

Rims1 Cas9-KO Strategy

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Project Overview



Project Name

Rims1

Project type

Cas9-KO

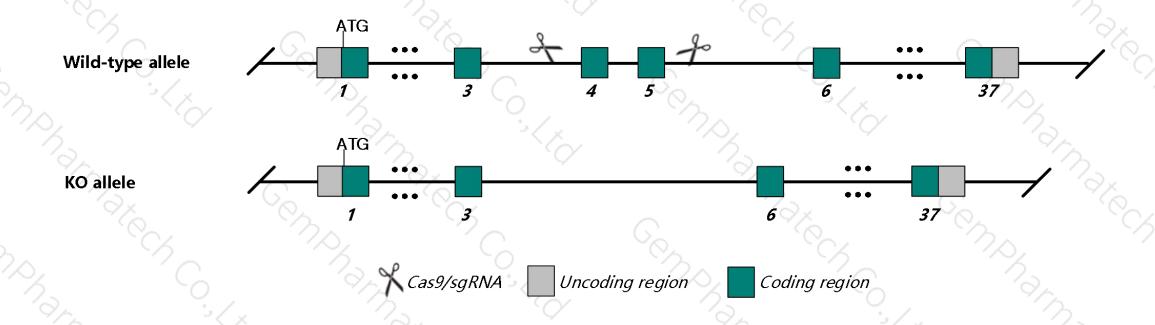
Strain background

C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Rims1* gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Rims1* gene has 11 transcripts. According to the structure of *Rims1* gene, exon4-exon5 of *Rims1-205*(ENSMUST00000097811.9) transcript is recommended as the knockout region. The region contains 226bp of coding reigon. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Rims1* gene. The brief process is as follows: CRISPR/Cas9 system

Notice



- ➤ According to the existing MGI data, Mice homozygous for disruptions in this gene display defects in maternal care and abnormalities in synaptic transmission in the central nervous system.
- > Rims1-207, Rims1-208 and Rims1-210 transcripts are incomplete, so the effect on them are unknown.
- The *Rims1* 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)



Rims1 regulating synaptic membrane exocytosis 1 [Mus musculus (house mouse)]

Gene ID: 116837, updated on 13-Jan-2020

Summary

☆ ?

Official Symbol Rims1 provided by MGI

Official Full Name regulating synaptic membrane exocytosis 1 provided by MGI

Primary source MGI:MGI:2152971

See related Ensembl: ENSMUSG00000041670

Gene type protein coding
RefSeq status VALIDATED
Organism Mus musculus

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

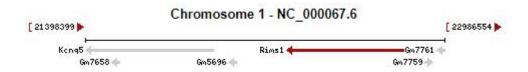
Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus

Also known as Rim; RIM1; RIM1a; Serg1; Rab3ip1; RIM1alpha; C030033M19Rik

Annotation information Annotation category: partial on reference assembly

Expression Biased expression in cortex adult (RPKM 23.6), frontal lobe adult (RPKM 17.0) and 5 other tissues See more

Orthologs human all



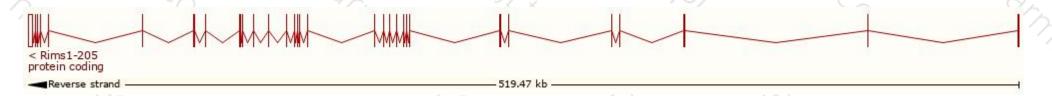
Transcript information (Ensembl)



The gene has 11 transcripts, all transcripts are shown below:

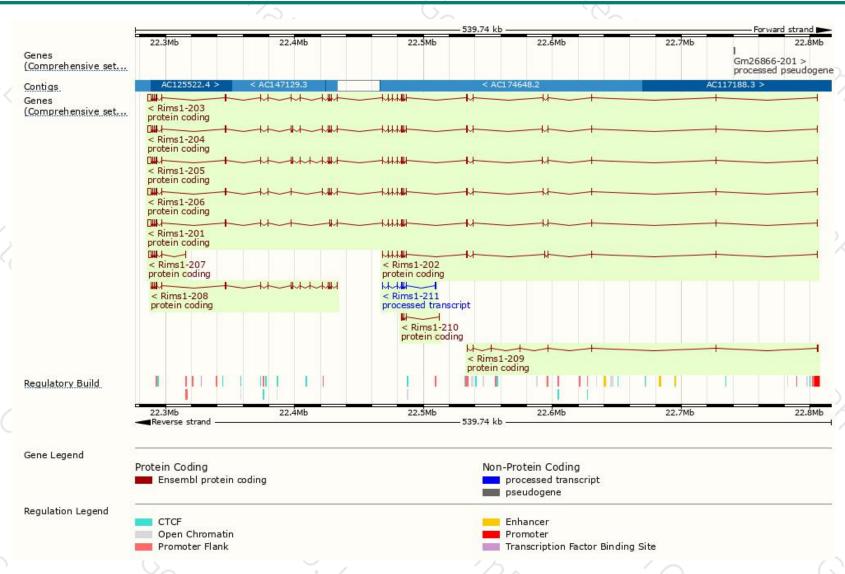
Name	Transcript ID .	bp 🍦	Protein	Biotype	CCDS	UniProt 👙	Flags
Rims1-205	ENSMUST00000097811.9	6446	1424aa	Protein coding	\$70X	F6VBR4₽	TSL:5 GENCODE basic APPRIS P5
Rims1-204	ENSMUST00000097810.9	6362	<u>1396aa</u>	Protein coding	\$7X	F6VBT9₽	TSL:5 GENCODE basic APPRIS ALT
Rims1-203	ENSMUST00000097809.9	6179	<u>1335aa</u>	Protein coding	5733	F6VBV0₽	TSL:5 GENCODE basic APPRIS ALT2
Rims1-206	ENSMUST00000115273.8	6107	<u>1311aa</u>	Protein coding	\$7X	<u>F6Y0S3</u> ₽	TSL:5 GENCODE basic APPRIS ALT2
Rims1-201	ENSMUST00000081544.12	5954	<u>1260aa</u>	Protein coding	57 8	F6TZK4₽	TSL:5 GENCODE basic APPRIS ALT2
Rims1-202	ENSMUST00000097808.6	2411	803aa	Protein coding	5700	E9PWP6函	CDS 3' incomplete TSL:5
Rims1-207	ENSMUST00000164877.7	2294	219aa	Protein coding	5700	Q7TPL9₽	TSL:1 GENCODE basic
Rims1-208	ENSMUST00000185942.1	2189	729aa	Protein coding	\$70	A0A087WSA4₽	CDS 5' incomplete TSL:1
Rims1-209	ENSMUST00000218140.1	1434	388aa	Protein coding	5755	A0A1W2P6S9₽	TSL:5 GENCODE basic
Rims1-210	ENSMUST00000239255.1	516	129aa	Protein coding	\$700	1570	CDS 3' incomplete
Rims1-211	ENSMUST00000239339.1	863	No protein	Processed transcript	8708	(85)	

The strategy is based on the design of Rims 1-205 transcript, The transcription is shown below



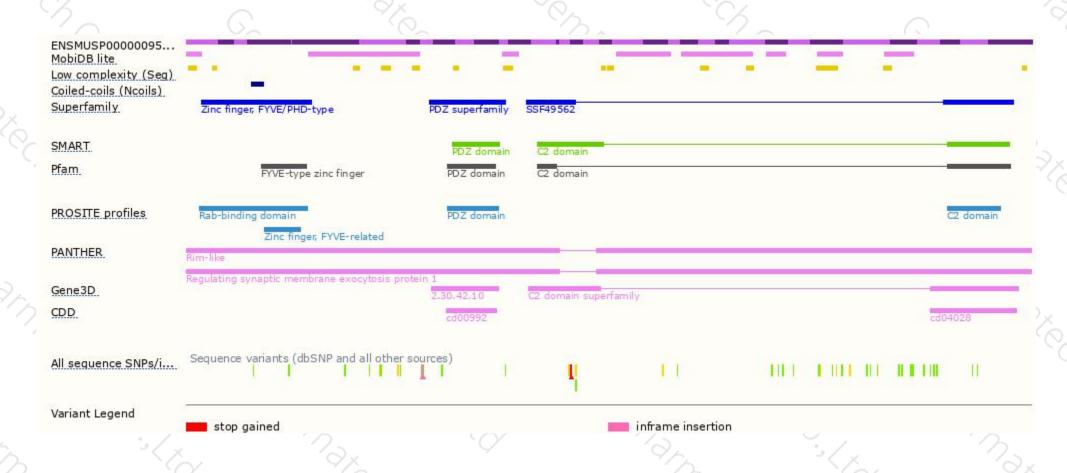
Genomic location distribution





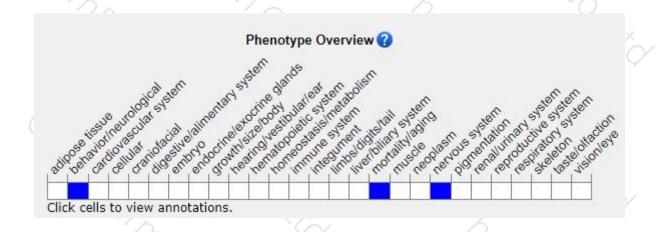
Protein domain





Mouse phenotype description(MGI)





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

Mice homozygous for disruptions in this gene display defects in maternal care and abnormalities in synaptic transmission in the central nervous system.



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





