

Cas9-CKO Strategy Compland and College

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Project Overview



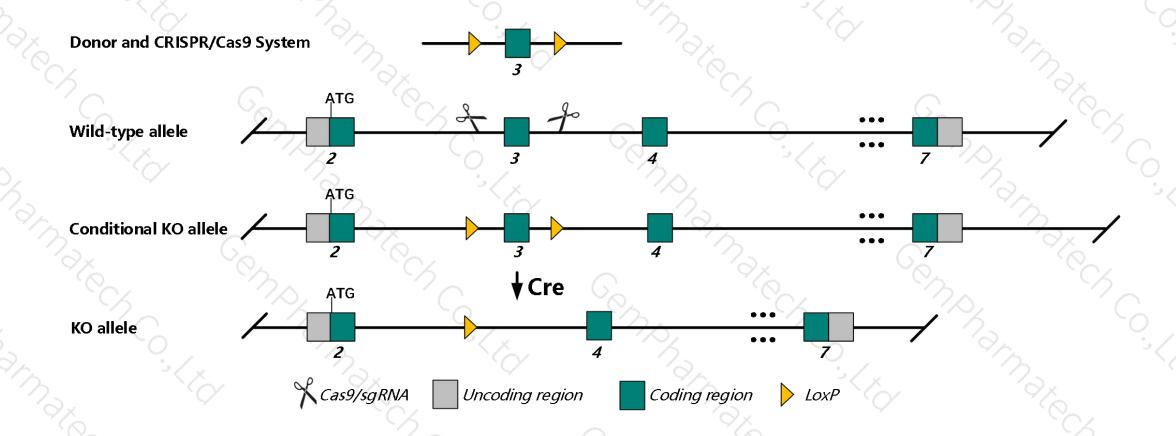
Project Name Ogn

Project type Cas9-CKO

Strain background C57BL/6JGpt

Conditional Knockout strategy

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



Technical routes



- The *Ogn* gene has 1 transcripts. According to the structure of *Ogn* gene, exon3 of *Ogn-201* (
 ENSMUST00000021822.6) transcript is recommended as the knockout region. The region contains 94bp coding sequence.

 Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Ogn* gene. The brief process is as follows:gRNA was transcribed in vitro, donor was constructed.Cas9, gRNA 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 disruption in this gene display reduced collagen fiber density and organization, as well as more variability in fibrilar diameter in both the skin and the cornea. Corneal clarity was unaffected whereas skin tensile strength was reduced.
- ➤ This gene was overlapped with the intron of *Cenpp* gene.
- The *Ogn* gene is located on the Chr13. If the conditional 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)



Ogn osteoglycin [Mus musculus (house mouse)]

Gene ID: 18295, updated on 18-Apr-2019

Summary

☆ ?

Official Symbol Ogn provided by MGI

Official Full Name osteoglycin provided by MGI

Primary source MGI:MGI:109278

See related Ensembl: ENSMUSG00000021390

Gene type protein coding
RefSeq status VALIDATED
Organism Mus musculus

Organism Mus musculus

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

Murinae; Mus; Mus

Also known as OG; OIF; SLRR3A; mimecan; mimican; 3110079A16Rik

Expression Broad expression in bladder adult (RPKM 41.4), limb E14.5 (RPKM 40.9) and 15 other tissues See more

Orthologs human all

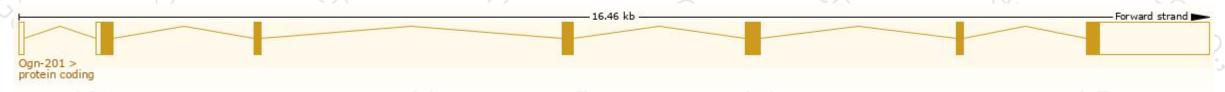
Transcript information (Ensembl)



The gene has 1 transcripts, all transcripts are shown below:

