

# Khdrbs2 Cas9-KO Strategy

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Reviewer: Huimin Su

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## **Project Overview**



**Project Name** 

Khdrbs2

**Project type** 

Cas9-KO

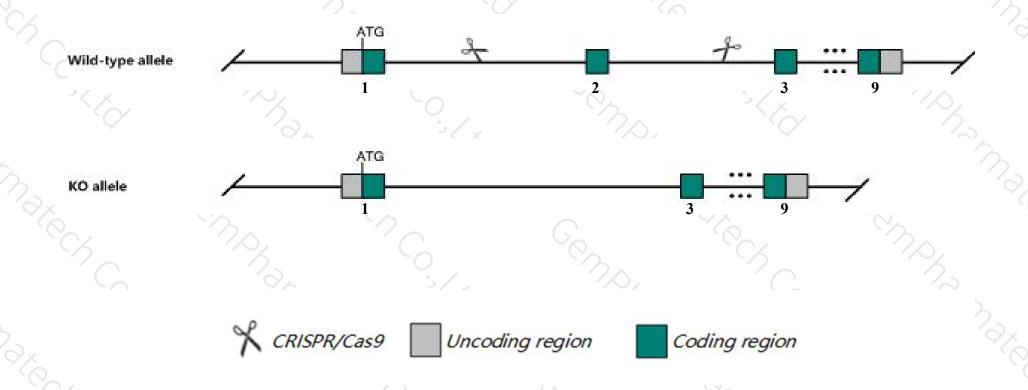
Strain background

C57BL/6JGpt

## **Knockout strategy**



This model will use CRISPR/Cas9 technology to edit the *Khdrbs2* gene. The schematic diagram is as follows:



## **Technical routes**



- ➤ The *Khdrbs2* gene has 5 transcripts. According to the structure of *Khdrbs2* gene, exon2 of *Khdrbs2-201*(ENSMUST00000027226.11) transcript is recommended as the knockout region. The region contains 128bp coding sequence Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Khdrbs2* gene. The brief process is as follows: CRISPR/Cas9 systematically systems.

### **Notice**



- > According to the existing MGI data, Homozygous mutant animals display smaller brain size and reduced weight in the cerebellum.
- The *Khdrbs2* 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)



#### Khdrbs2 KH domain containing, RNA binding, signal transduction associated 2 [Mus musculus (house mouse)]

Gene ID: 170771, updated on 31-Jan-2019

#### Summary



Official Symbol Khdrbs2 provided by MGI

Official Full Name KH domain containing, RNA binding, signal transduction associated 2 provided by MGI

Primary source MGI:MGI:2159649

See related Ensembl:ENSMUSG00000026058

Gene type protein coding
RefSeq status REVIEWED
Organism Mus musculus

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

Muroidea; Muridae; Murinae; Mus; Mus

Also known as 6330586C16Rik, Slim1, Slm-1, Slm1, TG-RP135, Tg(LRRK2\*R1441G)135Cjli, mSLM-1

Summary The protein encoded by this gene is similar to the src associated in mitosis, 68 kDa protein, which is an RNA-binding protein and a

substrate for Src-family tyrosine kinases during mitosis. This protein has a KH RNA-binding motif and proline-rich motifs which may be SH2 and SH3 domain binding sites. A similar rat protein is an RNA-binding protein which is tyrosine phosphorylated by Src during mitosis. These studies also suggest that the rat protein may function as an adaptor protein for Src by binding the SH2 and SH3 domains of various other

proteins. [provided by RefSeq, Jul 2008]

Expression Biased expression in CNS E18 (RPKM 5.5), whole brain E14.5 (RPKM 5.1) and 6 other tissuesSee more

Orthologs <u>human all</u>

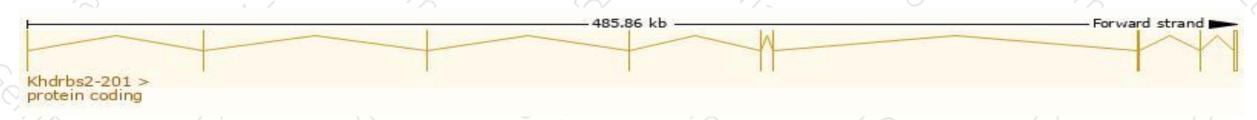
## Transcript information (Ensembl)



