

Prph2 Cas9-CKO Strategy

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Reviewer:

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Project Overview

Project Name

Prph2

Project type

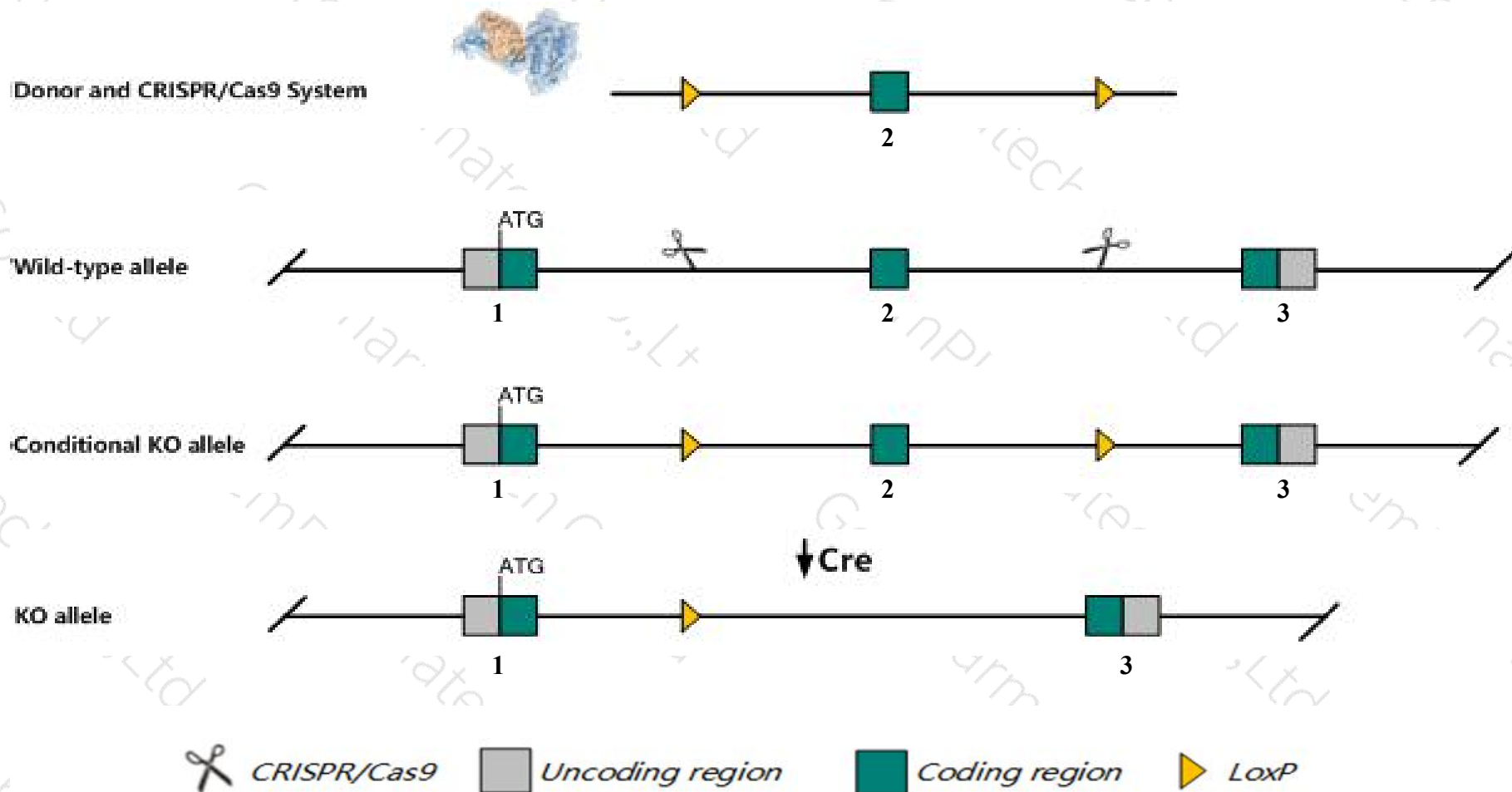
Cas9-CKO

Strain background

C57BL/6JGpt

Conditional Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Prph2* gene. The schematic diagram is as follows:



Technical routes

- The *Prph2* gene has 2 transcripts. According to the structure of *Prph2* gene, exon2 of *Prph2-201* (ENSMUST00000024773.5) transcript is recommended as the knockout region. The region contains 247bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Prph2* 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.

