

# Kcnq5 Cas9-CKO Strategy

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**Reviewer:** Xueting Zhang

**Design Date:** 2019-10-28

## **Project Overview**



**Project Name** 

Kcnq5

**Project type** 

Cas9-CKO

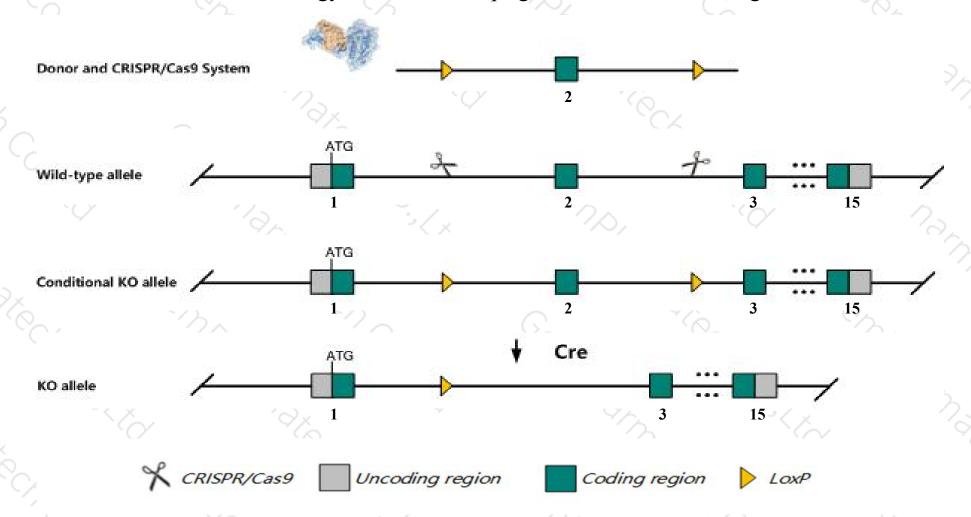
Strain background

C57BL/6JGpt

## Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Kcnq5* gene. The schematic diagram is as follows:



### Technical routes



- ➤ The *Kcnq5* gene has 7 transcripts. According to the structure of *Kcnq5* gene, exon2 of *Kcnq5-203*(ENSMUST00000115300.7) transcript is recommended as the knockout region. The region contains 91bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Kcnq5* 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 dominant negative knock-in mutation in this gene exhibit partial prenatal lethality and abnormal afterhyperpolarization in the in the CA3 area of hippocampus.
- > The *Kcnq5* gene is located on the Chr1. 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)



#### Kcnq5 potassium voltage-gated channel, subfamily Q, member 5 [ Mus musculus (house mouse) ]

Gene ID: 226922, updated on 12-Aug-2019

#### Summary



Official Symbol Kcnq5 provided by MGI

Official Full Name potassium voltage-gated channel, subfamily Q, member 5 provided by MGI

Primary source MGI:MGI:1924937

See related Ensembl: ENSMUSG00000028033

Gene type protein coding
RefSeq status VALIDATED
Organism Mus musculus

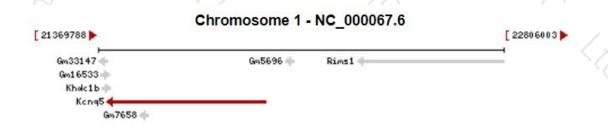
**Lineage** Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha;

Muroidea; Muridae; Murinae; Mus; Mus

Also known as D1Mgi1; AA589396; 7730402H11; 9230107O05Rik

Expression Biased expression in cortex adult (RPKM 3.9), frontal lobe adult (RPKM 3.2) and 9 other tissues See more

Orthologs human all



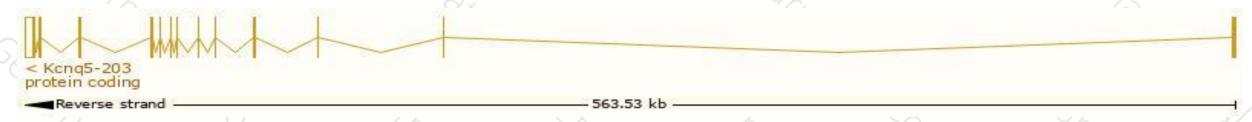
## Transcript information (Ensembl)