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

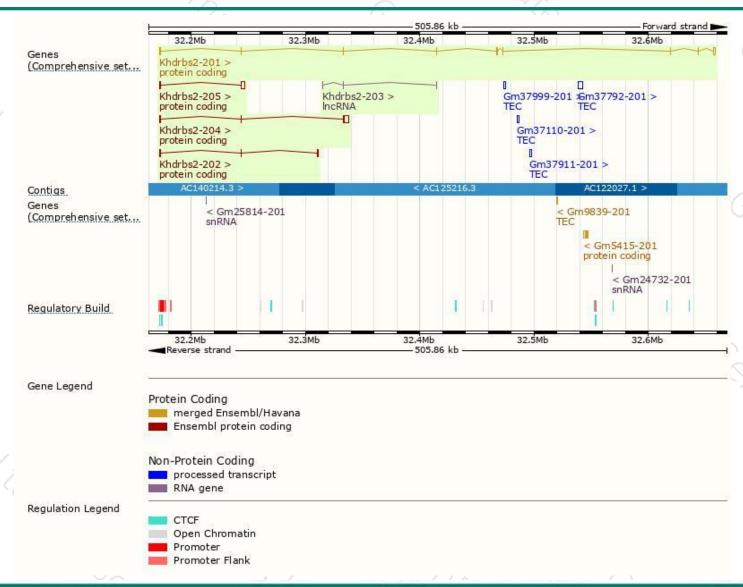
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Khdrbs2-201	ENSMUST00000027226.11	2281	<u>349aa</u>	Protein coding	CCDS14861	Q9WU01	TSL:1 GENCODE basic APPRIS P1
Khdrbs2-204	ENSMUST00000188257.6	5278	<u>193aa</u>	Protein coding	8 <del>-</del>	Q9WU01	TSL:1 GENCODE basic
Khdrbs2-205	ENSMUST00000189878.1	3259	91aa	Protein coding	¥ <del>-</del>	A0A087WSL3	TSL:1 GENCODE basic
Khdrbs2-202	ENSMUST00000185666.1	543	92aa	Protein coding	1 4	A0A087WR32	TSL:2 GENCODE basic
Khdrbs2-203	ENSMUST00000186984.1	275	No protein	IncRNA		12.	TSL:3

The strategy is based on the design of *Khdrbs2-201* transcript, The transcription is shown below



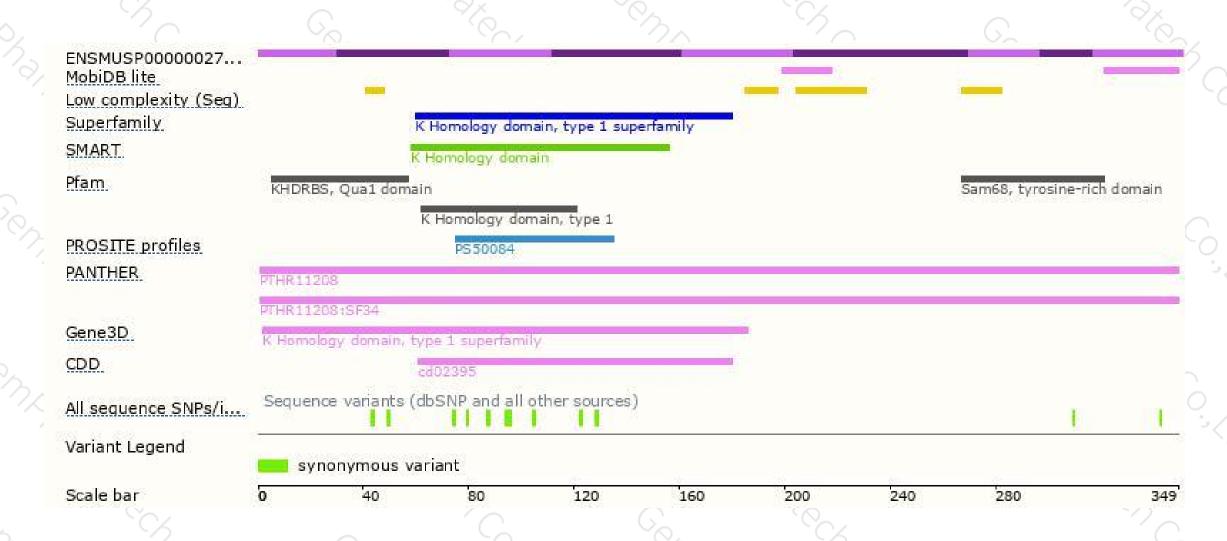
### Genomic location distribution





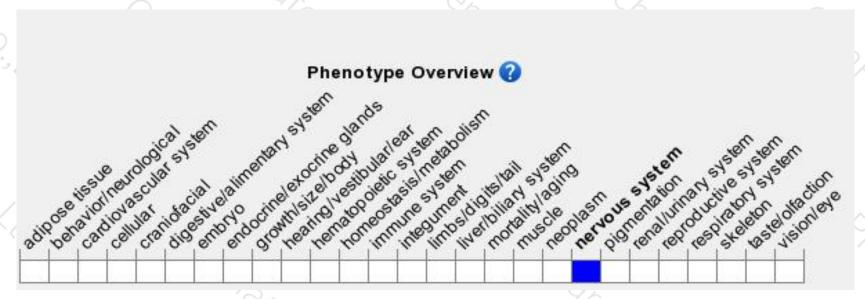
### Protein domain





## 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, Homozygous mutant animals display smaller brain size and reduced weight in the cerebellum.



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





