

# Prkg2 Cas9-CKO Strategy

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Reviewer: Ruirui Zhang

**Design Date:** 2019/8/29

## **Project Overview**



**Project Name** 

Prkg2

**Project type** 

Cas9-CKO

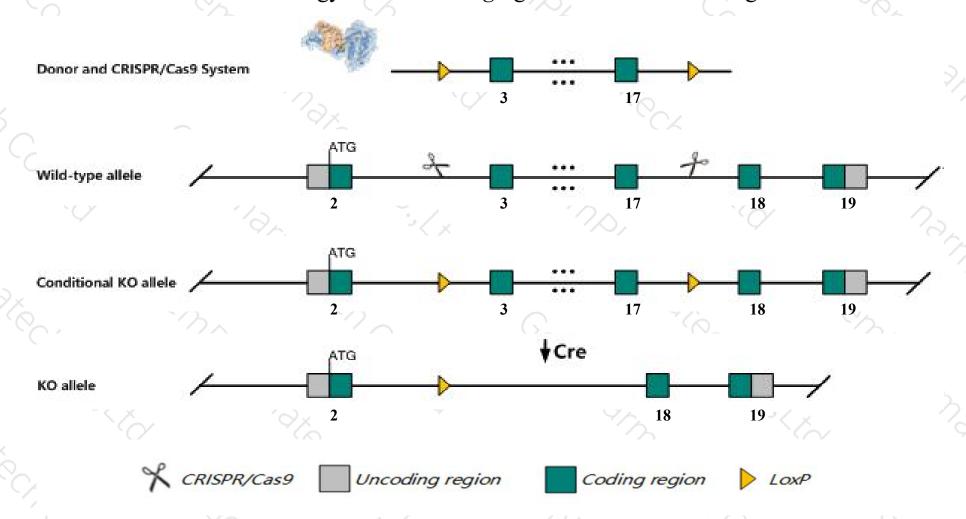
Strain background

C57BL/6JGpt

### Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Prkg2* gene. The schematic diagram is as follows:



### Technical routes



- The *Prkg2* gene has 5 transcripts. According to the structure of *Prkg2* gene, exon3-exon17 of *Prkg2-203* (ENSMUST00000161490.7) transcript is recommended as the knockout region. The region contains most 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 *Prkg2* 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, Homozygous null mice exhibit dwarfism, with abnormal skull morphology and short limbs and vertebrae. Defects in axial organization of the growth plates was evident as mice aged.

  Digestive secretion in response to enterotoxin was reduced.
- > The KO region deletes most of the coding sequence, but does not result in frameshift.
- > The *Prkg2* gene is located on the Chr5. 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)



#### Prkg2 protein kinase, cGMP-dependent, type II [ Mus musculus (house mouse) ]

Gene ID: 19092, updated on 27-Aug-2019

#### Summary

☆ ?

Official Symbol Prkg2 provided by MGI

Official Full Name protein kinase, cGMP-dependent, type II provided by MGI

Primary source MGI:MGI:108173

See related Ensembl: ENSMUSG00000029334

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

Also known as CGKII; Prkgr2; cGK-II; AW212535

Expression Broad expression in small intestine adult (RPKM 2.2), frontal lobe adult (RPKM 1.5) and 16 other tissues See more

