# Ackr3 Cas9-CKO Strategy

**Designer:** 

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**Reviewer:** 

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**Design Date:** 

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## **Project Overview**



**Project Name** 

Ackr3

**Project type** 

Cas9-CKO

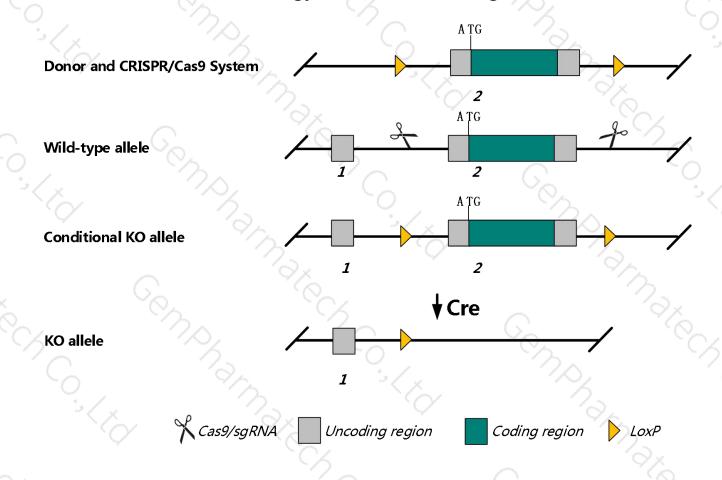
Strain background

C57BL/6JGpt

## Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Ackr3* gene. The schematic diagram is as follows:



### **Technical routes**



- The *Ackr3* gene has 2 transcripts. According to the structure of *Ackr3* gene, exon2 of *Ackr3*-201 (ENSMUST00000065587.4) transcript is recommended as the knockout region. The region contains all of the coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Ackr3* gene. The brief process is as follows: gRNA was transcribed in vitro, donor was constructed.Cas9, gRNA 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 or cell types.

### Notice



- According to the existing MGI data, Most homozygous null mutations result in perinatal lethality with cardiac defects including semilunar valve defects.
- The *Ackr3* gene is located on the Chr 1. 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 loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.

## Gene information (NCBI)



#### Ackr3 atypical chemokine receptor 3 [ Mus musculus (house mouse) ]

Gene ID: 12778, updated on 8-Oct-2019

#### Summary

☆ ?

Official Symbol Ackr3 provided by MGI

Official Full Name atypical chemokine receptor 3 provided by MGI

Primary source MGI:MGI:109562

See related Ensembl: ENSMUSG00000044337

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 Rdc1; Cxcr7; RDC-1; CXC-R7; CXCR-7; Cmkor1; AW541270

Expression Broad expression in adrenal adult (RPKM 169.3), heart adult (RPKM 116.2) and 20 other tissues See more

Orthologs human all

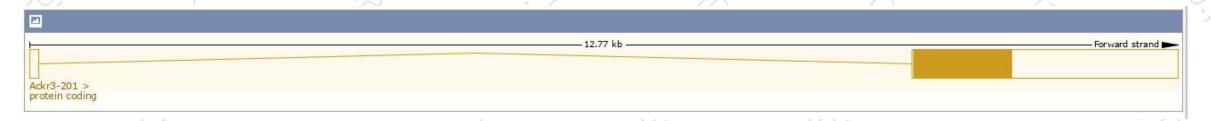
## Transcript information (Ensembl)



The gene has 2 transcripts, and all transcripts are shown below:

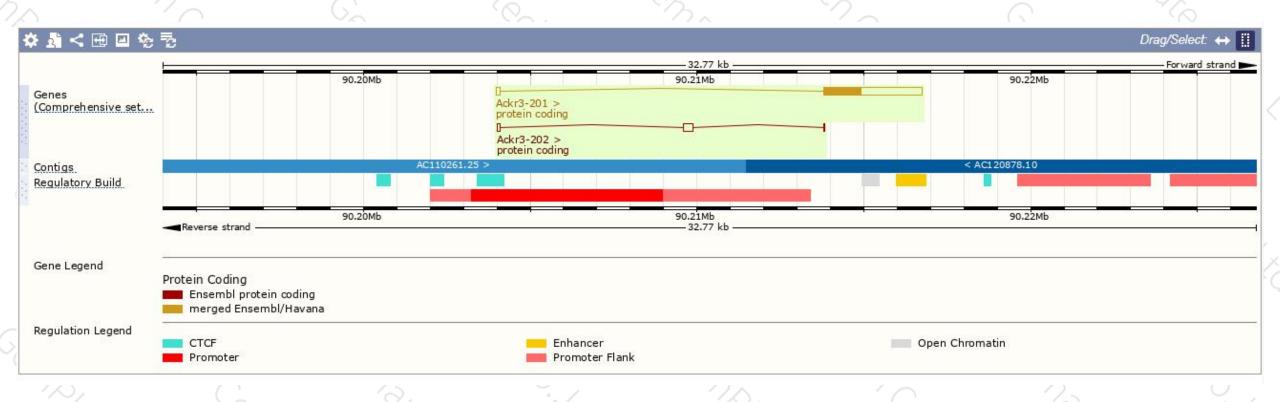
Show/hide columns (1 hidden)								Filter	
Name 🍦	Transcript ID	bp 🍦	Protein 4	Biotype	CCDS 🍦	UniProt 🍦	Flags		
Ackr3-201	ENSMUST00000065587.4	3065	<u>362aa</u>	Protein coding	CCDS15152₽	<u>P56485</u> ₽	TSL:1	GENCODE basic	APPRIS P1
Ackr3-202	ENSMUST00000159654.1	404	<u>4aa</u>	Protein coding	58	-		CDS 3' incomplete	TSL:3

The strategy is based on the design of *Ackr3*-201 transcript, The transcription is shown below



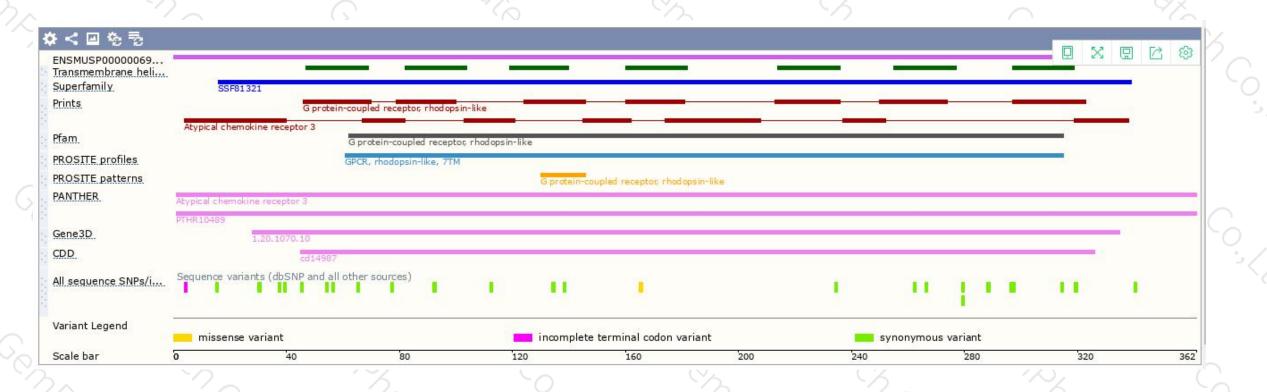
### Genomic location distribution





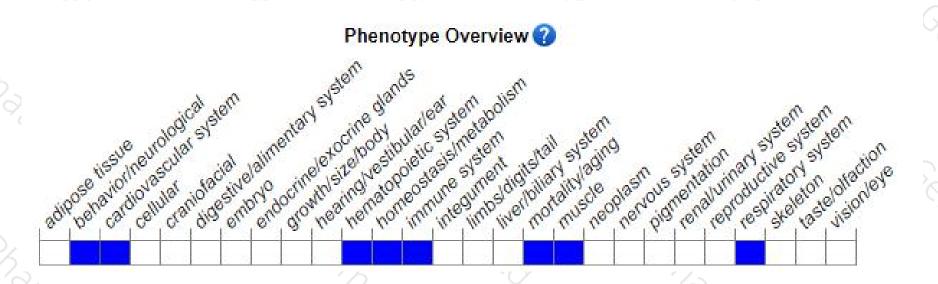
### Protein domain





## 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, Mutations in this locus affect cell-cycle regulation and apoptos is. Null homozygotes show high, early-onset tumor incidence; some have persistent hyaloid vasculature and cataracts. Truncated or temperature-sensitive alleles cause early aging phenotypes.

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





