

Gprc6a Cas9-KO Strategy

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Project Overview



Project Name

Gprc6a

Project type

Cas9-KO

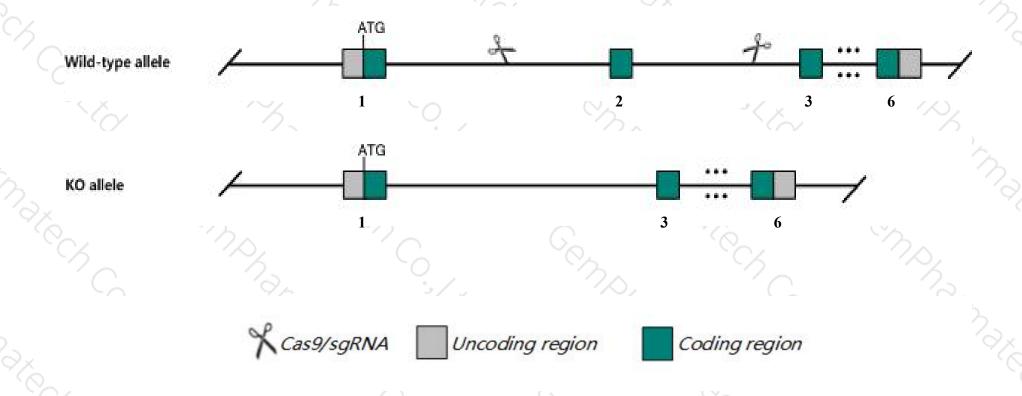
Strain background

C57BL/6J

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Gprc6a* gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Gprc6a* gene has 3 transcripts. According to the structure of *Gprc6a* gene, exon2 of *Gprc6a-201*(ENSMUST00000020062.3) transcript is recommended as the knockout region. The region contains 304bp coding sequence.

 Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Gprc6a* gene. The brief process is as follows: sgRNA was transcribed in vitro.Cas9 and sgRNA were microinjected into the fertilized eggs of C57BL/6J 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/6J mice.

Notice



- ➤ According to the existing MGI data, Mice homozygous for a knock-out allele show a metabolic syndrome characterized by impaired bone mineralization, increased fat mass, abnormal renal handling of calcium and phosphorus, fatty liver, glucose intolerance, testicular feminization and abnormal steroidogenesis.
- ➤ The *Gprc6a* gene is located on the Chr10. 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 gene transcription and translation processes, all risks cannot be predicted under existing information.

Gene information (NCBI)



Gprc6a G protein-coupled receptor, family C, group 6, member A [Mus musculus (house mouse)]

Gene ID: 210198, updated on 5-Mar-2019

Summary

↑ ?

Official Symbol Gprc6a provided by MGI

Official Full Name G protein-coupled receptor, family C, group 6, member A provided by MGI

Primary source MGI:MGI:2429498

See related Ensembl:ENSMUSG00000019905

Gene type protein coding
RefSeq status PROVISIONAL
Organism Mus musculus

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

Muroidea; Muridae; Murinae; Mus; Mus

Expression Low expression observed in reference datasetSee more

Orthologs <u>human</u> all

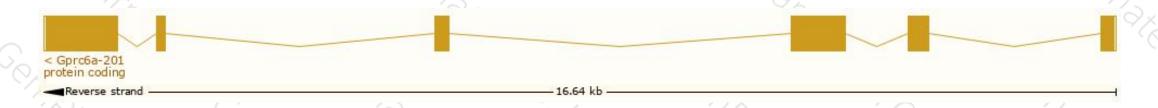
Transcript information (Ensembl)



The gene has 3 transcript, all transcripts are shown below:

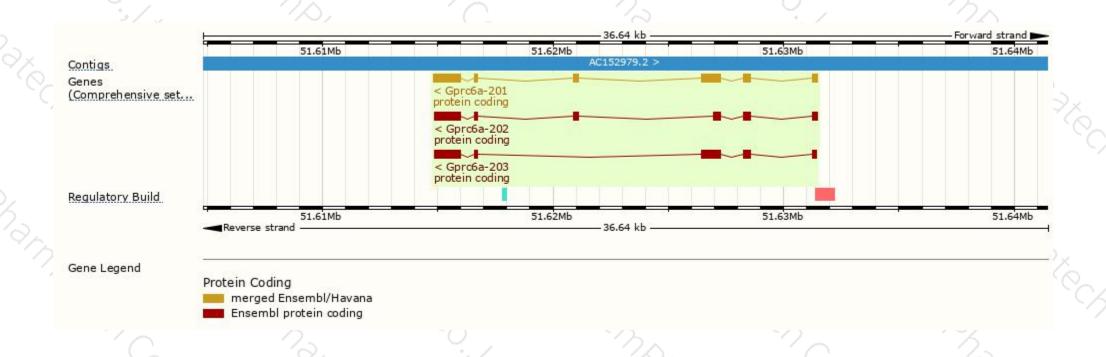
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Gprc6a-201	ENSMUST00000020062.3	2856	928aa	Protein coding	CCDS23834	Q8K4Z6	TSL:1 GENCODE basic APPRIS P2
Gprc6a-203	ENSMUST00000219286.1	2574	<u>857aa</u>	Protein coding	8-	Q8K4Z6	TSL:1 GENCODE basic
Gprc6a-202	ENSMUST00000218684.1	2292	<u>753aa</u>	Protein coding	-	Q8K4Z6	TSL:1 GENCODE basic APPRIS ALT2

The strategy is based on the design of *Gprc6a-201* transcript, the transcription is shown below



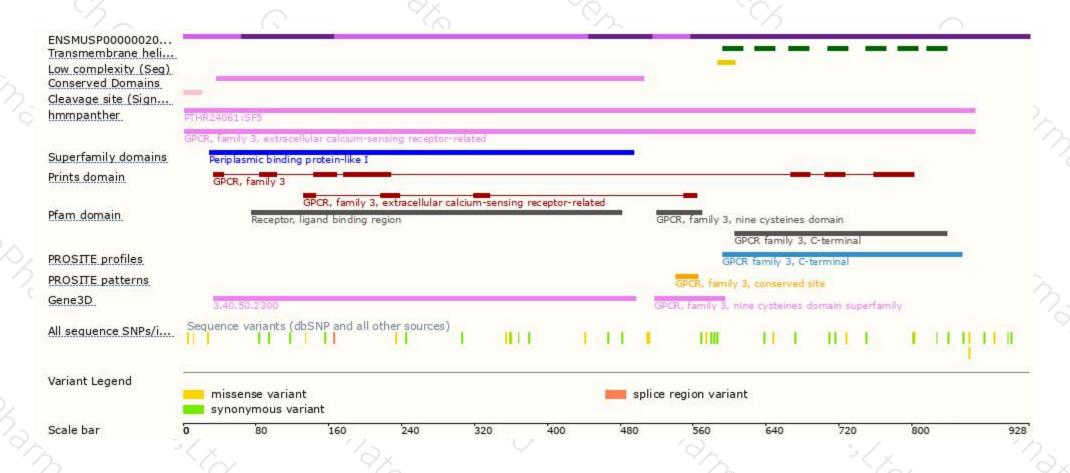
Genomic location distribution





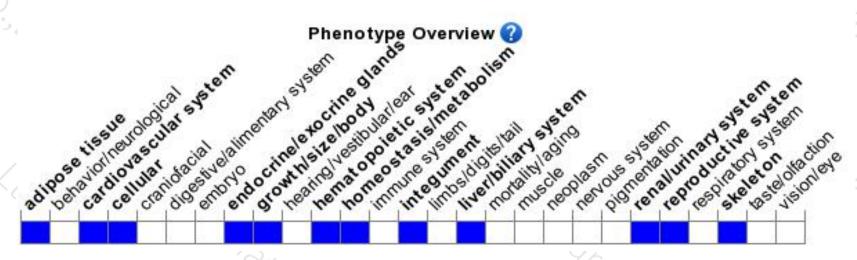
Protein domain





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 show a metabolic syndrome characterized by impaired bone mineralization, increased fat mass, abnormal renal handling of calcium and phosphorus, fatty liver, glucose intolerance, testicular feminization and abnormal steroidogenesis.



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

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