

# *Kcnq1* Cas9-KO Strategy

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# Project Overview

**Project Name**

***Kcnq1***

**Project type**

**Cas9-KO**

**Strain background**

**C57BL/6JGpt**

# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Kcnq1* gene. The schematic diagram is as follows:



- The *Kcnq1* gene has 7 transcripts. According to the structure of *Kcnq1* gene, exon2 of *Kcnq1-201* (ENSMUST00000009689.10) 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 *Kcnq1* gene. The brief process is as follows: CRISPR/Cas9 system

- According to the existing MGI data, Homozygous targeted null or spontaneous mutants show circling and head-tossing behavior and are deaf with inner ear dysmorphology. Paternal inheritance of a deletion of an imprinted control region within an intron of this gene results in small body size.
- The *Kcnq1* gene is located on the Chr7. 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)

## Kcnq1 potassium voltage-gated channel, subfamily Q, member 1 [Mus musculus (house mouse)]

Gene ID: 16535, updated on 31-Jan-2019

### Summary



<b>Official Symbol</b>	Kcnq1 provided by <a href="#">MGI</a>
<b>Official Full Name</b>	potassium voltage-gated channel, subfamily Q, member 1 provided by <a href="#">MGI</a>
<b>Primary source</b>	<a href="#">MGI:MGI:108083</a>
<b>See related</b>	<a href="#">Ensembl:ENSMUSG000000009545</a>
<b>Gene type</b>	protein coding
<b>RefSeq status</b>	VALIDATED
<b>Organism</b>	<a href="#">Mus musculus</a>
<b>Lineage</b>	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
<b>Also known as</b>	AW559127, KVLQT1, Kcna9
<b>Expression</b>	Biased expression in large intestine adult (RPKM 20.0), colon adult (RPKM 19.5) and 10 other tissues <a href="#">See more</a>
<b>Orthologs</b>	<a href="#">human</a> <a href="#">all</a>

