

# Bmp1 Cas9-KO Strategy

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Reviewer: JiaYu

## **Project Overview**



**Project Name** 

Bmp1

**Project type** 

Cas9-KO

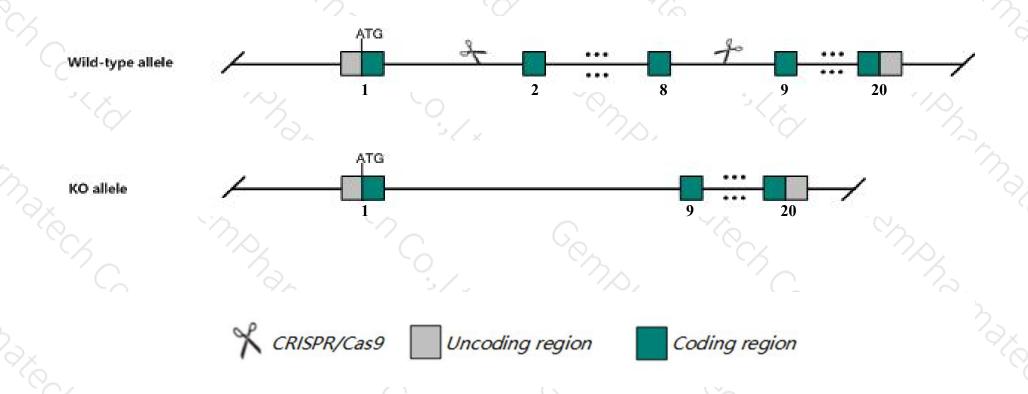
Strain background

C57BL/6JGpt

## **Knockout strategy**



This model will use CRISPR/Cas9 technology to edit the *Bmp1* gene. The schematic diagram is as follows:



## **Technical routes**



- ➤ The *Bmp1* gene has 7 transcripts. According to the structure of *Bmp1* gene, exon2-exon8 of *Bmp1-201*(ENSMUST00000022693.8) transcript is recommended as the knockout region. The region contains 923bp coding sequence.

  Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Bmp1* gene. The brief process is as follows: CRISPR/Cas9 system

### **Notice**



- ➤ According to the existing MGI data, Homozygous targeted mutant embryos have reduced ossification of the skull, persistent herniation of the gut, abnormal collagen fibrils in the amnion, and die at birth.
- The *Bmp1* gene is located on the Chr14. 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 the gene knockout on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.

## Gene information (NCBI)



#### Bmp1 bone morphogenetic protein 1 [Mus musculus (house mouse)]

Gene ID: 12153, updated on 12-Mar-2019

#### Summary

☆ ?

Official Symbol Bmp1 provided by MGI

Official Full Name bone morphogenetic protein 1 provided by MGI

Primary source MGI:MGI:88176

See related Ensembl: ENSMUSG00000022098

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 Pcp, Tld

Summary This gene encodes a metalloproteinase that plays an essential role in the formation of the extracellular matrix and is also able to induce

ectopic bone formation. Unlike other bone morphogenetic proteins, the protein encoded by this gene is not closely related to transforming growth factor-beta. This protein plays in role several developmental processes. In humans, mutations in this gene are associated with osteogenesis imperfecta and with increased bone mineral density and multiple recurrent fractures. Alternative splicing results in multiple

transcript variants. [provided by RefSeq, Dec 2013]

Expression Ubiquitous expression in limb E14.5 (RPKM 81.0), ovary adult (RPKM 50.8) and 25 other tissuesSee more

Orthologs <u>human</u> all

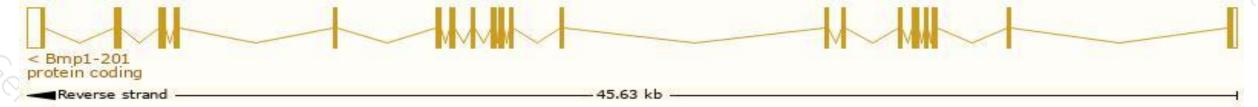
## Transcript information (Ensembl)



The gene has 7 transcripts, all transcripts are shown below:

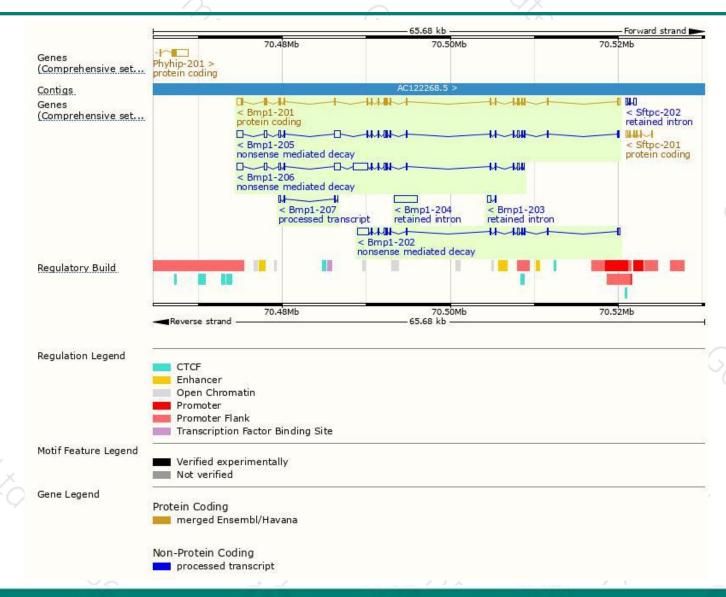
		7			F		[-
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Bmp1-201	ENSMUST00000022693.8	3684	<u>991aa</u>	Protein coding	CCDS36972	P98063	TSL:1 GENCODE basic APPRIS P1
Bmp1-206	ENSMUST00000227944.1	5365	<u>57aa</u>	Nonsense mediated decay	698	A0A2I3BQB3	CDS 5' incomplete
Bmp1-205	ENSMUST00000226906.1	4340	<u>194aa</u>	Nonsense mediated decay	140	A0A2I3BRJ8	
Bmp1-202	ENSMUST00000226246.1	3472	<u>194aa</u>	Nonsense mediated decay	V-50	A0A2I3BRJ8	
Bmp1-207	ENSMUST00000228501.1	507	No protein	Processed transcript	1783	5	
Bmp1-204	ENSMUST00000226601.1	2736	No protein	Retained intron	677	- 8	
Bmp1-203	ENSMUST00000226539.1	452	No protein	Retained intron		20	

The strategy is based on the design of *Bmp1-201* transcript, The transcription is shown below



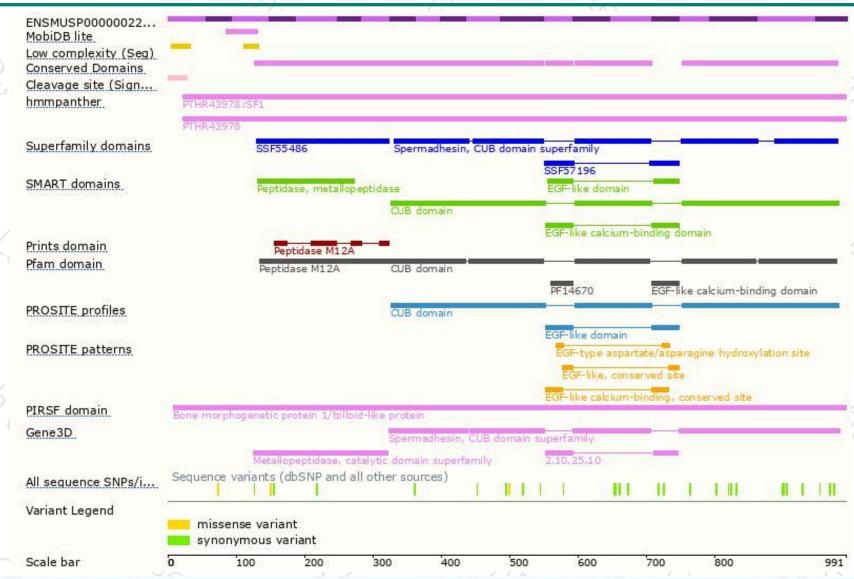
### Genomic location distribution





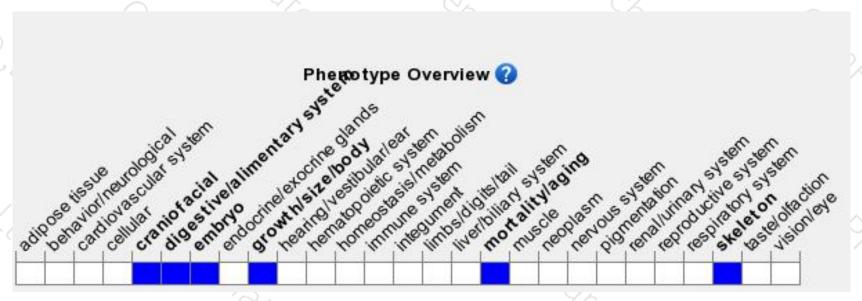
### Protein domain





## 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, Homozygous targeted mutant embryos have reduced ossification of the skull, persistent herniation of the gut, abnormal collagen fibrils in the amnion, and die at birth.



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





