

# Bmp7 Cas9-CKO Strategy

Designer: Yanhua Shen

Reviewer:Xueting Zhang

Design Date:2019-09-03

# **Project Overview**



**Project Name** 

Bmp7

**Project type** 

Cas9-CKO

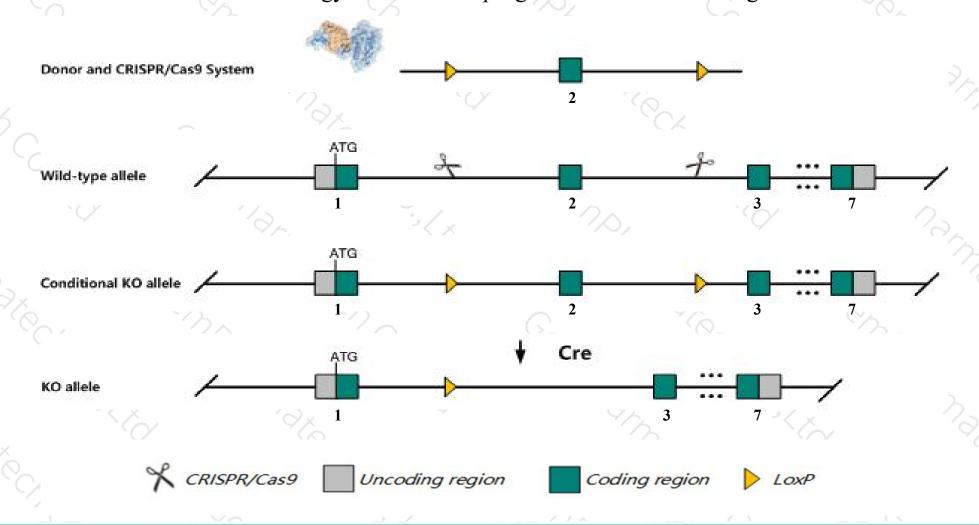
Strain background

C57BL/6JGpt

## Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Bmp7* gene. The schematic diagram is as follows:



### Technical routes



- The *Bmp7* gene has 2 transcripts. According to the structure of *Bmp7* gene, exon2 of *Bmp7-201*(ENSMUST0000009143.7) transcript is recommended as the knockout region. The region contains 193bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Bmp7* 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, Various homozygous targeted mutations result in postnatal lethality, a wide range of skeletal and cartilage abnormalities, renal dysplasia and polycystic kidney, and eye defects.
- > Some amino acids will remain at the N-terminus and some functions may be retained.
- > The *Bmp7* 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)



#### Bmp7 bone morphogenetic protein 7 [ Mus musculus (house mouse) ]

Gene ID: 12162, updated on 27-Aug-2019

Summary

↑ ?

Official Symbol Bmp7 provided by MGI

Official Full Name bone morphogenetic protein 7 provided by MGI

Primary source MGI:MGI:103302

See related Ensembl: ENSMUSG00000008999

Gene type protein coding
RefSeq status REVIEWED
Organism Mus musculus

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

Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus

Also known as OP1

Summary This gene encodes a secreted ligand of the TGF-beta (transforming growth factor-beta) superfamily of proteins. Ligands of this family

bind various TGF-beta receptors leading to recruitment and activation of SMAD family transcription factors that regulate gene

expression. The encoded preproprotein is proteolytically processed to generate each subunit of the disulfide-linked homodimer. Mutation

of this gene results in skeletal, kidney, and other developmental defects. [provided by RefSeq, Jul 2016]

Expression Broad expression in adrenal adult (RPKM 31.1), kidney adult (RPKM 17.3) and 20 other tissues See more

Orthologs human all

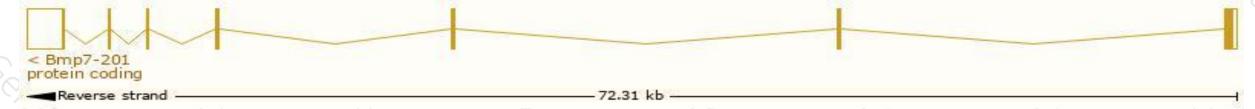
# Transcript information (Ensembl)



