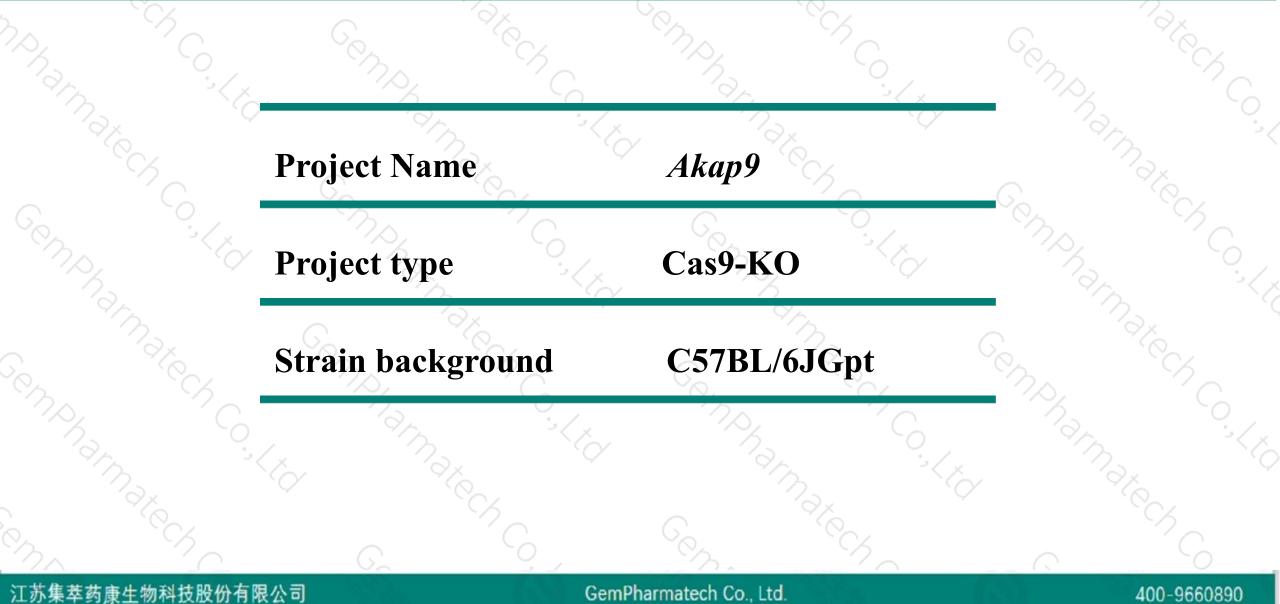


Akap9 Cas9-KO Strategy

Designer: Xiaojing Li Design Date: 2020-1-8 Reviewer: JiaYu

Project Overview

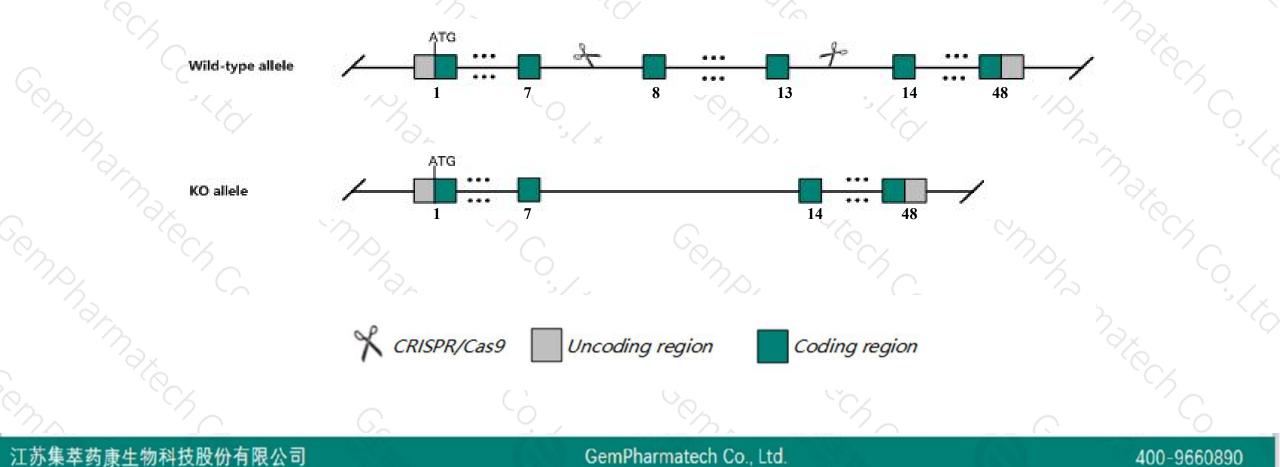




Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Akap9 gene. The schematic diagram is as follows:





- The Akap9 gene has 16 transcripts. According to the structure of Akap9 gene, exon8-exon13 of Akap9-201 (ENSMUST00000044492.9) transcript is recommended as the knockout region. The region contains 821bp coding sequence. Knock out the region will result in disruption of protein function.
- > In this project we use CRISPR/Cas9 technology to modify Akap9 gene. The brief process is as follows: CRISPR/Cas9 system

- According to the existing MGI data, Mice homozygous for a chemically induced allele exhibit male infertily with abnormal spermatogenesis and Sertoli maturation.
- The Akap9 gene is located on the Chr5. 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)



Akap9 A kinase (PRKA) anchor protein (yotiao) 9 [Mus musculus (house mouse)]

Gene ID: 100986, updated on 12-Aug-2019

Summary

Official SymbolAkap9 provided by MGIOfficial Full NameA kinase (PRKA) anchor protein (yotiao) 9 provided by MGIPrimary sourceMGI:MGI:2178217See relatedEnsembl:ENSMUSG0000040407Gene typeprotein codingRefSeq statusVALIDATEDOrganismMus musculusLineageEukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;
Myomorpha; Muroidea; Murinae; Mus; MusAlso known asPRKA9; AKAP-9; C79026; mei2-5; AKAP450; repro12; AW545847; G1-448-15; mKIAA0803; 5730481H23RikExpressionBroad expression in CNS E14 (RPKM 12.4), CNS E18 (RPKM 10.9) and 23 other tissues See more
human all

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Transcript information (Ensembl)



Name 🔺	Transcript ID 👙	bp 🖕	Protein 🖕	Biotype 🝦	CCDS 🝦	UniProt 🝦	Flags
Akap9-201	ENSMUST0000044492.9	13123	<u>3779aa</u>	Protein coding	<u>CCDS19070</u> 교	<u>E9QQ10</u> 교	TSL:1 GENCODE basic APPRIS P1
Akap9-202	ENSMUST00000123119.1	670	No protein	Retained intron	15	11.5.1	TSL:3
Akap9-203	ENSMUST00000132058.1	3646	No protein	Retained intron	17	858	TSL:1
Akap9-204	ENSMUST00000133952.1	499	<u>113aa</u>	Protein coding	17	A0A0G2JFW0@	CDS 5' incomplete TSL:3
Akap9-205	ENSMUST00000140977.7	375	No protein	IncRNA	15	11.5.1	TSL:3
Akap9-206	ENSMUST00000141817.1	491	No protein	Retained intron	17	150	TSL:2
Akap9-207	ENSMUST00000143365.7	2010	<u>581aa</u>	Protein coding	17	A0A140T8S1@	CDS 3' incomplete TSL:1
Akap9-208	ENSMUST00000147629.1	718	<u>56aa</u>	Nonsense mediated decay	<i></i>	<u>G3UX24</u> 🗗	CDS 5' incomplete TSL:5
Akap9-209	ENSMUST00000176863.1	595	<u>198aa</u>	Protein coding	17	H3BLL7@	CDS 5' and 3' incomplete TSL:3
Akap9-210	ENSMUST00000177448.1	491	No protein	Retained intron	17	11.50	TSL:2
Akap9-211	ENSMUST00000196350.1	2323	No protein	Retained intron	17	11.50	TSL:NA
Akap9-212	ENSMUST00000196837.1	460	No protein	Retained intron	5	11.54	TSL:NA
Akap9-213	ENSMUST00000196918.1	1676	No protein	Retained intron	17	10.50	TSL:NA
Akap9-214	ENSMUST00000197366.1	479	No protein	Retained intron	17	15.5	TSL:NA
Akap9-215	ENSMUST00000198973.1	581	No protein	Retained intron	5	10.750	TSL:NA
Akap9-216	ENSMUST00000238634.1	1352	<u>282aa</u>	Protein coding	5	11-11	CDS 3' incomplete

The gene has 16 transcripts, all transcripts are shown below:

The strategy is based on the design of Akap9-201 transcript, The transcription is shown below

