

Kcnq5 Cas9-KO Strategy

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Reviewer: Xueting Zhang

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Project Overview



Project Name Kcnq5

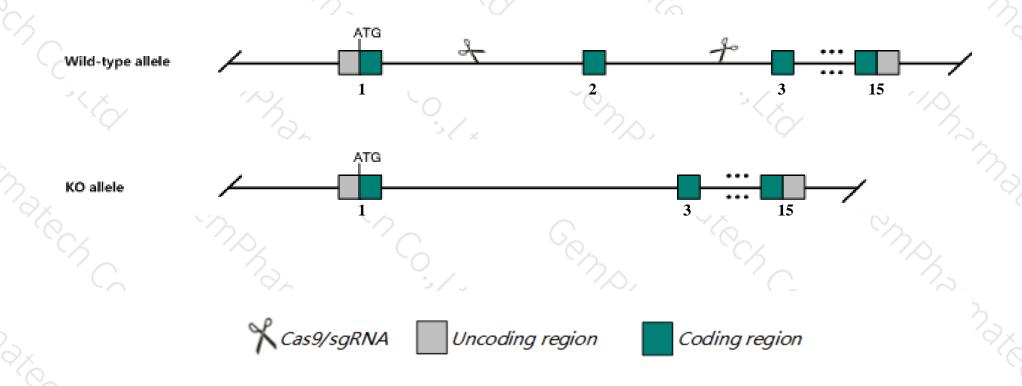
Project type Cas9-KO

Strain background C57BL/6JGpt

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: sgRNA was transcribed in vitro.Cas9 and sgRNA 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.

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 the gene knockout on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology 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

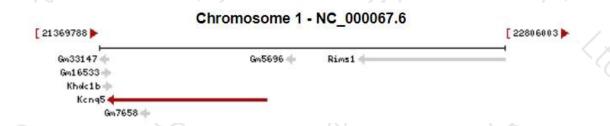
Lineage Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha;

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



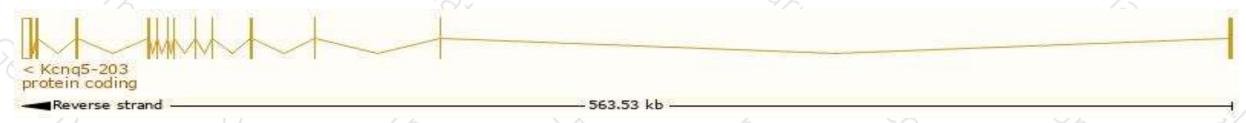
Transcript information (Ensembl)



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

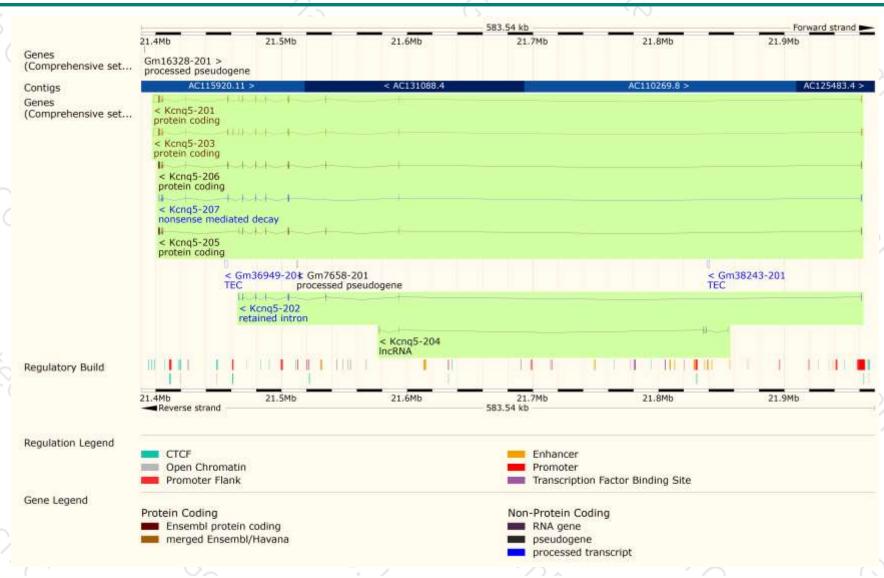
| Name 🍦 | Transcript ID 🗼 | bp 🌲 | Protein 🍦 | Translation ID 🗼 | Biotype | CCDS | UniProt | Flags 🝦 |
|-----------|-----------------------|------|--------------|----------------------|-------------------------|-------------|-----------------|---------------------------------|
| Kcnq5-203 | ENSMUST00000115300.7 | 6975 | <u>952aa</u> | 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 | <u>924aa</u> | 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 | - | <u>G3UZ84</u> ₽ | TSL:5 |
| Kcnq5-202 | ENSMUST00000115299.7 | 1472 | No protein | - | Retained intron | - | - | TSL:5 |
| Kcnq5-204 | ENSMUST00000134505.1 | 662 | No protein | - | IncRNA | - | - | TSL:3 |

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



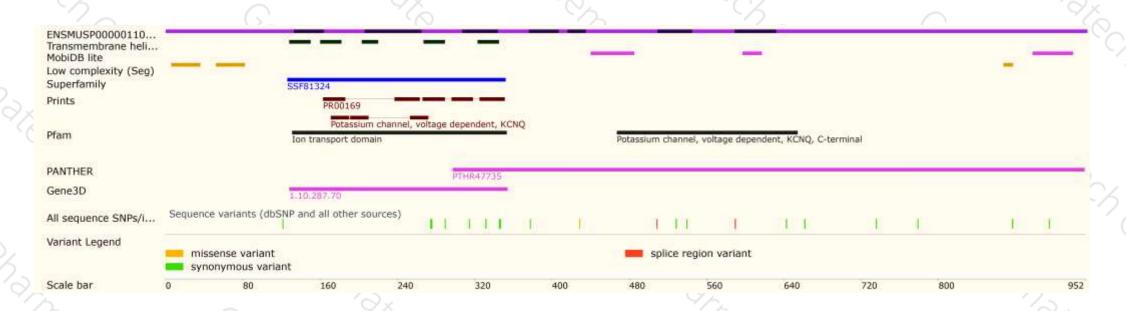
Genomic location distribution





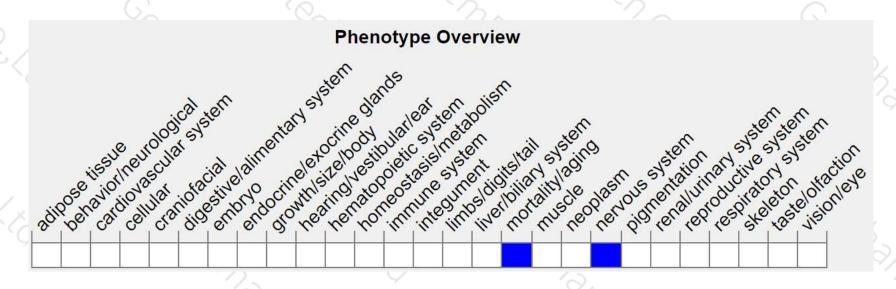
Protein domain





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.

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