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

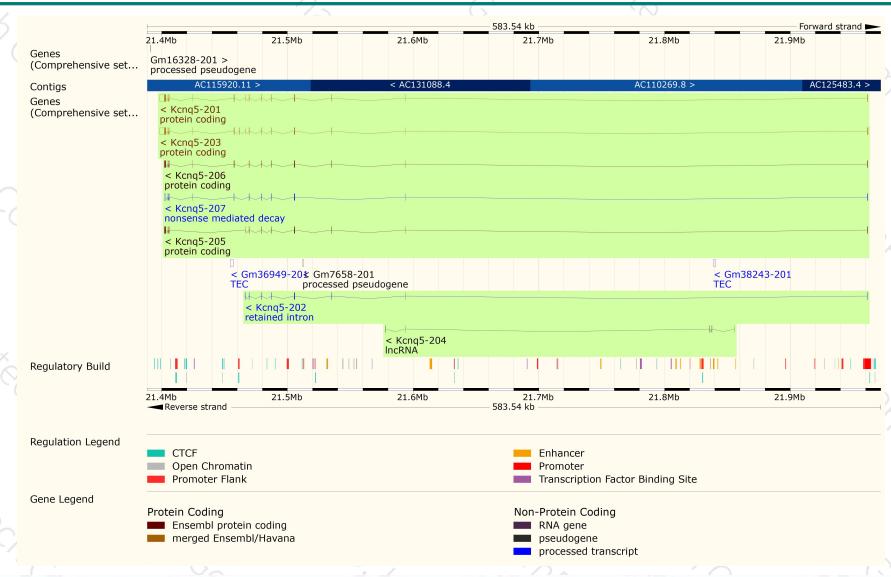
Name	Transcript ID	bp 🛊	Protein	Translation ID	Biotype	CCDS .	UniProt	Flags
Kcnq5-203	ENSMUST00000115300.7	6975	952aa	ENSMUSP00000110955.1	Protein coding	CCDS48230₺	E9Q9F1@	TSL:1 GENCODE basic APPRIS ALT2
Kcnq5-201	ENSMUST00000029667.12	6562	<u>933aa</u>	ENSMUSP00000029667.6	Protein coding	CCDS35525 ₺	Q9JK45₺₽	TSL:1 GENCODE basic APPRIS P3
Kcnq5-206	ENSMUST00000173404.1	2775	924aa	ENSMUSP00000134076.1	Protein coding	CCDS78554₺	G3UYG5₽	TSL:5 GENCODE basic APPRIS ALT2
Kcnq5-205	ENSMUST00000173058.7	2472	<u>823aa</u>	ENSMUSP00000134166.1	Protein coding	-	G3UYP4₽	TSL:5 GENCODE basic
Kcnq5-207	ENSMUST00000174183.7	2648	<u>172aa</u>	ENSMUSP00000134389.1	Nonsense mediated decay	15	G3UZ84₽	TSL:5
Kcnq5-202	ENSMUST00000115299.7	1472	No protein	140	Retained intron	-	= =	TSL:5
Kcnq5-204	ENSMUST00000134505.1	662	No protein	17	IncRNA	9.73	8	TSL:3

The strategy is based on the design of *Kcnq5-203* transcript, The transcription is shown below



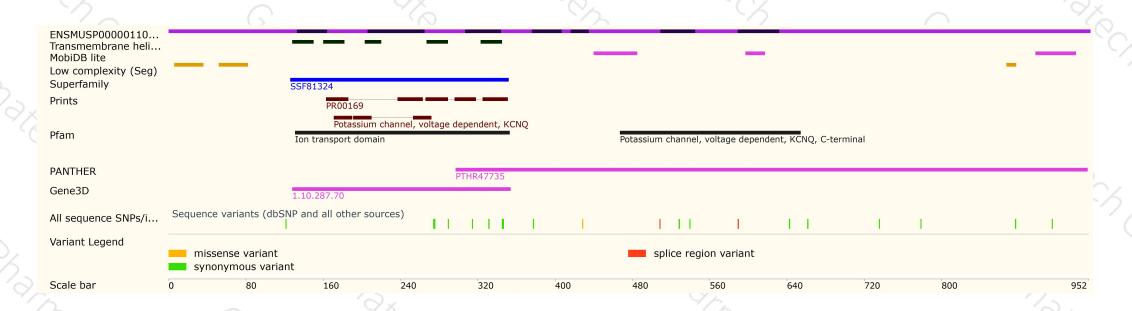
### Genomic location distribution





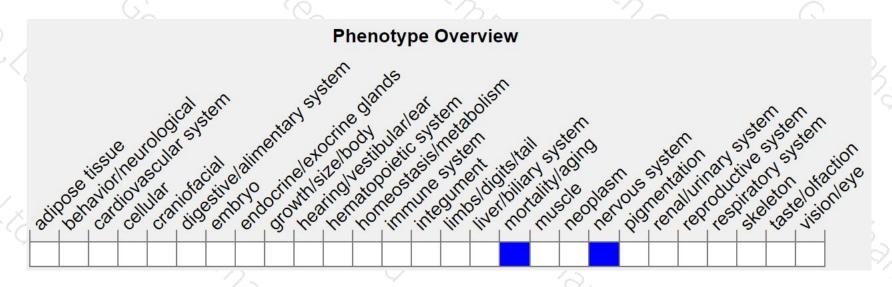
## Protein domain





## 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, Mice homozygous for a dominant negative knock-in mutation in this gene exhibit partial prenatal lethality and abnormal afterhyperpolarization in the in the CA3 area of hippocampus.



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