Akap9-201 > protein coding

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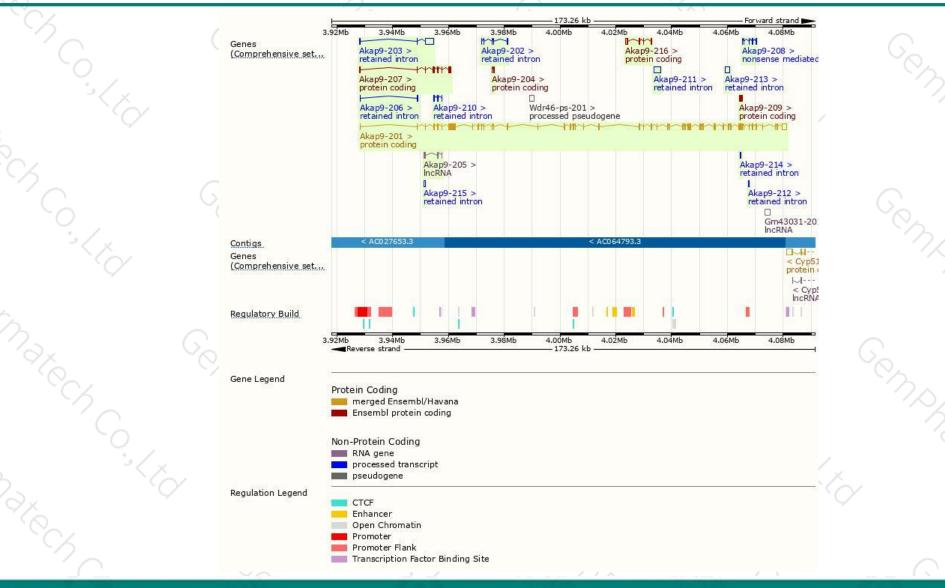
-153.04 kb

400-9660890

Forward strand

Genomic location distribution



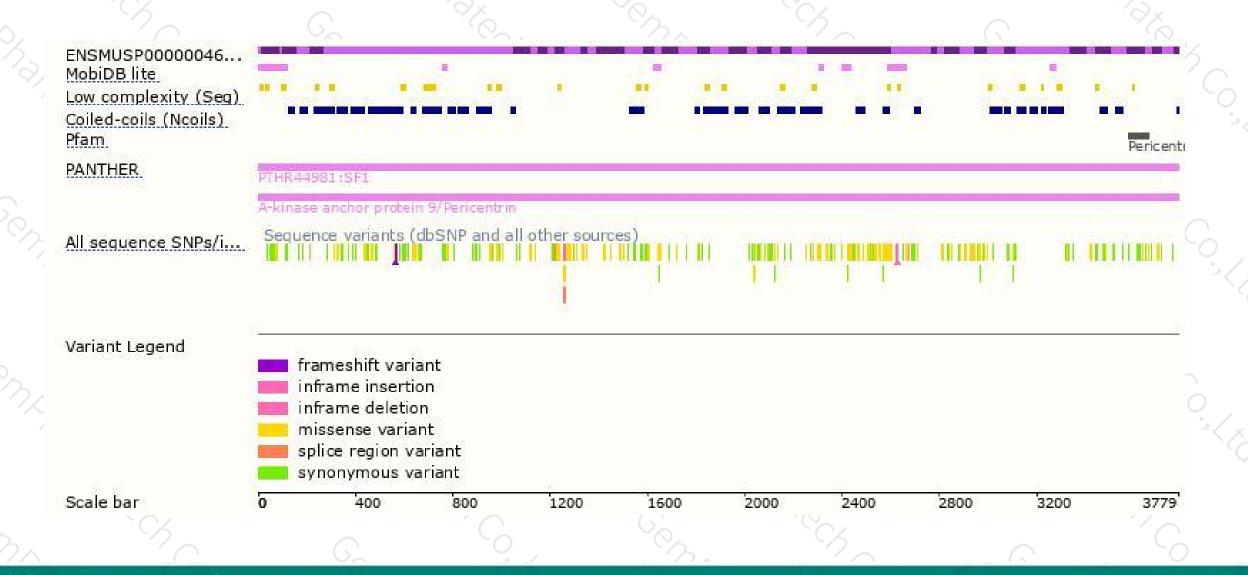


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Protein domain



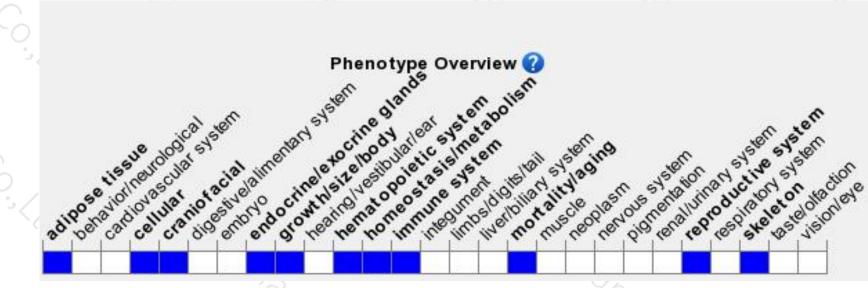


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Mouse phenotype description(MGI)





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 chemically induced allele exhibit male infertily with abnormal spermatogenesis and Sertoli maturation.

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



