

# Arhgef6 Cas9-KO Strategy

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Reviewer: Ruirui Zhang

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## **Project Overview**



**Project Name** 

Arhgef6

**Project type** 

Cas9-KO

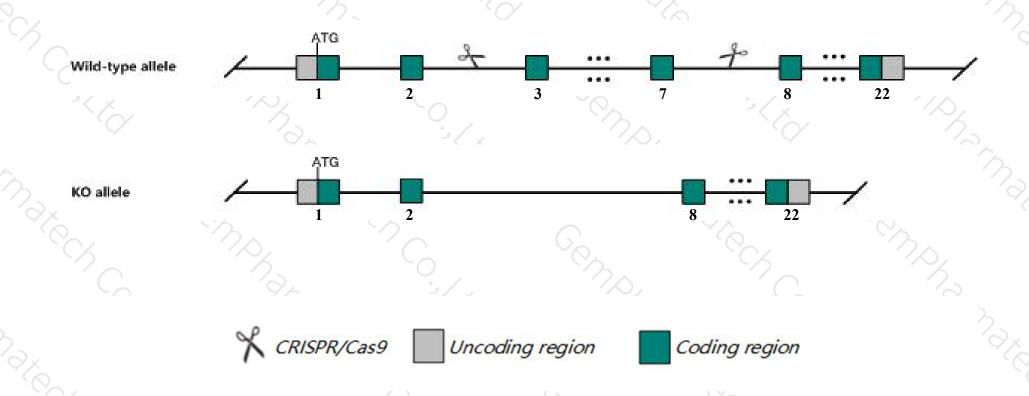
Strain background

C57BL/6JGpt

# **Knockout strategy**



This model will use CRISPR/Cas9 technology to edit the Arhgef6 gene. The schematic diagram is as follows:



### **Technical routes**



- The *Arhgef6* gene has 8 transcripts. According to the structure of *Arhgef6* gene, exon3-exon7 of *Arhgef6-201* (ENSMUST00000033468.10) transcript is recommended as the knockout region. The region contains 575bp coding sequence Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Arhgef6* gene. The brief process is as follows: CRISPR/Cas9 syste

### **Notice**



- ➤ According to the existing MGI data, Mice homozygous for a truncated allele exhibit decreased mature lymphocyte cell numbers, decreased B and T cell proliferation, and defective humeral response. Mice homozygous for a reporter allele exhibit abnormal dendrite morphology and synaptic plasticity and cognitive defects.
- ➤ The transcript *Arhgef6-207* is incomplete, so the effect on it is unknown.
- > The *Arhgef6* gene is located on the ChrX. 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)



#### Arhgef6 Rac/Cdc42 guanine nucleotide exchange factor (GEF) 6 [ Mus musculus (house mouse) ]

Gene ID: 73341, updated on 12-Aug-2019

#### Summary



Official Symbol Arhgef6 provided by MGI

Official Full Name Rac/Cdc42 guanine nucleotide exchange factor (GEF) 6 provided by MGI

Primary source MGI:MGI:1920591

See related Ensembl: ENSMUSG00000031133

Gene type protein coding
RefSeq status VALIDATED
Organism <u>Mus musculus</u>

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

Muroidea; Muridae; Murinae; Mus; Mus

Also known as alpha-PIX; 1600028C08Rik; 1700038J06Rik; 4930592P22Rik

Expression Biased expression in placenta adult (RPKM 38.2), thymus adult (RPKM 12.8) and 12 other tissues See more

Orthologs <u>human</u> <u>all</u>

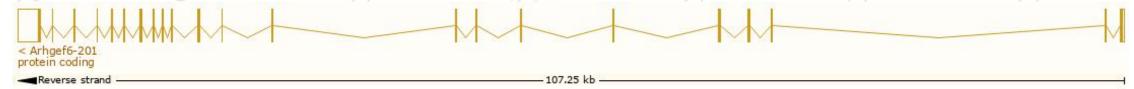
# Transcript information (Ensembl)