# Transcript information (Ensembl)

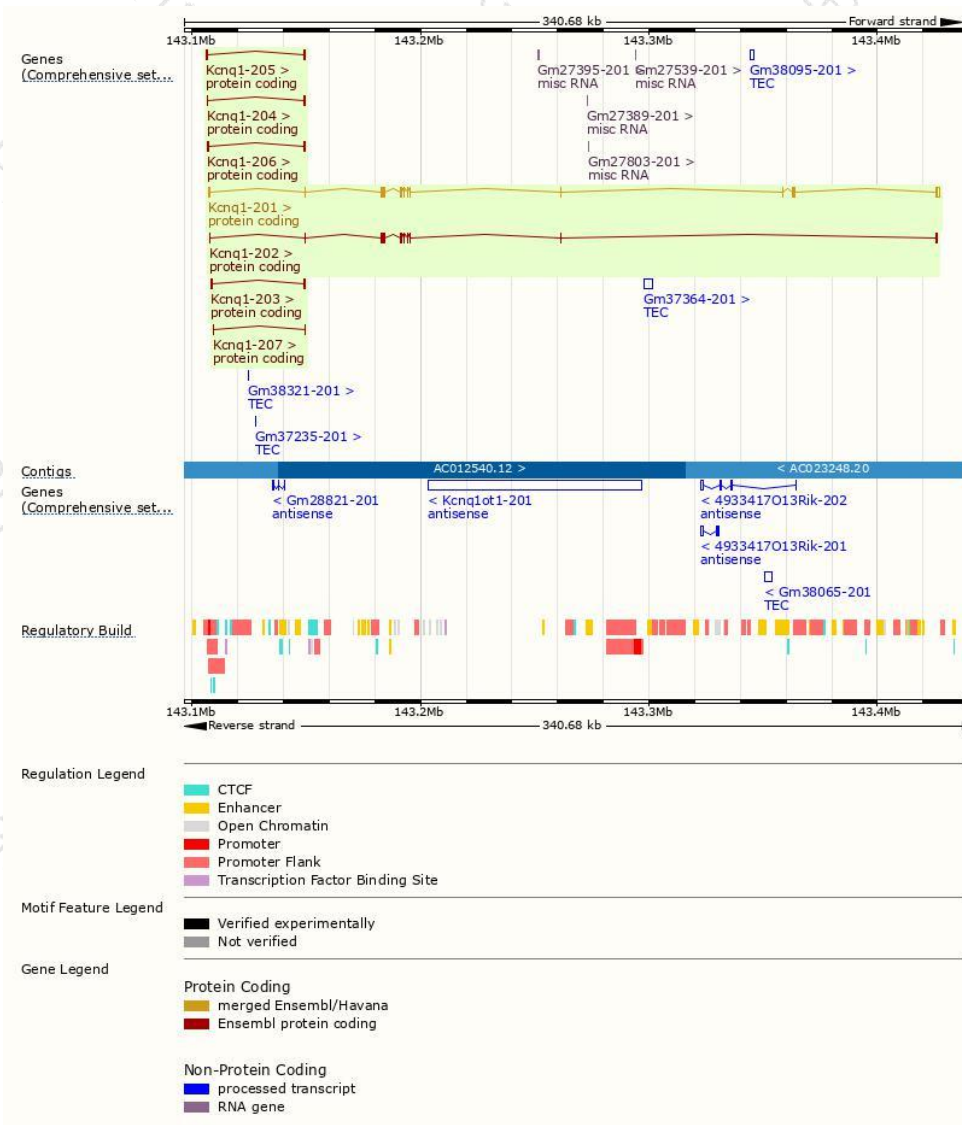
The gene has 7 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Kcnq1-201	<a href="#">ENSMUST00000009689.10</a>	3029	<a href="#">668aa</a>	Protein coding	<a href="#">CCDS22039</a>	<a href="#">P97414</a>	TSL:1 GENCODE basic APPRIS P1
Kcnq1-202	<a href="#">ENSMUST00000185383.1</a>	1535	<a href="#">440aa</a>	Protein coding	-	<a href="#">A0A087WNY7</a>	TSL:1 GENCODE basic
Kcnq1-203	<a href="#">ENSMUST00000186284.6</a>	490	<a href="#">77aa</a>	Protein coding	-	<a href="#">A0A087WQE3</a>	CDS 3' incomplete TSL:1
Kcnq1-206	<a href="#">ENSMUST00000186798.1</a>	294	<a href="#">49aa</a>	Protein coding	-	<a href="#">A0A087WPJ2</a>	CDS 3' incomplete TSL:1
Kcnq1-207	<a href="#">ENSMUST00000187213.1</a>	282	<a href="#">39aa</a>	Protein coding	-	<a href="#">A0A087WPA8</a>	CDS 3' incomplete TSL:1
Kcnq1-204	<a href="#">ENSMUST00000186288.6</a>	196	<a href="#">30aa</a>	Protein coding	-	<a href="#">A0A087WRU8</a>	CDS 3' incomplete TSL:1
Kcnq1-205	<a href="#">ENSMUST00000186488.6</a>	182	<a href="#">60aa</a>	Protein coding	-	<a href="#">A0A087WRK7</a>	CDS 3' incomplete TSL:1

The strategy is based on the design of *Kcnq1-201* transcript,The transcription is shown below

