

Rere Cas9-KO Strategy

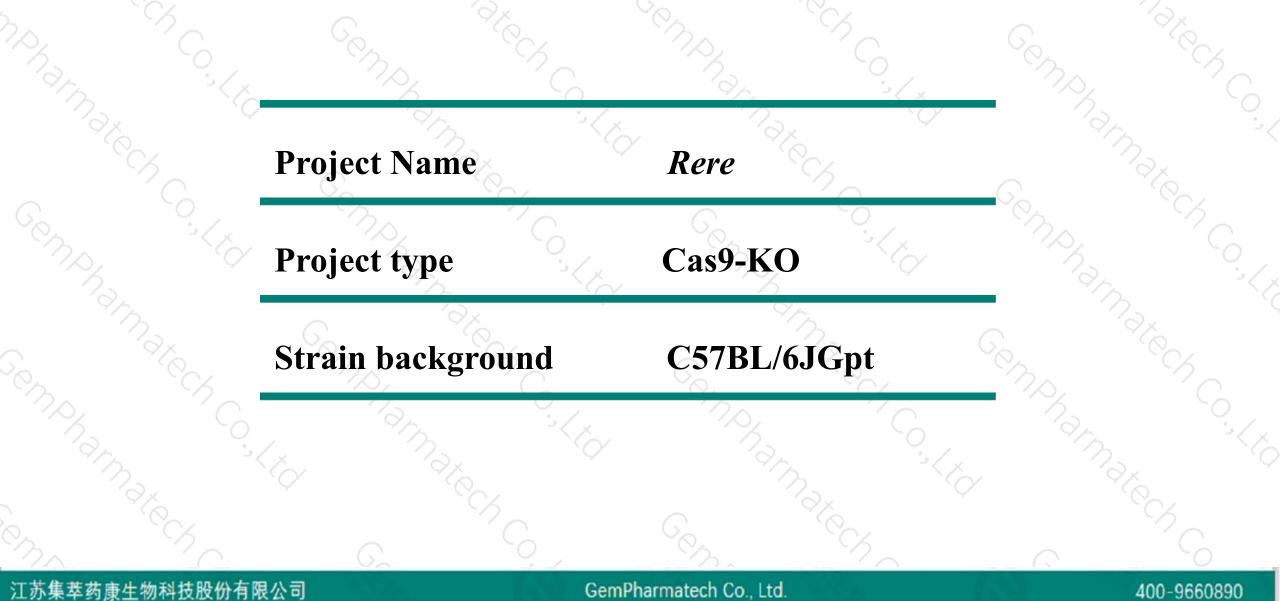
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Reviewer: Xiaojing Li

Design Date: 2020-7-27

Project Overview

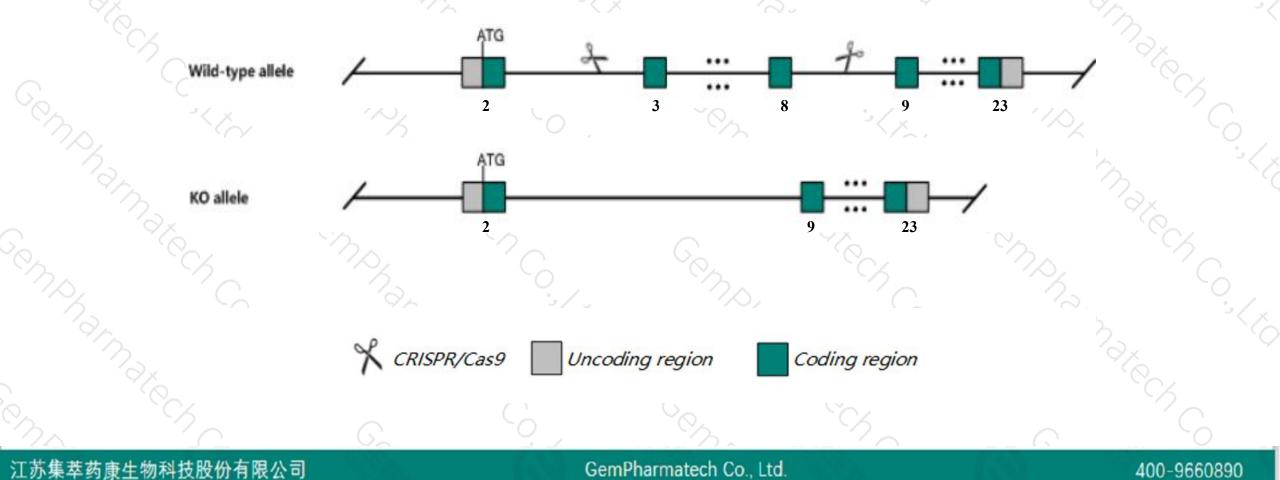




Knockout strategy



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





> The *Rere* gene has 8 transcripts. According to the structure of *Rere* gene, exon3-exon8 of *Rere-*202(ENSMUST00000105682.8) transcript is recommended as the knockout region. The region contains 554bp coding sequence. Knock out the region will result in disruption of protein function.