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

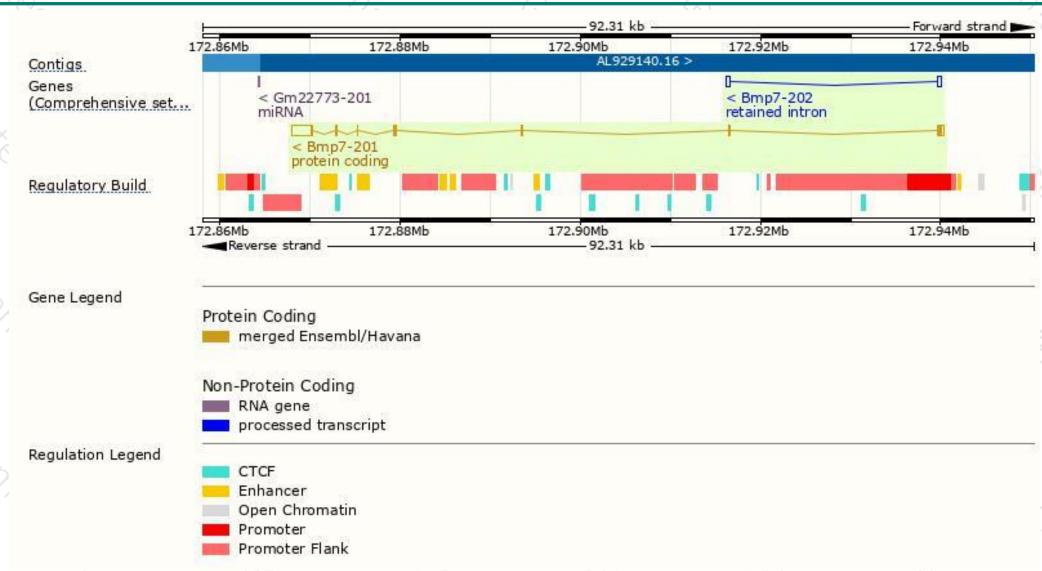
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Bmp7-201	ENSMUST00000009143.7	3670	<u>430aa</u>	Protein coding	CCDS17136	P23359	TSL:1 GENCODE basic APPRIS P1
Bmp7-202	ENSMUST00000137247.1	980	No protein	Retained intron	-	-	TSL:2

The strategy is based on the design of *Bmp7-201* transcript, The transcription is shown below



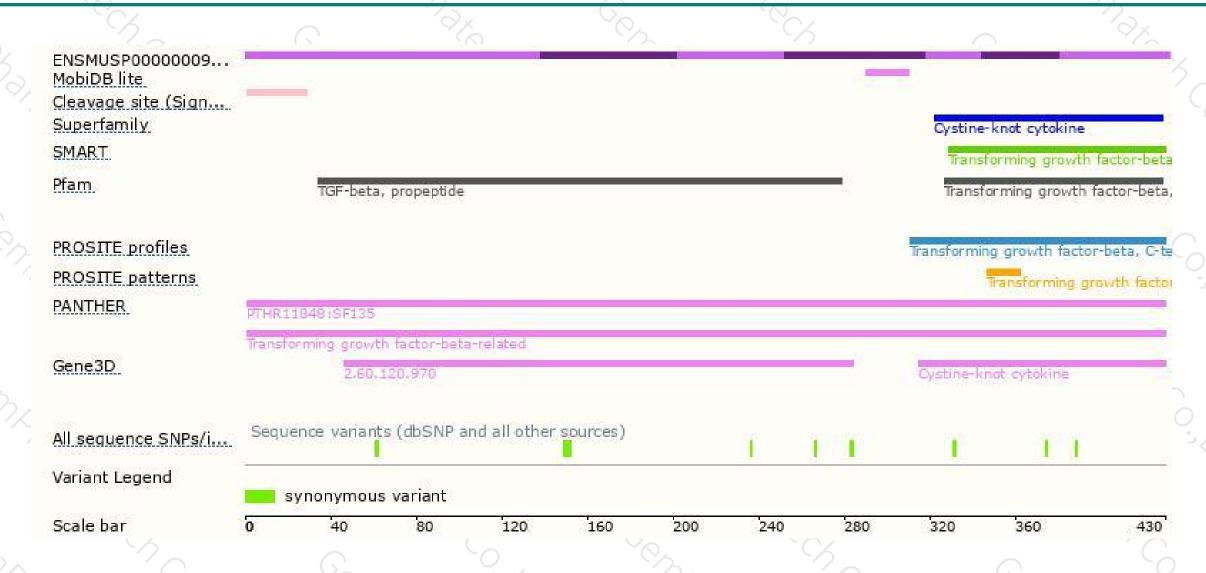
### Genomic location distribution





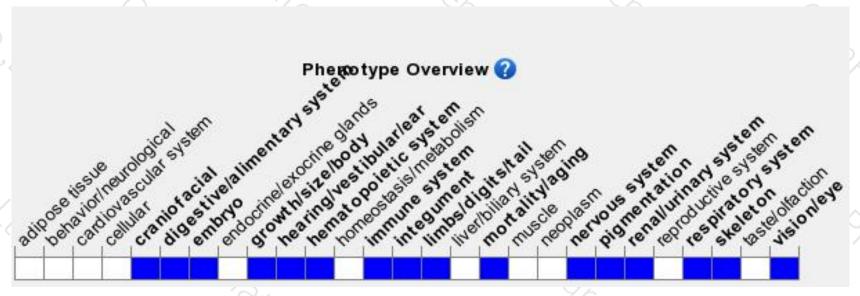
### Protein domain





# 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, Various homozygous targeted mutations result in postnatal lethality, a wide range of skeletal and cartilage abnormalities, renal dysplasia and polycystic kidney, and eye defects.



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