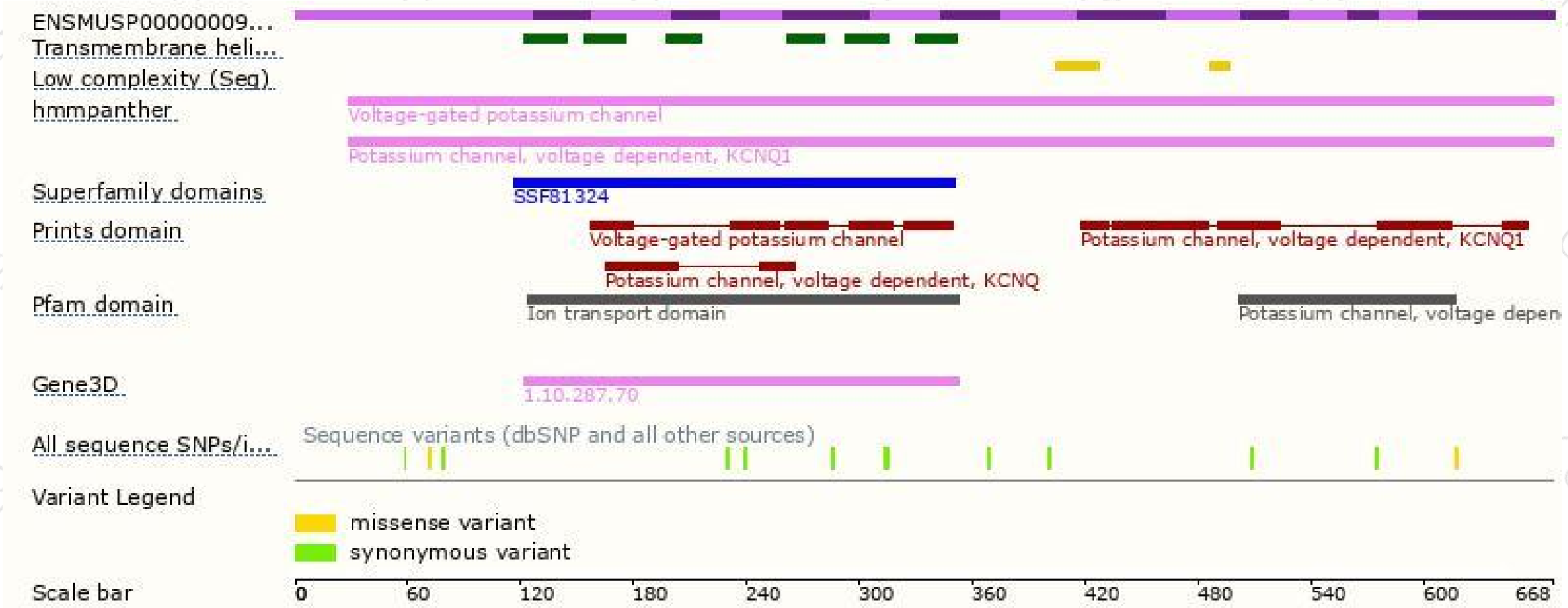


# Genomic location distribution

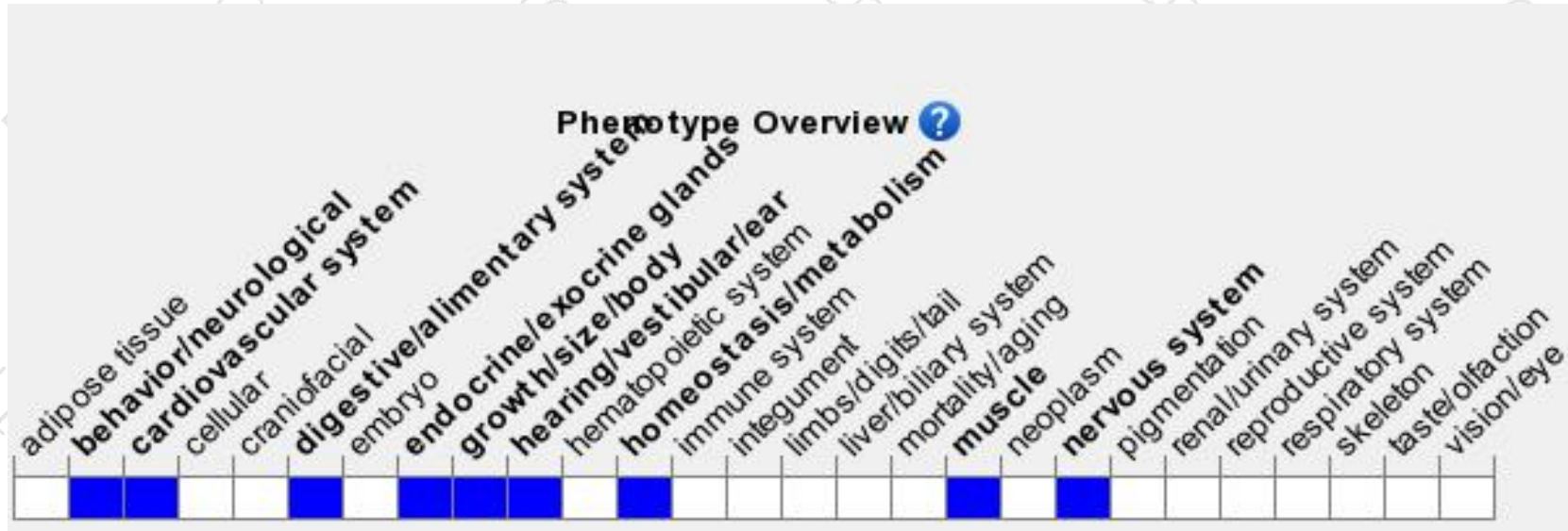




# Protein domain



# Mouse phenotype description(MGI )



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

According to the existing MGI data, Homozygous targeted null or spontaneous mutants show circling and head-tossing behavior and are deaf with inner ear dysmorphology. Paternal inheritance of a deletion of an imprinted control region within an intron of this gene results in small body size.

If you have any questions, you are welcome to inquire.

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