Orthologs human all

## Transcript information (Ensembl)



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

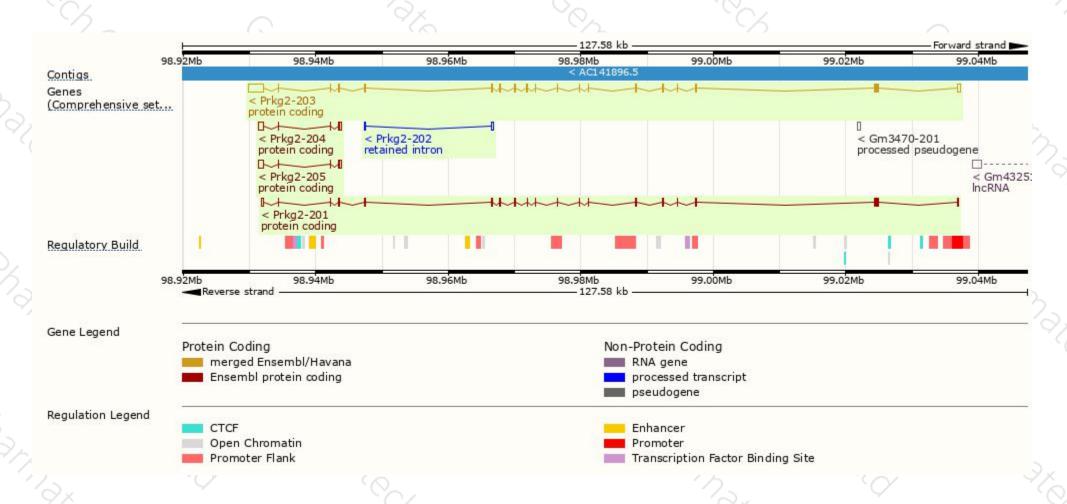
Name	Transcript ID	bp 🌲	Protein 🍦	Biotype #	CCDS .	UniProt 🝦	Flags
Prkg2-203	ENSMUST00000161490.7	4968	762aa	Protein coding	CCDS19459₽	Q8CAH8₽	TSL:1 APPRIS P1
Prkg2-201	ENSMUST00000031277.6	2618	733aa	Protein coding	7	E9QPH0₽	TSL:5 GENCODE basic
Prkg2-204	ENSMUST00000162147.5	1389	107aa	Protein coding		Q3TMJ2₽	TSL:1 GENCODE basic
Prkg2-205	ENSMUST00000162619.7	1308	<u>94aa</u>	Protein coding	9.7	A0A0G2JEE8₽	TSL:5 GENCODE basic
Prkg2-202	ENSMUST00000160765.1	387	No protein	Retained intron	-	65	TSL:3

The strategy is based on the design of *Prkg2-203* transcript, The transcription is shown below



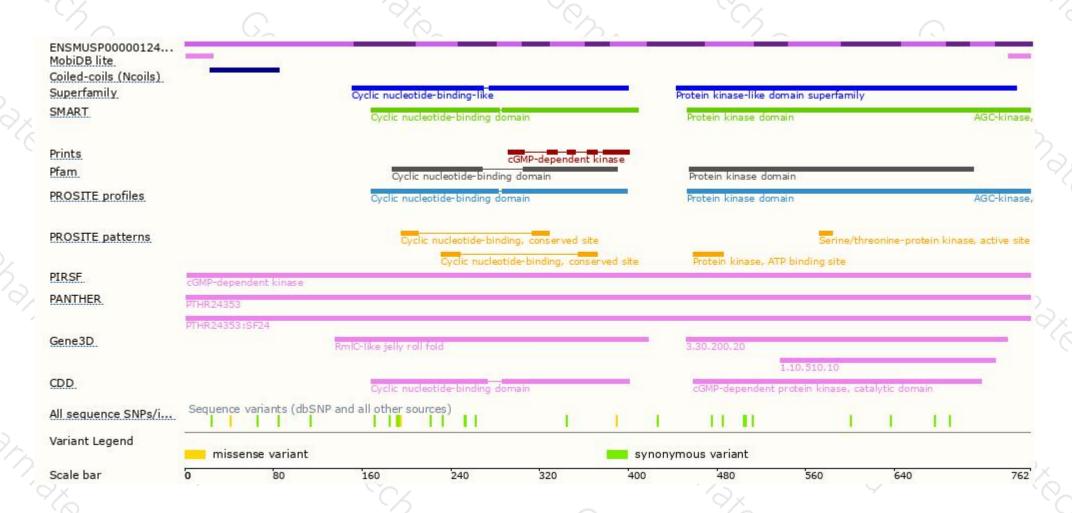
### Genomic location distribution





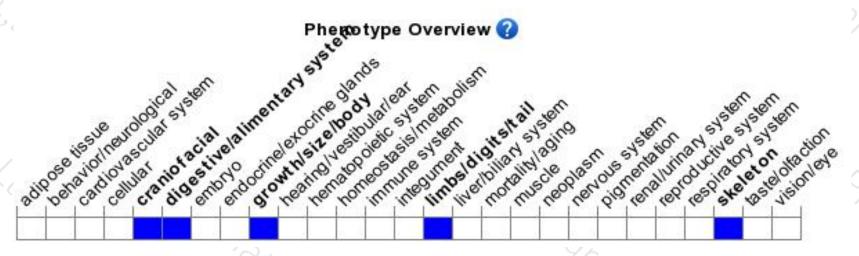
### Protein domain





## 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 null mice exhibit dwarfism, with abnormal skull morphology and short limbs and vertebrae. Defects in axial organization of the growth plates was evident as mice aged. Digestive secretion in response to enterotoxin was reduced.



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





