

# Rab11a Cas9-CKO Strategy

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**Reviewer:** Huimin Su

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## **Project Overview**



**Project Name** 

Rab11a

**Project type** 

Cas9-CKO

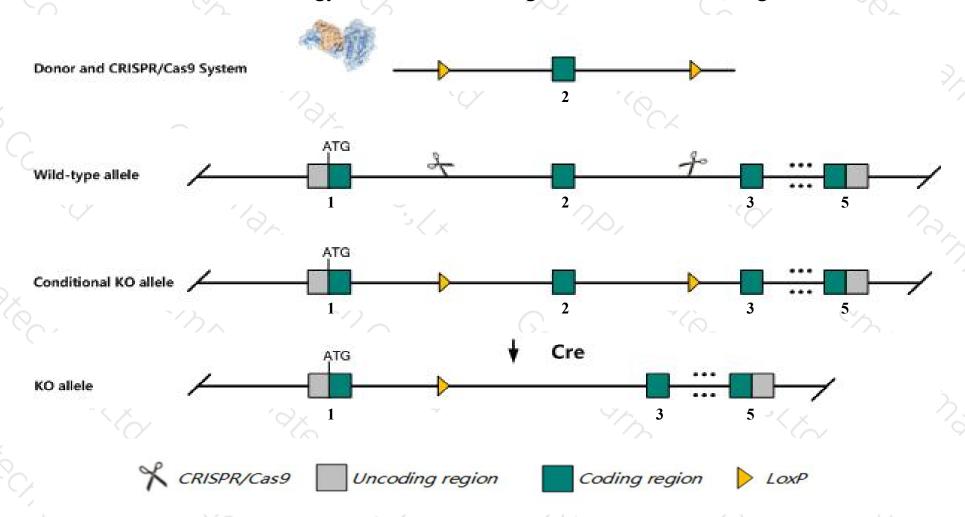
Strain background

C57BL/6JGpt

## Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Rab11a* gene. The schematic diagram is as follows:



### Technical routes



- The *Rab11a* gene has 9 transcripts. According to the structure of *Rab11a* gene, exon2 of *Rab11a-208*(ENSMUST00000172298.7) transcript is recommended as the knockout region. The region contains 196bp coding sequence.

  Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Rab11a* 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.

### **Notice**



- According to the existing MGI data, mice homozygous for a knock-out allele exhibit decreased embryo size, a rudimentary egg cylinder, failure of primitive streak formation, absent primitive node and head folds, failure to gastrulate, and complete lethality prior to organogenesis.
- > Transcript Rab11a-204,205,207 may not be affected.
- The *Rab11a* gene is located on the Chr9. 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.

## Gene information (NCBI)



#### Rab11a RAB11A, member RAS oncogene family [ Mus musculus (house mouse) ]

Gene ID: 53869, updated on 10-Dec-2019

#### Summary

☆ ?

Official Symbol Rab11a provided by MGI

Official Full Name RAB11A, member RAS oncogene family provided by MGI

Primary source MGI:MGI:1858202

See related Ensembl: ENSMUSG00000004771

Gene type protein coding
RefSeq status VALIDATED
Organism Mus musculus

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

Murinae; Mus; Mus

Expression Ubiquitous expression in CNS E18 (RPKM 50.9), bladder adult (RPKM 49.3) and 28 other tissues See more

Orthologs <u>human</u> all

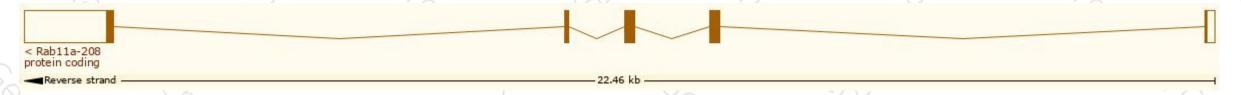
## Transcript information (Ensembl)



The gene has 9 transcripts, all transcripts are shown below:

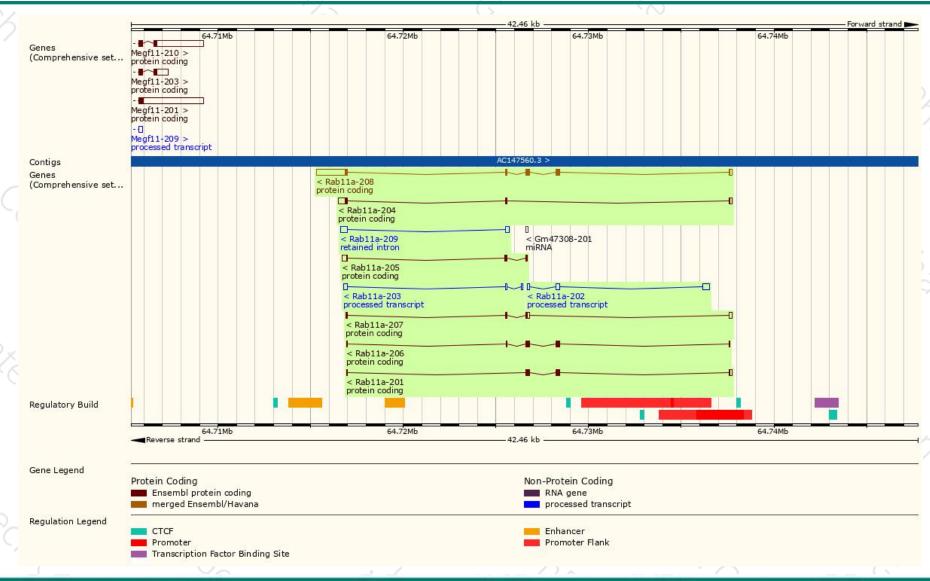
Name 🍦	Transcript ID 🍦	bp 🍦	Protein	Biotype A	CCDS	UniProt	Flags
Rab11a-209	ENSMUST00000172444.1	603	No protein	Retained intron		-	TSL:1
Rab11a-208	ENSMUST00000172298.7	2336	216aa	Protein coding	CCDS23282 ₽	P62492 & Q0PD45 &	TSL:1 GENCODE basic APPRIS P1
Rab11a-204	ENSMUST00000167569.7	726	86aa	Protein coding	(57)	E9Q6B3 ₽	TSL:3 GENCODE basic
Rab11a-201	ENSMUST00000004892.6	595	<u>155aa</u>	Protein coding	74	F8WGS1₽	CDS 3' incomplete TSL:5
Rab11a-207	ENSMUST00000171100.1	508	80aa	Protein coding	(37)	E9PZB2 ₽	CDS 3' incomplete   TSL:2
Rab11a-205	ENSMUST00000168366.1	500	<u>87aa</u>	Protein coding	74	<u>F6R2Z5</u> ₽	CDS 5' incomplete TSL:3
Rab11a-206	ENSMUST00000169058.7	462	<u>153aa</u>	Protein coding	(57)	E9Q3P9 ₽	TSL:5 GENCODE basic
Rab11a-202	ENSMUST00000165083.1	695	No protein	Processed transcript	754	-	TSL:5
Rab11a-203	ENSMUST00000166857.1	397	No protein	Processed transcript	878		TSL:3

The strategy is based on the design of *Rab11a-208* transcript, the transcription is shown below:



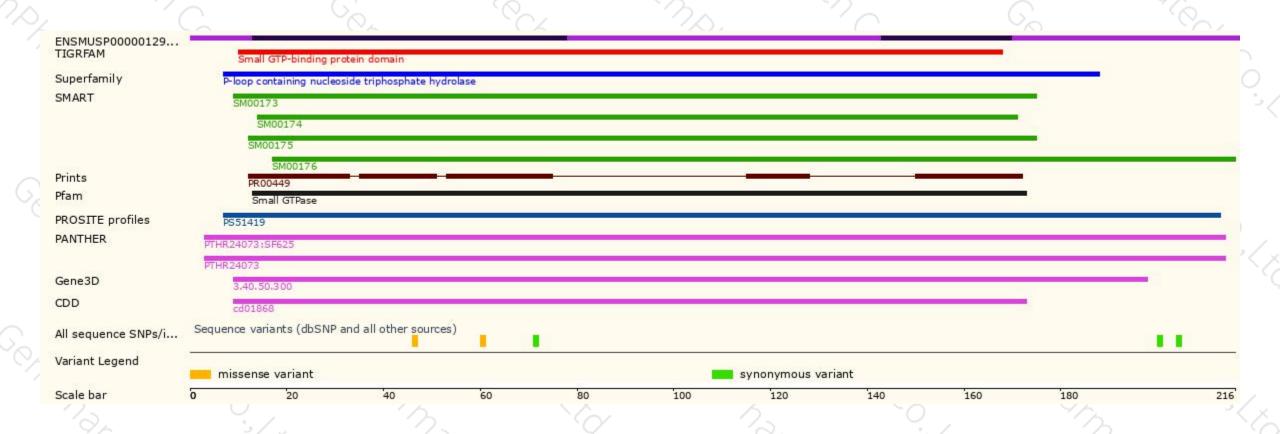
### Genomic location distribution





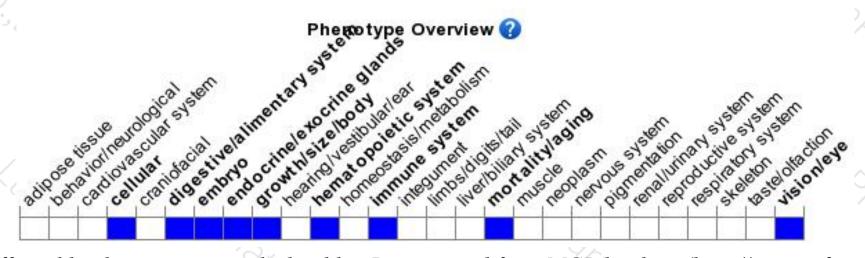
### Protein domain





## 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 knock-out allele exhibit decreased embryo size, a rudimentary egg cylinder, failure of primitive streak formation, absent primitive node and head folds, failure to gastrulate, and complete lethality prior to organogenesis.



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