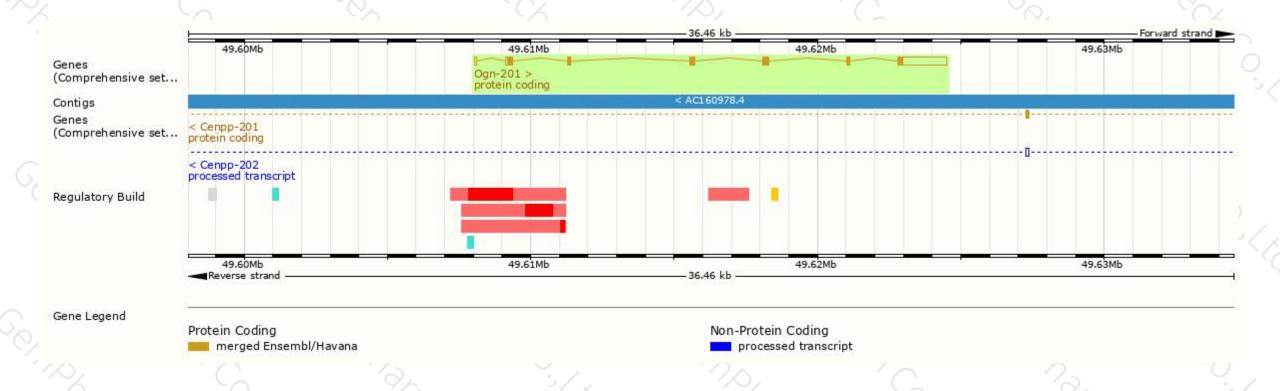
Name	Transcript ID	bp 🛊	Protein	Biotype	CCDS	UniProt	Flags		
Ogn-201	ENSMUST00000021822.6	2566	298aa	Protein coding	CCDS26505₺	Q543C5&Q62000&	TSL:1	GENCODE basic	APPRIS P1

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



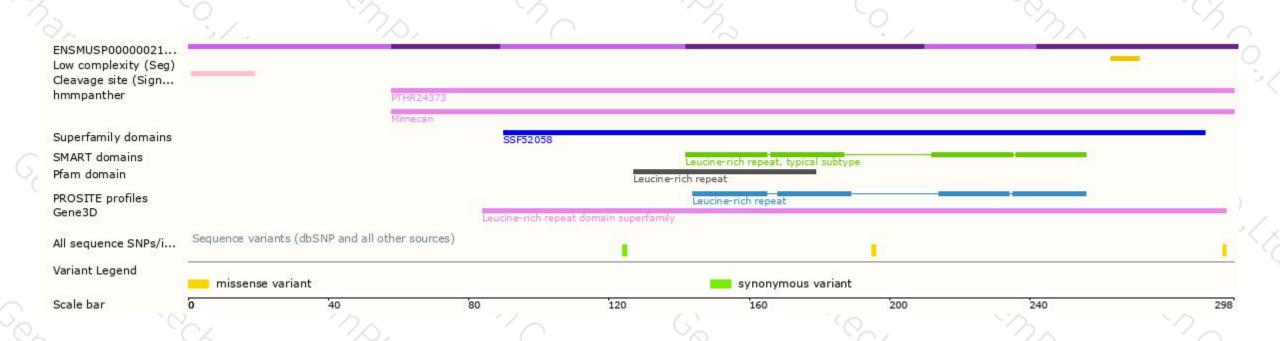
Genomic location distribution





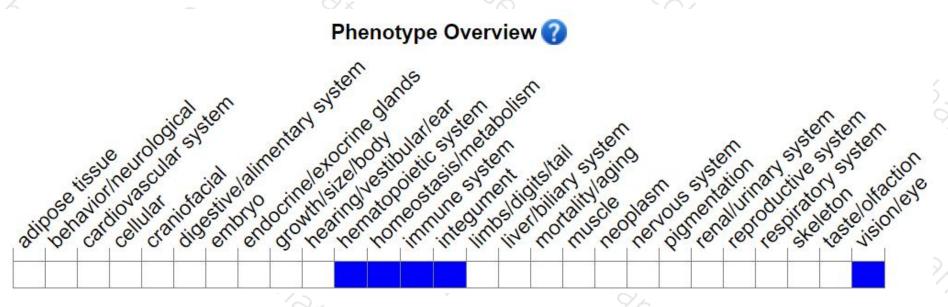
Protein domain





Mouse phenotype description(MGI)





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

Mice homozygous for a disruption in this gene display reduced collagen fiber density and organization, as well as more variability in fibrilar diameter in both the skin and the cornea. Corneal clarity was unaffected whereas skin tensile strength was reduced.



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





