

Phf20 Cas9-CKO Strategy

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Project Overview



Project Name

Phf20

Project type

Cas9-CKO

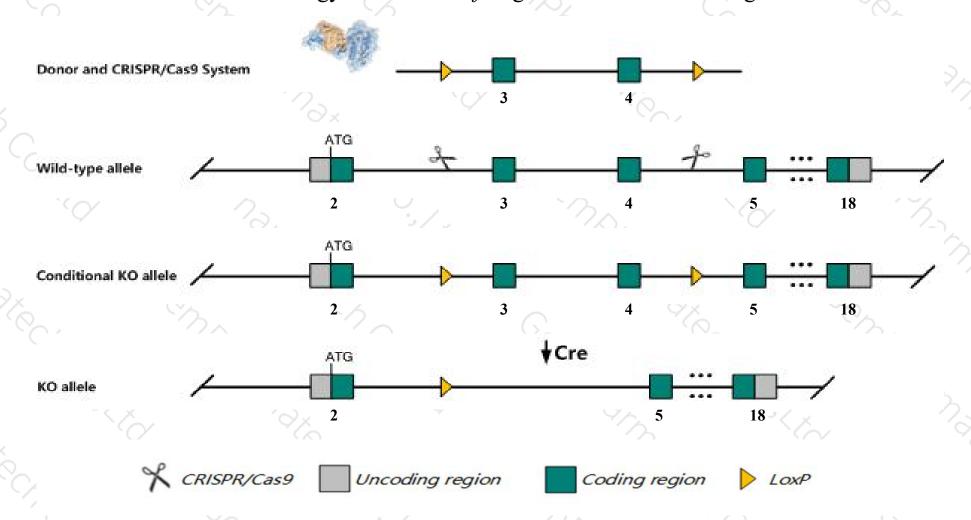
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Phf20* gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Phf20* gene has 11 transcripts. According to the structure of *Phf20* gene, exon3-exon4 of *Phf20*201(ENSMUST00000037401.9) transcript is recommended as the knockout region. The region contains 257bp coding sequence.

 Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Phf20* 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 neonatal lethality, decreased body size and total body fat amount, and abnormal skeletal and hematopoietic development.
- ➤ Transcript 207 CDS 3' incomplete the influences is unknown.
- > The *Phf20* gene is located on the Chr2. 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)



Phf20 PHD finger protein 20 [Mus musculus (house mouse)]

Gene ID: 228829, updated on 13-Mar-2020

Summary

☆ ?

Official Symbol Phf20 provided by MGI

Official Full Name PHD finger protein 20 provided by MGI

Primary source MGI:MGI:2444148

See related Ensembl:ENSMUSG00000038116

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 6820402020Rik

Expression Ubiquitous expression in cerebellum adult (RPKM 9.8), CNS E18 (RPKM 9.6) and 28 other tissuesSee more

Orthologs <u>human all</u>

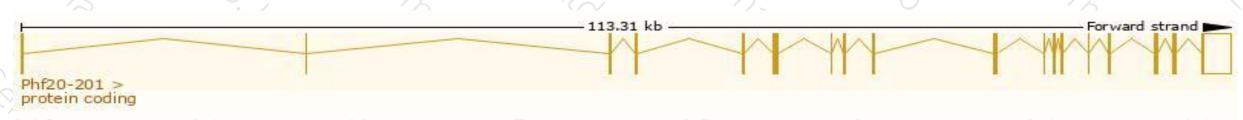
Transcript information (Ensembl)



The gene has 11 transcripts, all transcripts are shown below:

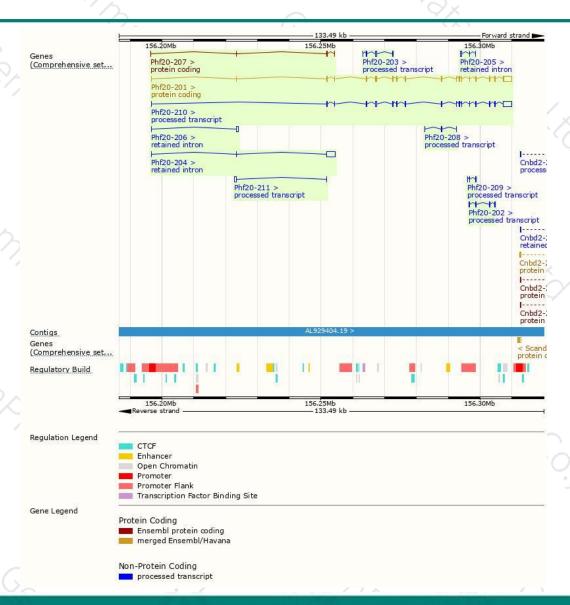
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Phf20-201	ENSMUST00000037401.9	5737	1010aa	Protein coding	CCDS38297	Q8BLG0	TSL:1 GENCODE basic APPRIS P1
Phf20-207	ENSMUST00000145731.7	453	<u>113aa</u>	Protein coding	2	A2AV92	CDS 3' incomplete TSL:3
Phf20-210	ENSMUST00000152617.7	5524	No protein	Processed transcript	2	2	TSL:5
Phf20-202	ENSMUST00000122814.1	903	No protein	Processed transcript	70	-	TSL:5
Phf20-203	ENSMUST00000131479.1	605	No protein	Processed transcript	2	-	TSL:3
Phf20-211	ENSMUST00000153822.1	605	No protein	Processed transcript	-	-	TSL:2
Phf20-208	ENSMUST00000147234.1	377	No protein	Processed transcript	-	-	TSL:3
Phf20-209	ENSMUST00000147242.7	368	No protein	Processed transcript	2	-	TSL:5
Phf20-204	ENSMUST00000132129.1	2896	No protein	Retained intron	5	-	TSL:1
Phf20-205	ENSMUST00000132352.1	703	No protein	Retained intron	-	-	TSL:5
Phf20-206	ENSMUST00000135796.1	702	No protein	Retained intron	2	12	TSL:3

The strategy is based on the design of *Phf20-201* transcript, the transcription is shown below:



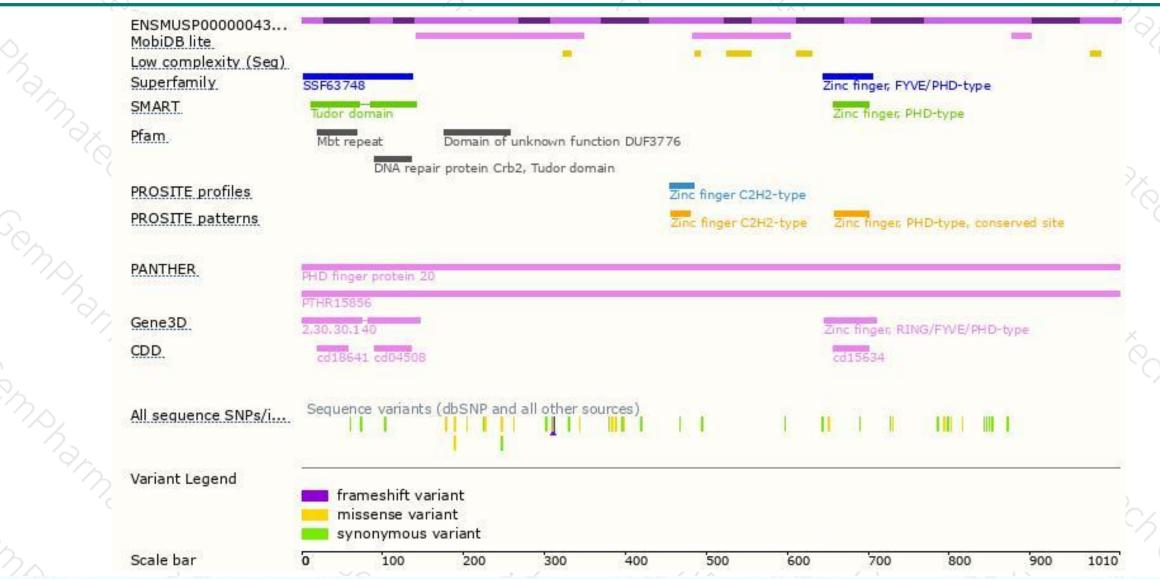
Genomic location distribution





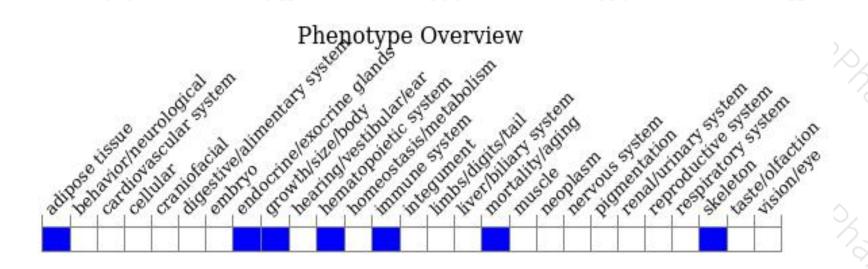
Protein domain





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 knock-out allele exhibit neonatal lethality, decreased body size and total body fat amount, and abnormal skeletal and hematopoietic development.



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