- According to the existing MGI data, Mice homozygous for a spontaneous mutation display slow retinal degeneration with thinning and loss of the outer nuclear layer, loss of photoreceptor outer segments, and increased numbers of Muller cells. Heterozygous mice also display retinal degeneration and Muller cell gliosis.
- The *Prph2* gene is located on the Chr17. 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)

Prph2 peripherin 2 [Mus musculus (house mouse)]

Gene ID: 19133, updated on 31-Jan-2019

Summary



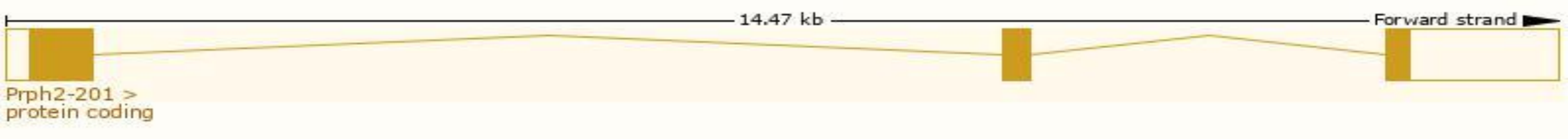
Official Symbol	Prph2 provided by MGI
Official Full Name	peripherin 2 provided by MGI
Primary source	MGI:MGI:102791
See related	Ensembl:ENSMUSG00000023978
Gene type	protein coding
RefSeq status	VALIDATED
Organism	Mus musculus
Lineage	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
Also known as	AOFMD, AVMD, Nmf193, PRPH, RP7, Rd-2, Rd2, Rds, Tspan22, rds
Expression	Restricted expression toward testis adult (RPKM 1.3) See more
Orthologs	human all

Transcript information (Ensembl)

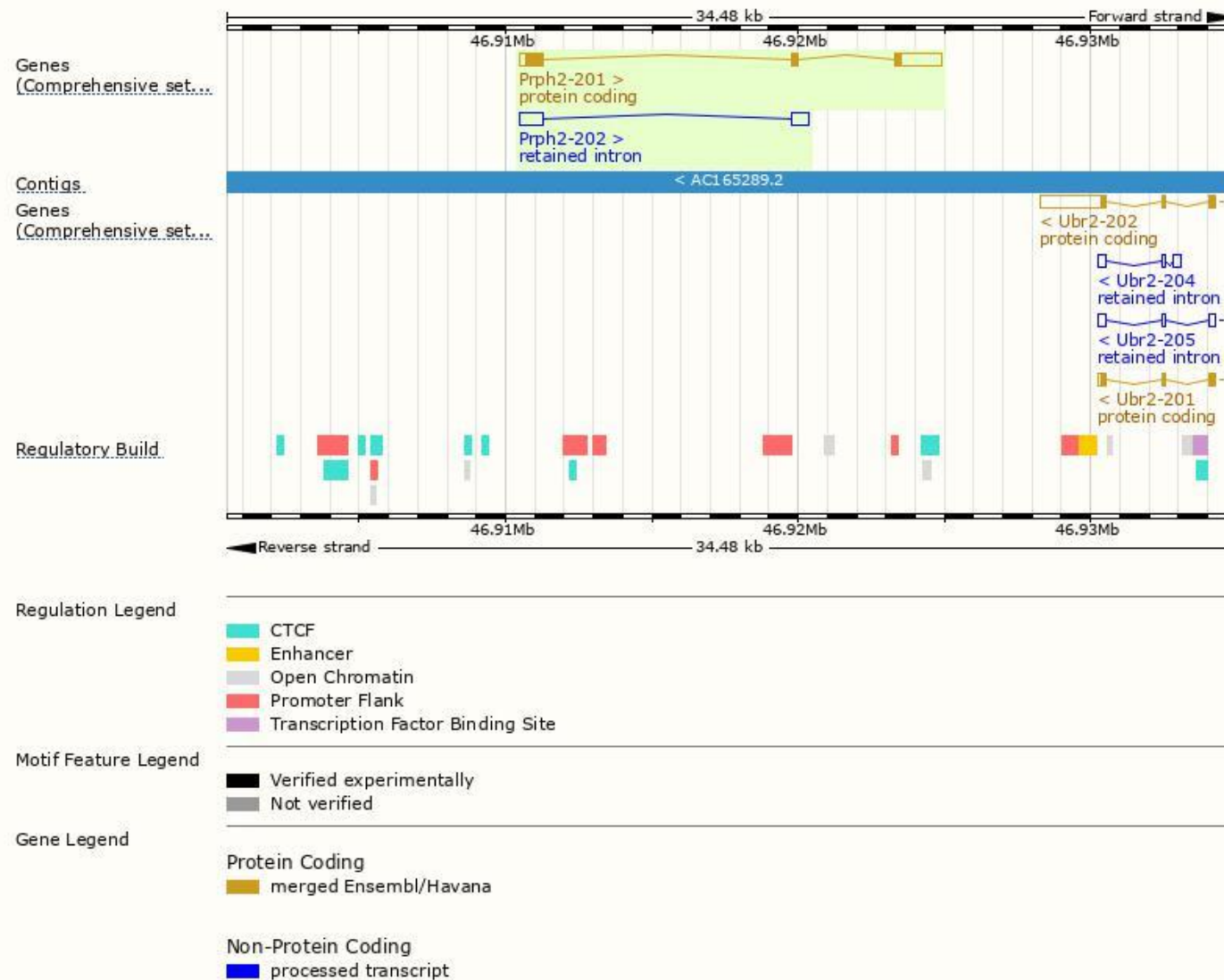
The gene has 2 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Prph2-201	ENSMUST00000024773.5	2665	346aa	Protein coding	CCDS28844	P15499 Q3UWK3	TSL:1 GENCODE basic APPRIS P1
Prph2-202	ENSMUST00000162469.1	1384	No protein	Retained intron	-	-	TSL:1

