ENSMUST00000161237.1	845	<u>165aa</u>	Nonsense mediated decay	120	E0CYL0	TSL:5
Kcp-206	ENSMUST00000160181.1	802	91aa	Nonsense mediated decay	725	F7B5X1	CDS 5' incomplete TSL:3
Kcp-209	ENSMUST00000161655.1	760	No protein	Retained intron	1.5	(5)	TSL:3
Kcp-210	ENSMUST00000162959.7	747	No protein	Retained intron	-	690	TSL:3

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



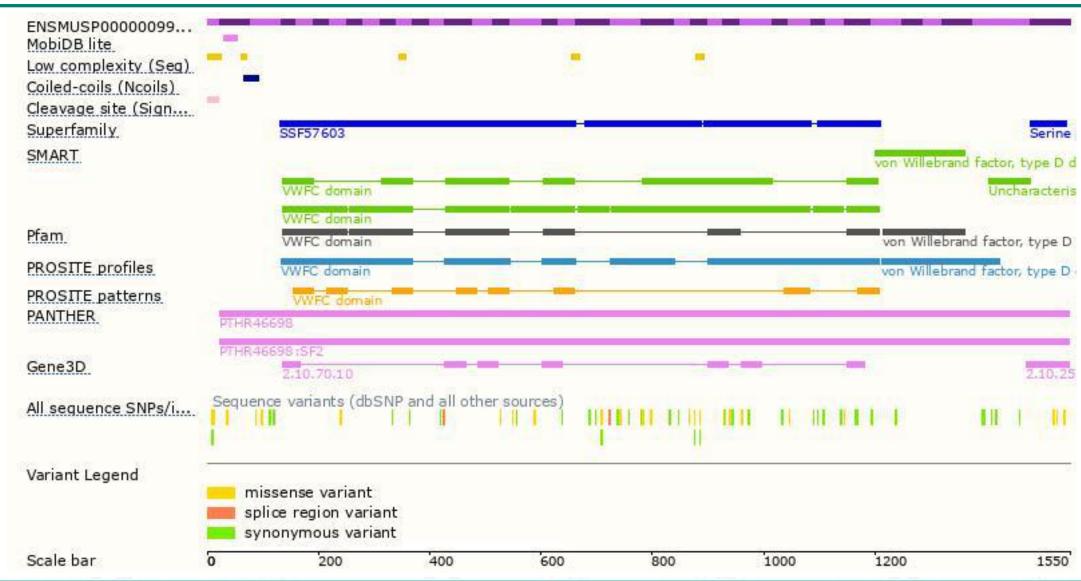
Genomic location distribution





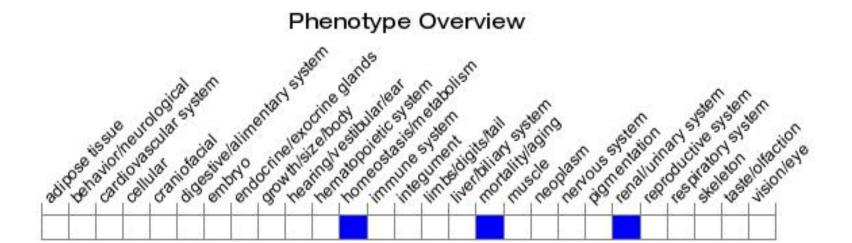
Protein domain





Mouse phenotype description(MGI)





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

According to the existing MGI data, Homozygous null mice display increased sensitivity to renal injury.



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





