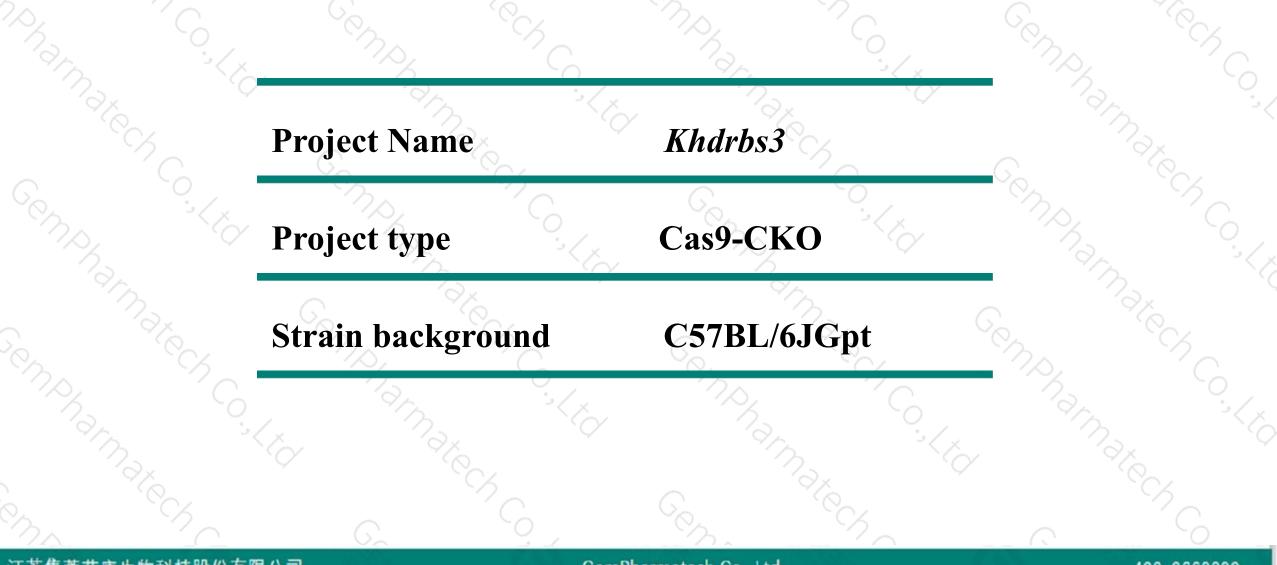


Khdrbs3 Cas9-CKO Strategy

Designer:Xiaojing Li Reviewer:JiaYu Design Date:2020-2-14

Project Overview





江苏集萃药康生物科技股份有限公司

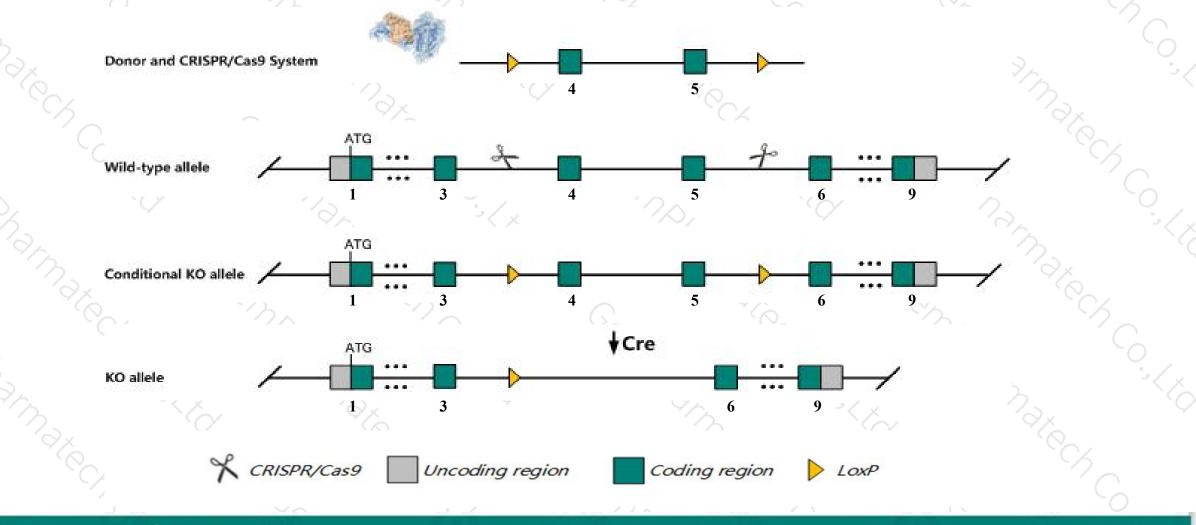
GemPharmatech Co., Ltd.

400-9660890

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Khdrbs3* gene. The schematic diagram is as follows:



江苏集萃药康生物科技股份有限公司

GemPharmatech Co., Ltd.