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

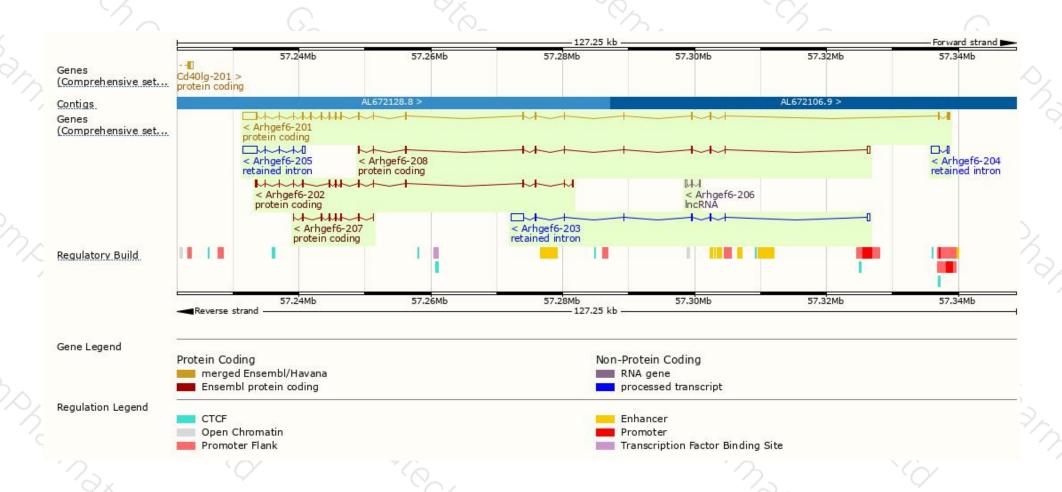
Name	Transcript ID	bp 🍦	Protein	Biotype	CCDS	UniProt 🍦	Flags
Arhgef6-201	ENSMUST00000033468.10	4569	<u>795aa</u>	Protein coding	CCDS40984₽	F6WMJ3₽	TSL:1 GENCODE basic APPRIS P1
Arhgef6-202	ENSMUST00000114735.8	1614	<u>483aa</u>	Protein coding	.=	A2AFJ8₽	TSL:1 GENCODE basic
Arhgef6-208	ENSMUST00000176986.7	1604	<u>302aa</u>	Protein coding	-	<u>H3BJ53</u> ₽	CDS 3' incomplete TSL:5
Arhgef6-207	ENSMUST00000175900.1	748	250aa	Protein coding	-	H3BKQ0₽	CDS 5' and 3' incomplete TSL:5
Arhgef6-203	ENSMUST00000134028.8	2951	No protein	Retained intron	=	27	TSL:1
Arhgef6-205	ENSMUST00000151768.1	2696	No protein	Retained intron	2	2	TSL:1
Arhgef6-204	ENSMUST00000135098.1	1301	No protein	Retained intron		20	TSL:1
Arhgef6-206	ENSMUST00000175636.1	675	No protein	IncRNA	-	-	TSL:1

The strategy is based on the design of Arhgef6-201 transcript, The transcription is shown below



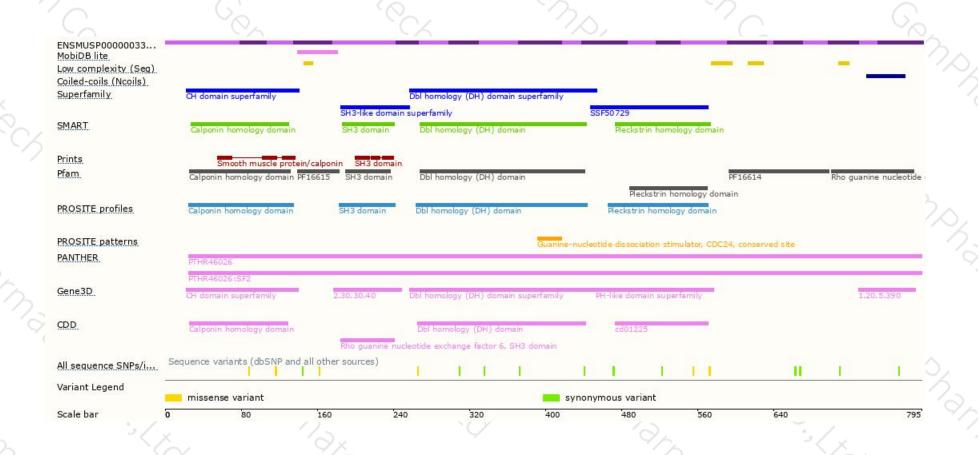
### Genomic location distribution





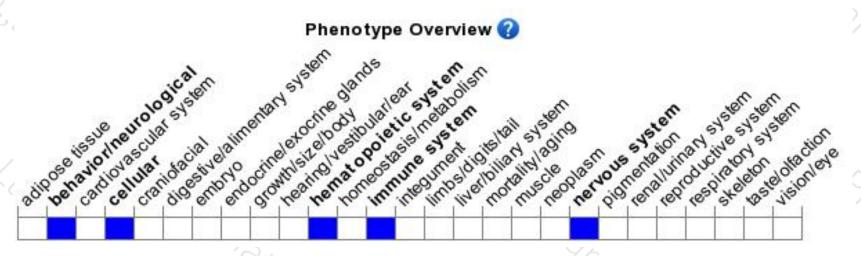
### Protein domain





# 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, Mice homozygous for a truncated allele exhibit decreased mature lymphocyte cell numbers, decreased B and T cell proliferation, and defective humeral response. Mice homozygous for a reporter allele exhibit abnormal dendrite morphology and synaptic plasticity and cognitive defects.



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





