

# Grm7 Cas9-KO Strategy

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## **Project Overview**



Project Name Grm7

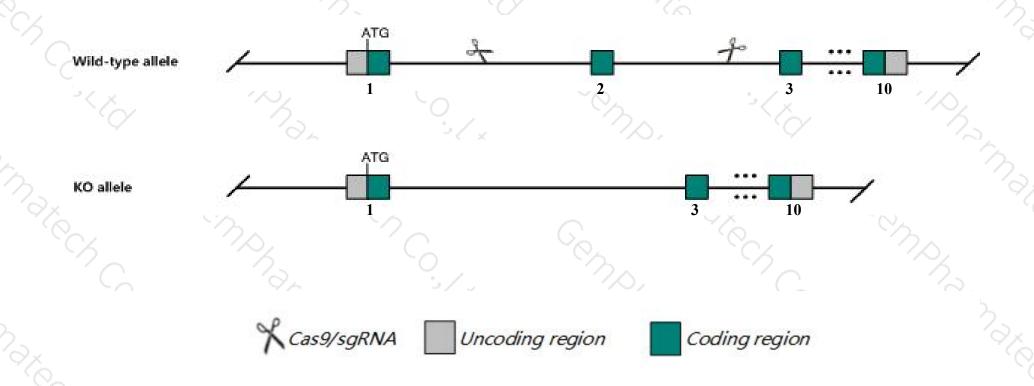
Project type Cas9-KO

Strain background C57BL/6J

## **Knockout strategy**



This model will use CRISPR/Cas9 technology to edit the *Grm7* gene. The schematic diagram is as follows:



### **Technical routes**



- ➤ The *Grm7* gene has 5 transcripts. According to the structure of *Grm7* gene, exon2 of *Grm7-202*(ENSMUST00000172951.1) transcript is recommended as the knockout region. The region contains 217bp coding sequence.

  Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Grm7* gene. The brief process is as follows: sgRNA was transcribed in vitro.Cas9 and sgRNA were microinjected into the fertilized eggs of C57BL/6J 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/6J mice.

### **Notice**



- ➤ According to the existing MGI data, Nullizygous mice exhibit epilepsy and deficits in fear response and conditioned taste aversion. Homozygotes for a knock-in allele show impaired spatial working memory and higher susceptibility to PTZ. Homozygotes for a reporter allele show impaired coordination and higher susceptibility to metrazol.
- > The *Grm7* 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 the gene knockout on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.

## Gene information (NCBI)



#### Grm7 glutamate receptor, metabotropic 7 [Mus musculus (house mouse)]

Gene ID: 108073, updated on 9-Apr-2019

#### Summary

☆ ?

Official Symbol Grm7 provided by MGI

Official Full Name glutamate receptor, metabotropic 7 provided by MGI

Primary source MGI:MGI:1351344

See related Ensembl:ENSMUSG00000056755

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 6330570A01Rik, BB176677, C030018L03, E130018M02Rik, Gpr1g, Gprc1g, SMN2, Tg(SMN2)89Ahmb, mGluR7

Expression Biased expression in cortex adult (RPKM 6.5), frontal lobe adult (RPKM 4.9) and 6 other tissues See more

Orthologs <u>human all</u>

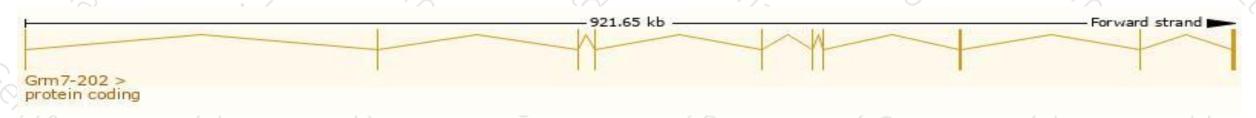
## Transcript information (Ensembl)



The gene has 5 transcripts, all transcripts are shown below:

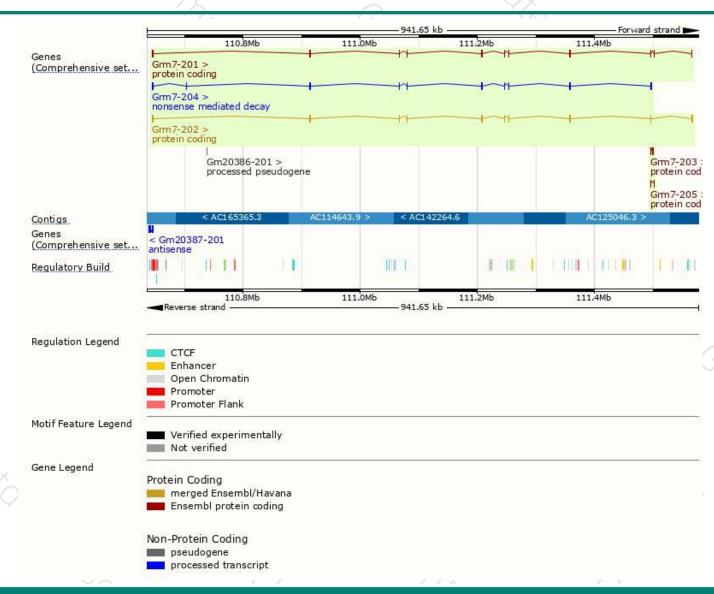
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Grm7-202	ENSMUST00000172951.1	4197	<u>915aa</u>	Protein coding	CCDS39587	Q68ED2	TSL:1 GENCODE basic APPRIS P3
Grm7-201	ENSMUST00000071076.12	3127	922aa	Protein coding	CCDS85110	G5E8D5	TSL:5 GENCODE basic APPRIS ALT1
Grm7-203	ENSMUST00000173001.2	418	74aa	Protein coding	84	G3UYV0	CDS 5' incomplete TSL:5
Grm7-205	ENSMUST00000174310.1	386	<u>75aa</u>	Protein coding	2	G3UY08	CDS 5' incomplete TSL:3
Grm7-204	ENSMUST00000174018.7	4210	188aa	Nonsense mediated decay	15	G3UZU3	TSL:1

The strategy is based on the design of *Grm7-202* transcript, The transcription is shown below



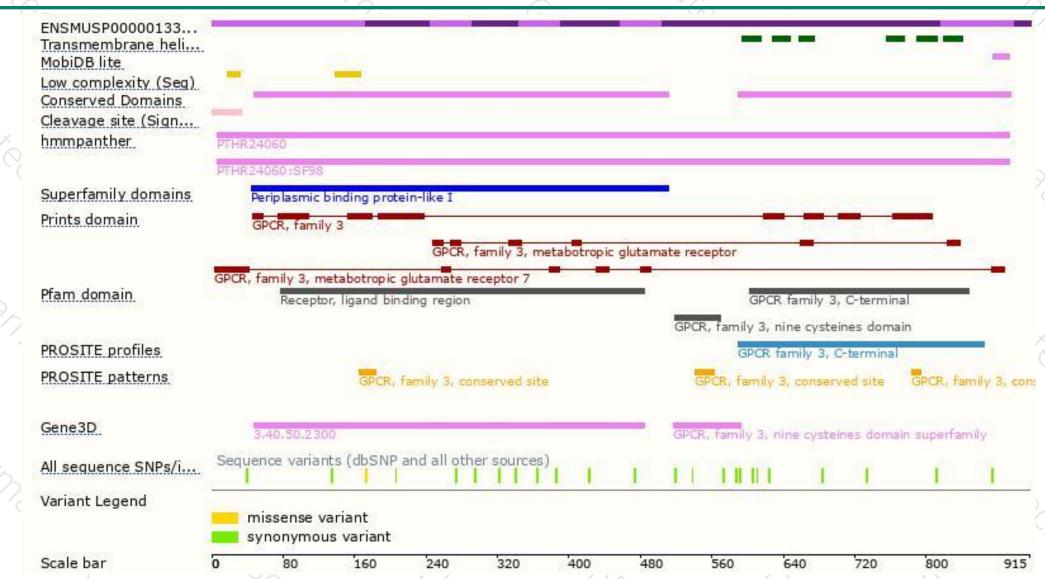
### Genomic location distribution





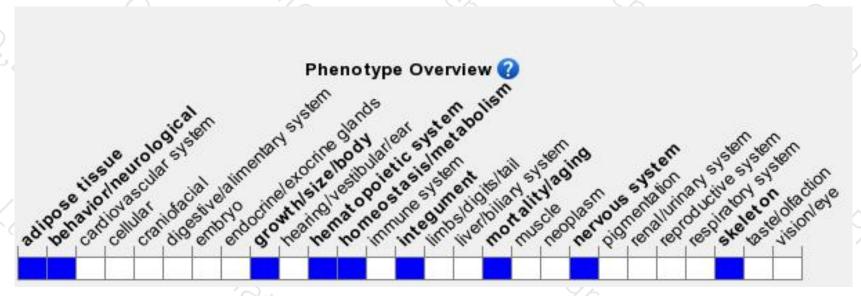
### Protein domain





## 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, Nullizygous mice exhibit epilepsy and deficits in fear response and conditioned taste aversion. Homozygotes for a knock-in allele show impaired spatial working memory and higher susceptibility to PTZ. Homozygotes for a reporter allele show impaired coordination and higher susceptibility to metrazol.



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

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