

# Hsd17b11 Cas9-CKO Strategy

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## **Project Overview**



**Project Name** 

Hsd17b11

**Project type** 

Cas9-CKO

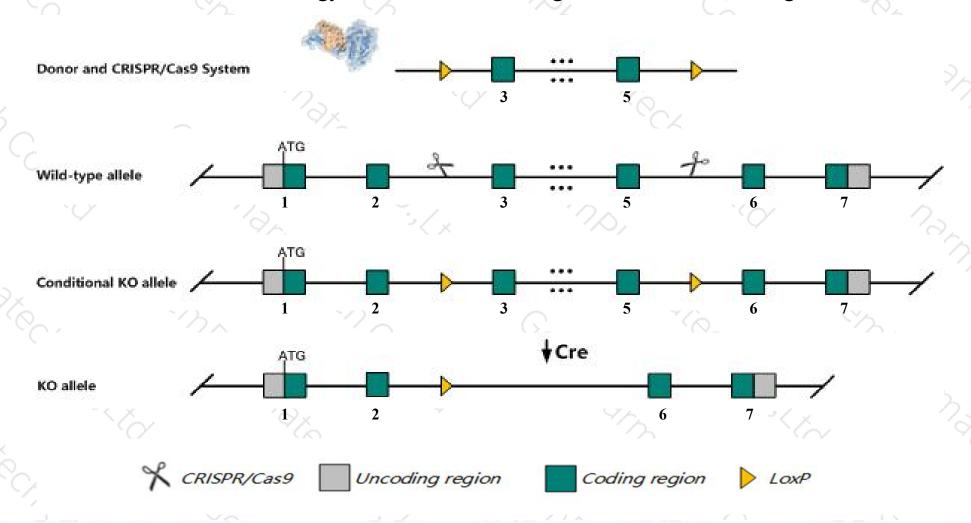
Strain background

C57BL/6JGpt

## Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Hsd17b11* gene. The schematic diagram is as follows:



### Technical routes



- The *Hsd17b11* gene has 2 transcripts. According to the structure of *Hsd17b11* gene, exon3-exon5 of *Hsd17b11-201* (ENSMUST00000031251.15) transcript is recommended as the knockout region. The region contains 377bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Hsd17b11* 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, Homozygous mutant mice exhibit an increased mean serum IgG2a response to ovalbumin challenge and an increased mean percentage of immature B cells in bone marrow.
- The *Hsd17b11* gene is located on the Chr5. 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)



#### Hsd17b11 hydroxysteroid (17-beta) dehydrogenase 11 [ Mus musculus (house mouse) ]

Gene ID: 114664, updated on 12-Aug-2019

#### Summary

☆ ?

Official Symbol Hsd17b11 provided by MGI

Official Full Name hydroxysteroid (17-beta) dehydrogenase 11 provided by MGI

Primary source MGI:MGI:2149821

See related Ensembl: ENSMUSG00000029311

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

Also known as SDR2; Dhrs8; Pan1b; retSDR2

Expression Ubiquitous expression in large intestine adult (RPKM 49.0), duodenum adult (RPKM 48.7) and 27 other tissues See more

Orthologs human all

#### Genomic context

☆ ?

Location: 5; 5 E5

See Hsd17b11 in Genome Data Viewer

Exon count: 7

Annotation release	Status	Assembly	Chr	Location	
108	current	GRCm38.p6 (GCF_000001635.26)	5	NC_000071.6 (103989765104021796, complement)	
Build 37.2	previous assembly	MGSCv37 (GCF_000001635.18)	5	NC_000071.5 (104418784104450815, complement)	

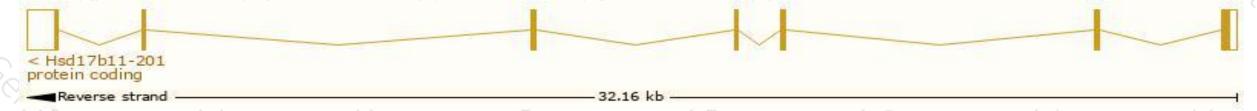
## Transcript information (Ensembl)



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

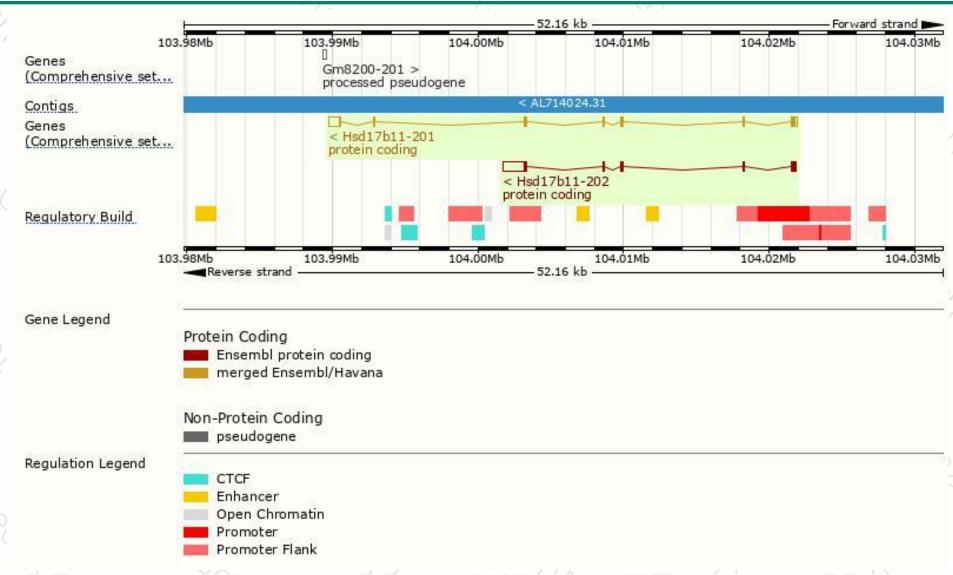
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Hsd17b11-201	ENSMUST00000031251.15	1825	298aa	Protein coding	CCDS19481	Q9EQ06	TSL:1 GENCODE basic APPRIS P2
Hsd17b11-202	ENSMUST00000119025.1	2234	232aa	Protein coding	690	Q9EQ06	TSL:1 GENCODE basic APPRIS ALT2

The strategy is based on the design of *Hsd17b11-201* transcript, The transcription is shown below



### Genomic location distribution





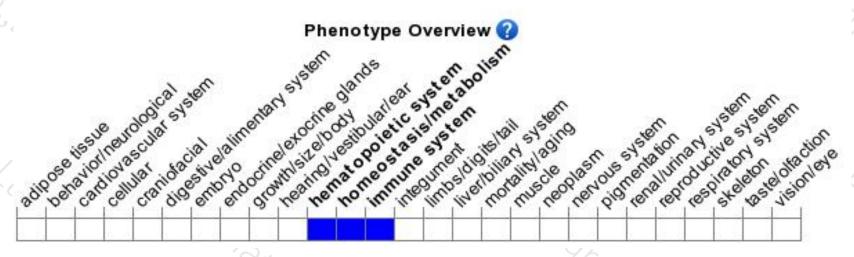
### Protein domain





## 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 mutant mice exhibit an increased mean serum IgG2a response to ovalbuming challenge and an increased mean percentage of immature B cells in bone marrow.



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