400-9660890



The *Khdrbs3* gene has 6 transcripts. According to the structure of *Khdrbs3* gene, exon4-exon5 of *Khdrbs3-201* (ENSMUST00000022954.6) transcript is recommended as the knockout region. The region contains 287bp coding sequence. Knock out the region will result in disruption of protein function.

In this project we use CRISPR/Cas9 technology to modify *Khdrbs3* gene. The brief process is as follows:CRISPR/Cas9 system and Donor 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.

The flox mice will be knocked out after mating with mice expressing Cre recombinase, resulting in the loss of function of the target gene in specific tissues and cell types.



- According to the existing MGI data, Mice homozygous for a knock-out allele are viable and fertile with no detectable spatial memory deficits. Males sire slightly smaller litters than control males.
 - > The *Khdrbs3* gene is located on the Chr15. 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 loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at existing technological level.

江苏集萃药康生物科技股份有限公司

Gene information (NCBI)

Khdrbs3 KH domain containing, RNA binding, signal transduction associated 3 [Mus musculus (house mouse)]

Gene ID: 13992, updated on 12-Aug-2019

Summary

Official Symbol Khdrbs3 provided by <u>MGI</u> Official Full Name KH domain containing, RNA binding, signal transduction associated 3 provided by <u>MGI</u>

Primary source MGI:MGI:1313312

See related Ensembl:ENSMUSG00000022332

Gene type protein coding

RefSeq status VALIDATED

Organism Mus musculus

Lineage Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Murinae; Mus; Mus

Also known as Etle; Salp; Slm2; SLM-2; T-STAR

Expression Broad expression in CNS E18 (RPKM 36.8), cortex adult (RPKM 31.0) and 19 other tissues See more Orthologs human all

GemPharmatech Co., Ltd.





☆ ?

Transcript information (Ensembl)



400-9660890

The gene has 6 transcripts, all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Khdrbs3-201	ENSMUST00000022954.6	1887	<u>346aa</u>	Protein coding	CCDS27513	Q9R226	TSL:1 GENCODE basic APPRIS P1
Khdrbs3-206	ENSMUST00000230847.1	1344	<u>258aa</u>	Protein coding	3 7	A0A2R8VJR4	GENCODE basic
Khdrbs3-205	ENSMUST00000230073.1	501	<u>126aa</u>	Protein coding	84	A0A2R8VHM1	CDS 3' incomplete
Khdrbs3-202	ENSMUST00000229234.1	2861	<u>346aa</u>	Nonsense mediated decay	CCDS27513	<u>Q9R226</u>	
Khdrbs3-204	ENSMUST00000229683.1	1740	<u>346aa</u>	Nonsense mediated decay	CCDS27513	Q9R226	
Khdrbs3-203	ENSMUST00000229534.1	688	No protein	Retained intron		-	

The strategy is based on the design of *Khdrbs3-201* transcript, The transcription is shown below

protein coding	Khdrbs3-201 > protein coding		-16	5.09 kb	Fo	rward strand
	protein coding	 Va		(D.)	(\)	

Genomic location distribution



400-9660890



江苏集萃药康生物科技股份有限公司

GemPharmatech Co., Ltd.

Protein domain



		~`C.4		7A.				
ENSMUSP00000022 MobiDB lite						-		20
Low complexity (Seg)						1		
Superfamily		K Homology doma	in, type 1 supe	rfamily				
SMART	1	Homology domain	1	- 1 C				
Pfam	KHDRBS, Qua1 dom	ain				s	Sam68, tyrosine-i	rich domain
		K Homology dom	ain, type 1					0
PANTHER	PTHR11208	17945						
	PTHR11208:SF29							
Gene3D	K Homology domain,	type 1 superfamily	/					
		cd02395						
All sequence SNPs/i	Sequence variants	(dbSNP and all o	ther sources)	14	1. 10	60		2
Variant Legend	missense var							O
Scale bar	0 40	80	120	160	200	240	280	346
2°CX		70		G.	0			20
-70		\sim		~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~				

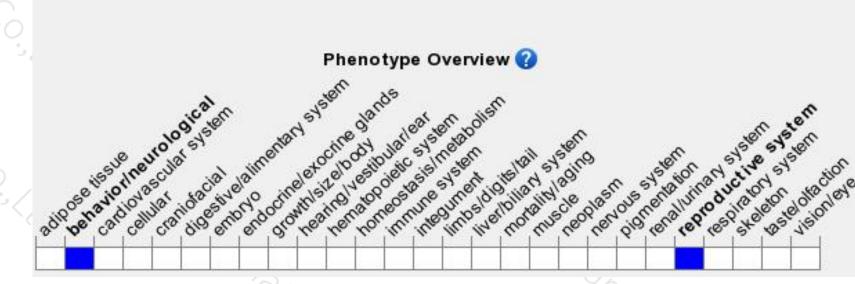
江苏集萃药康生物科技股份有限公司

GemPharmatech Co., Ltd.

400-9660890

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 knock-out allele are viable and fertile with no detectable spatial memory deficits. Males sire slightly smaller litters than control males.



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



