

# Bbs9 Cas9-CKO Strategy

Designer: JiaYu

Reviewer: Xiaojing Li

**Design Date:** 2020-2-28

## **Project Overview**



**Project Name** 

Bbs9

**Project type** 

Cas9-CKO

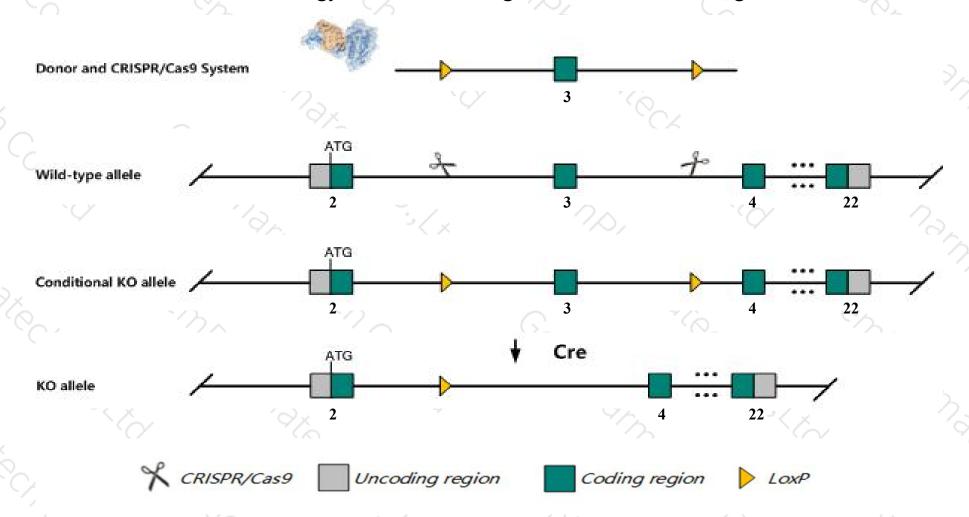
Strain background

C57BL/6JGpt

## Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Bbs9* gene. The schematic diagram is as follows:



### Technical routes



- The *Bbs9* gene has 12 transcripts. According to the structure of *Bbs9* gene, exon3 of *Bbs9-210*(ENSMUST00000147712.7) transcript is recommended as the knockout region. The region contains 151bp coding sequence.

  Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Bbs9* 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**



- > The *Bbs9* gene is located on the Chr9. 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.
- ➤ Transcript 206 CDS 5' incomplete the influences is unknown.
- > 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)



#### Bbs9 Bardet-Biedl syndrome 9 (human) [Mus musculus (house mouse)]

Gene ID: 319845, updated on 31-Jan-2019

#### Summary

☆ ?

Official Symbol Bbs9 provided by MGI

Official Full Name Bardet-Biedl syndrome 9 (human) provided by MGI

Primary source MGI:MGI:2442833

See related Ensembl: ENSMUSG00000035919

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 E130103I17Rik

Expression Ubiquitous expression in testis adult (RPKM 9.5), CNS E18 (RPKM 5.2) and 26 other tissuesSee more

Orthologs <u>human</u> all

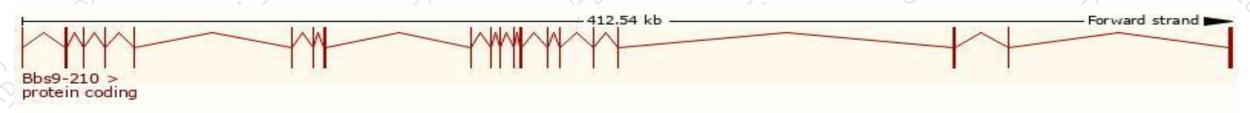
## Transcript information (Ensembl)



The gene has 12 transcripts, all transcripts are shown below:

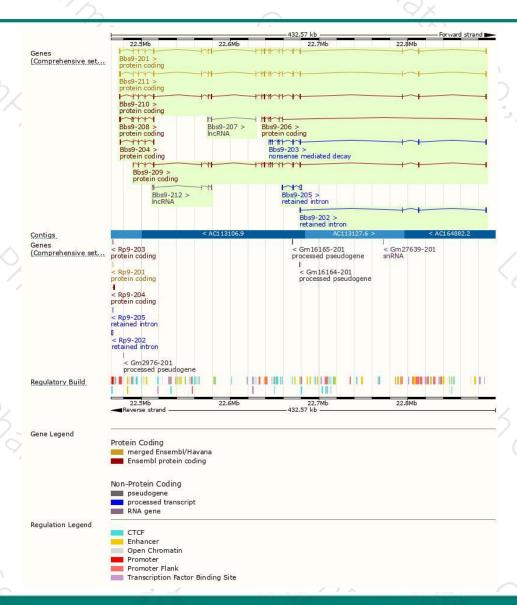
Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
ENSMUST00000147712.7	3495	880aa	Protein coding	CCDS22928	Q811G0	TSL:1 GENCODE basic APPRIS P2
ENSMUST00000150395.7	3425	880aa	Protein coding	CCDS22928	Q811G0	TSL:1 GENCODE basic APPRIS P2
ENSMUST00000039798.15	3382	880aa	Protein coding	CCDS22928	Q811G0	TSL:1 GENCODE basic APPRIS P2
ENSMUST00000147405.7	3045	885aa	Protein coding	29	Q811G0	TSL:5 GENCODE basic APPRIS ALT1
ENSMUST00000128812.7	833	<u>147aa</u>	Protein coding	- Bá	D3Z389	CDS 3' incomplete TSL:5
ENSMUST00000142313.7	833	<u>132aa</u>	Protein coding	+8	D3Z6T1	CDS 3' incomplete TSL:3
ENSMUST00000136084.7	638	<u>163aa</u>	Protein coding	<b>1</b> 8	F6XSQ3	CDS 5' incomplete TSL:5
ENSMUST00000127296.7	1640	<u>61aa</u>	Nonsense mediated decay	29	D6RIM6	CDS 5' incomplete TSL:5
ENSMUST00000130479.1	2641	No protein	Retained intron	- 5ú	-	TSL:1
ENSMUST00000124076.1	1012	No protein	Retained intron	<del>.</del> 8	*	TSL:3
ENSMUST00000152719.7	716	No protein	IncRNA	49	-	TSL:5
ENSMUST00000137547.1	542	No protein	IncRNA	29	12	TSL:3
	ENSMUST00000147712.7 ENSMUST00000150395.7 ENSMUST00000039798.15 ENSMUST00000147405.7 ENSMUST00000128812.7 ENSMUST00000142313.7 ENSMUST00000136084.7 ENSMUST00000127296.7 ENSMUST00000130479.1 ENSMUST00000124076.1 ENSMUST00000152719.7	ENSMUST00000147712.7 3495 ENSMUST00000150395.7 3425 ENSMUST00000039798.15 3382 ENSMUST00000147405.7 3045 ENSMUST00000128812.7 833 ENSMUST00000142313.7 833 ENSMUST00000136084.7 638 ENSMUST00000127296.7 1640 ENSMUST00000130479.1 2641 ENSMUST00000124076.1 1012 ENSMUST00000152719.7 716	ENSMUST00000147712.7         3495         880aa           ENSMUST00000150395.7         3425         880aa           ENSMUST00000039798.15         3382         880aa           ENSMUST00000147405.7         3045         885aa           ENSMUST00000128812.7         833         147aa           ENSMUST00000142313.7         833         132aa           ENSMUST00000136084.7         638         163aa           ENSMUST00000127296.7         1640         61aa           ENSMUST00000130479.1         2641         No protein           ENSMUST00000124076.1         1012         No protein           ENSMUST00000152719.7         716         No protein	ENSMUST00000147712.7         3495         880aa         Protein coding           ENSMUST00000150395.7         3425         880aa         Protein coding           ENSMUST00000039798.15         3382         880aa         Protein coding           ENSMUST00000147405.7         3045         885aa         Protein coding           ENSMUST00000128812.7         833         147aa         Protein coding           ENSMUST00000142313.7         833         132aa         Protein coding           ENSMUST00000136084.7         638         163aa         Protein coding           ENSMUST00000127296.7         1640         61aa         Nonsense mediated decay           ENSMUST00000130479.1         2641         No protein         Retained intron           ENSMUST00000152719.7         716         No protein         IncRNA	ENSMUST00000147712.7         3495         880aa         Protein coding         CCDS22928           ENSMUST00000150395.7         3425         880aa         Protein coding         CCDS22928           ENSMUST00000039798.15         3382         880aa         Protein coding         CCDS22928           ENSMUST00000147405.7         3045         885aa         Protein coding         -           ENSMUST00000128812.7         833         147aa         Protein coding         -           ENSMUST00000142313.7         833         132aa         Protein coding         -           ENSMUST00000136084.7         638         163aa         Protein coding         -           ENSMUST00000127296.7         1640         61aa         Nonsense mediated decay         -           ENSMUST00000130479.1         2641         No protein         Retained intron         -           ENSMUST00000124076.1         1012         No protein         Retained intron         -           ENSMUST00000152719.7         716         No protein         IncRNA         -	ENSMUST00000147712.7         3495         880aa         Protein coding         CCDS22928         Q811G0           ENSMUST00000150395.7         3425         880aa         Protein coding         CCDS22928         Q811G0           ENSMUST00000039798.15         3382         880aa         Protein coding         CCDS22928         Q811G0           ENSMUST00000147405.7         3045         885aa         Protein coding         -         Q811G0           ENSMUST00000128812.7         833         147aa         Protein coding         -         D3Z389           ENSMUST00000142313.7         833         132aa         Protein coding         -         D3Z6T1           ENSMUST00000136084.7         638         163aa         Protein coding         -         F6XSQ3           ENSMUST00000127296.7         1640         61aa         Nonsense mediated decay         -         D6RIM6           ENSMUST00000124076.1         1012         No protein         Retained intron         -         -           ENSMUST00000152719.7         716         No protein         IncRNA         -         -