➤ In this project we use CRISPR/Cas9 technology to modify *Rere* gene. The brief process is as follows: CRISPR/Cas9 system 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.

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According to the existing MGI data,mice homozygous for disruptions in this gene display embryonic lethality with abnormalities in neural tube development, somite development, and in the embryonic heart. Mice homozygous for an ENU-induced allele exhibit narrow snouts, decreased body weight, renal agenesis and small eyes.
The *Rere* gene is located on the Chr4. 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.

Notice

Gene information (NCBI)



\$?

Rere arginine glutamic acid dipeptide (RE) repeats [Mus musculus (house mouse)]

Gene ID: 68703, updated on 13-Mar-2020

Summary

Rere provided by MGI
arginine glutamic acid dipeptide (RE) repeats provided by MGI
MGI:MGI:2683486
Ensembl:ENSMUSG0000039852
protein coding
VALIDATED
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;
Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
1110033A15Rik, Al414665, ARG, ARP, ATN1L, AW742570, Atr2, DNB1, eye, eyes3, mKIAA0458
Ubiquitous expression in adrenal adult (RPKM 15.0), spleen adult (RPKM 12.9) and 28 other tissues See more
human all

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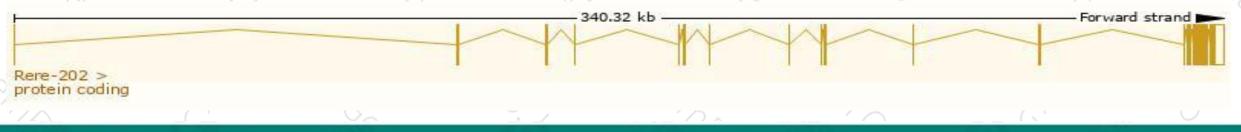
Transcript information (Ensembl)



The gene has 8 transcripts, all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Rere-202	ENSMUST00000105682.8	7728	<u>1558aa</u>	Protein coding	CCDS38979	<u>Q80TZ9</u>	TSL:5 GENCODE basic APPRIS P2
Rere-201	ENSMUST00000105680.8	6503	<u>1290aa</u>	Protein coding	-	<u>A2A7T3</u>	TSL:5 GENCODE basic APPRIS ALT2
Rere-208	ENSMUST00000219467.1	1305	<u>435aa</u>	Protein coding	-	A0A1W2P787	CDS 5' and 3' incomplete TSL:2
Rere-205	ENSMUST0000136646.1	939	<u>199aa</u>	Protein coding	3	<u>A2A7T1</u>	CDS 5' incomplete TSL:1
Rere-204	ENSMUST00000131600.1	535	<u>178aa</u>	Protein coding	2:	<u>B1AS11</u>	CDS 5' and 3' incomplete TSL:2
Rere-206	ENSMUST00000137112.1	496	No protein	Processed transcript	-	1574	TSL:5
Rere-203	ENSMUST00000129188.1	417	No protein	Processed transcript	-		TSL:2
Rere-207	ENSMUST00000142379.1	379	No protein	Processed transcript	21	1.12	TSL:3

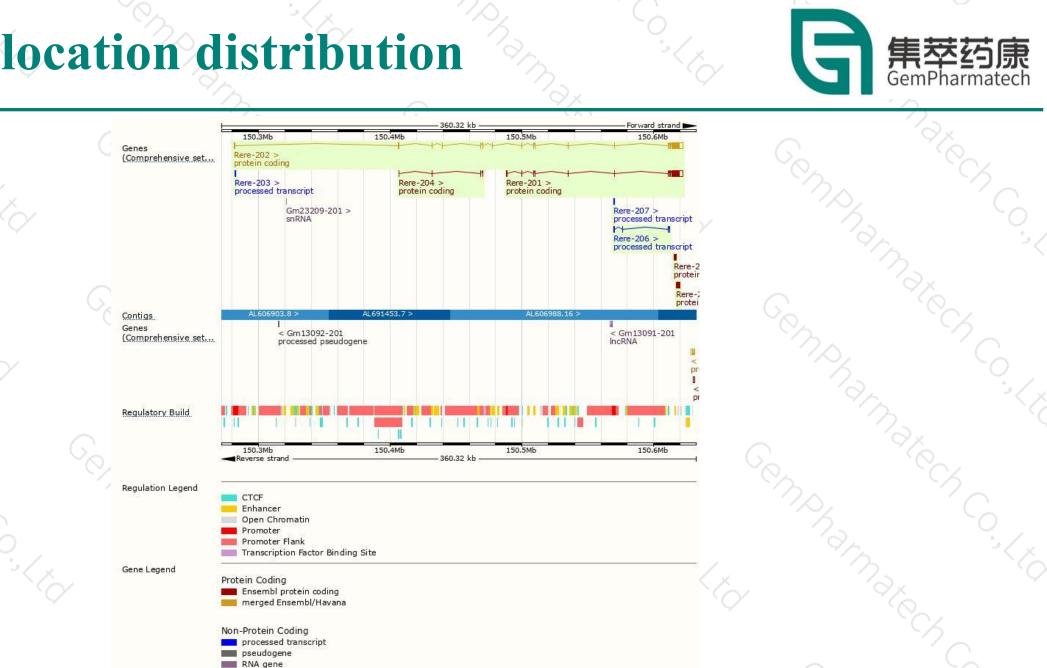
The strategy is based on the design of *Rere-202* transcript, the transcription is shown below:



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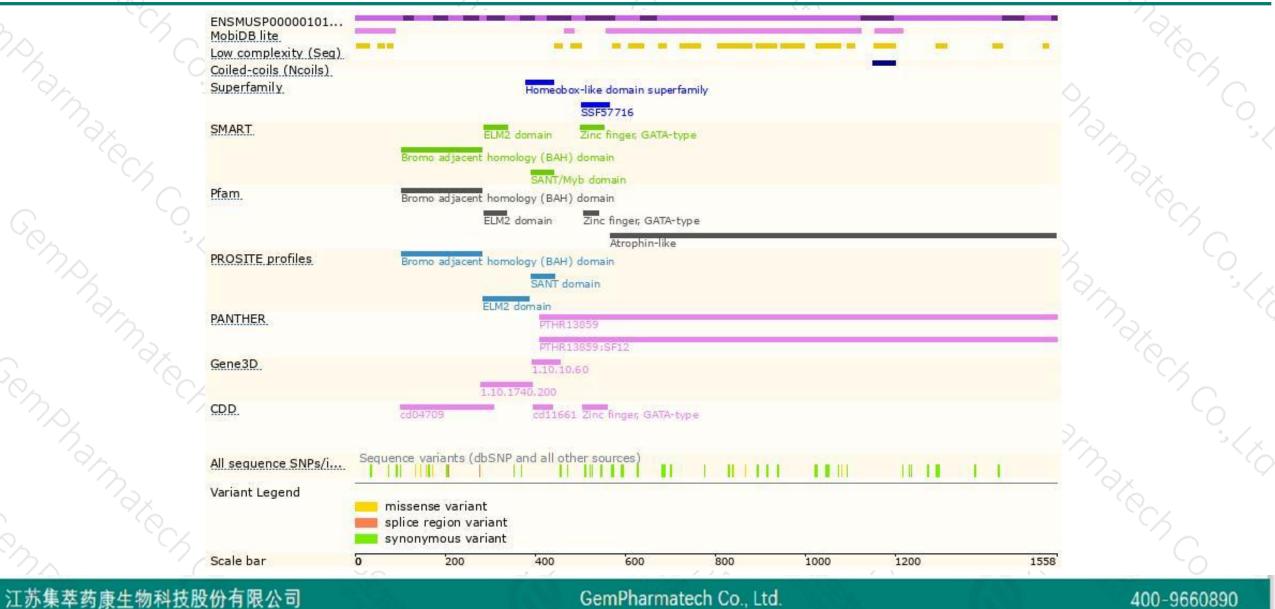
Genomic location distribution



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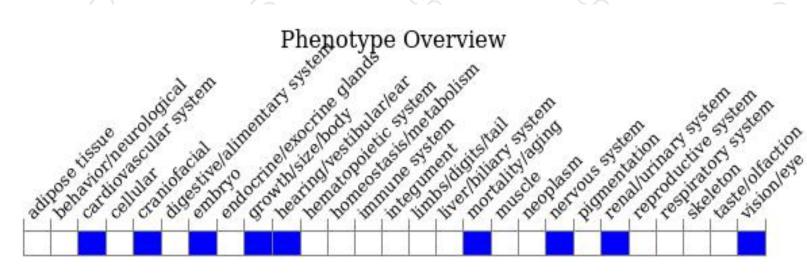
Protein domain





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 disruptions in this gene display embryonic lethality with abnormalities in neural tube development, somite development, and in the embryonic heart. Mice homozygous for an ENU-induced allele exhibit narrow snouts, decreased body weight, renal agenesis and small eyes.

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