The strategy is based on the design of *Prph2-201* transcript,The transcription is shown below



Genomic location distribution



Protein domain

ENSMUSP00000024...

Transmembrane heli...

Low complexity (Seq)

Conserved Domains

hmmpanther

PTHR19282

PTHR19282;SF202

Superfamily domains

Prints domain

Pfam domain

Tetraspanin, EC2 domain superfamily

Peripherin/rom-1

Tetraspanin/ Peripherin

PROSITE patterns

Gene3D

Tetraspanin, EC2 domain superfamily

All sequence SNPs/i....

Sequence variants (dbSNP and all other sources)

Variant Legend

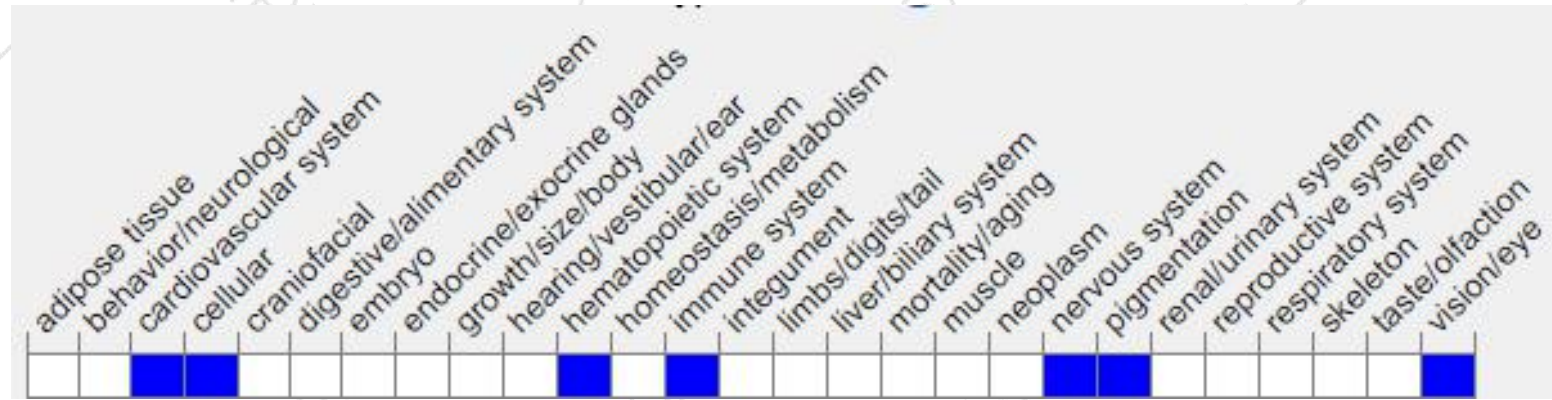
missense variant

synonymous variant

Scale bar

0 40 80 120 160 200 240 280 346

Mouse phenotype description(MGI)



Phenotypes affected by the gene are marked in blue. Data quoted from MGI database(<http://www.informatics.jax.org/>).

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If you have any questions, you are welcome to inquire.

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