The strategy is based on the design of *Bbs9-210* transcript, The transcription is shown below



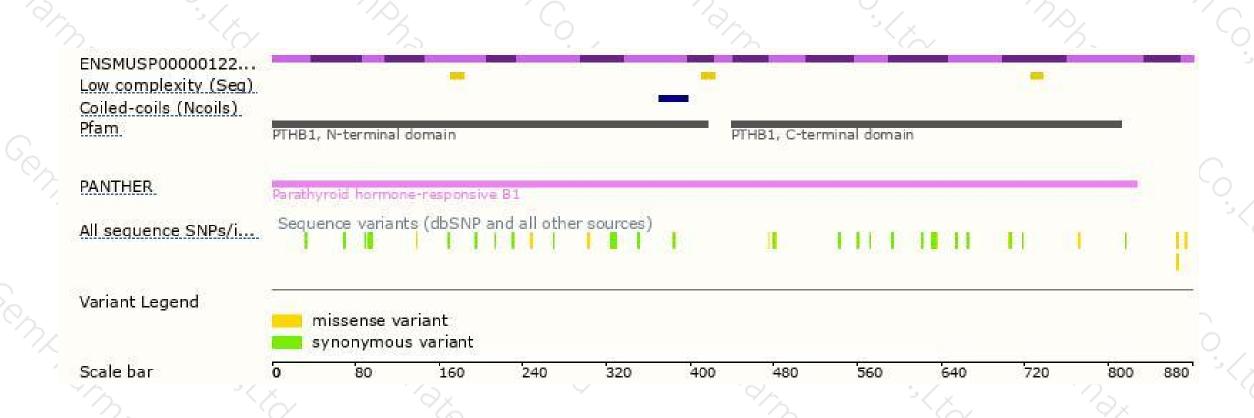
### Genomic location distribution





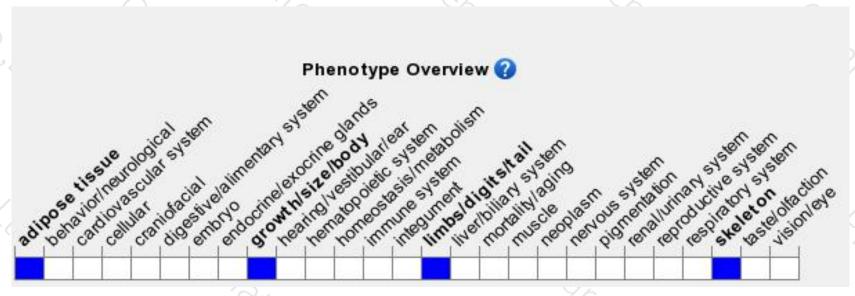
## Protein domain





## Mouse phenotype description(MGI)





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



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





