

# Lonp1 Cas9-CKO Strategy

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Reviewer: JiaYu

# **Project Overview**



**Project Name** 

Lonp1

**Project type** 

Cas9-CKO

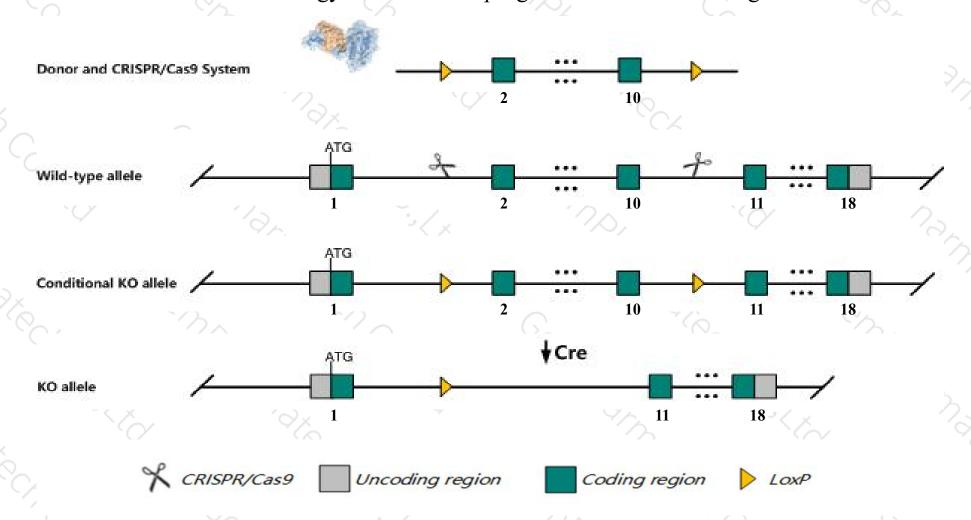
Strain background

C57BL/6JGpt

## Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Lonp1* gene. The schematic diagram is as follows:



### Technical routes



- ➤ The *Lonp1* gene has 2 transcripts. According to the structure of *Lonp1* gene, exon2-exon10 of *Lonp1-201*(ENSMUST00000047226.9) transcript is recommended as the knockout region. The region contains 1259bp coding sequence.

  Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Lonp1* 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, Mice homozygous for a knock-out allele exhibit embryonic lethality with embryonic growth retardation, small size and decreased mitochondrial DNA content. Mice heterozygous for this allele exhibit reduced chemically-induced tumors.
- > The *Lonp1* gene is located on the Chr17. 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)



#### Lonp1 Ion peptidase 1, mitochondrial [Mus musculus (house mouse)]

Gene ID: 74142, updated on 7-Apr-2019

#### Summary

☆ ?

Official Symbol Lonp1 provided by MGI

Official Full Name Ion peptidase 1, mitochondrial provided by MGI

Primary source MGI:MGI:1921392

See related Ensembl: ENSMUSG00000041168

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 1200017E13Rik, LON, Prss15

Expression Broad expression in adrenal adult (RPKM 284.2), ovary adult (RPKM 104.2) and 27 other tissuesSee more

Orthologs human all

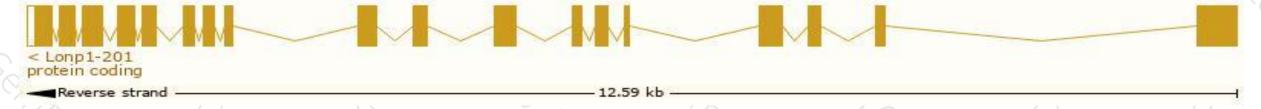
# Transcript information (Ensembl)



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

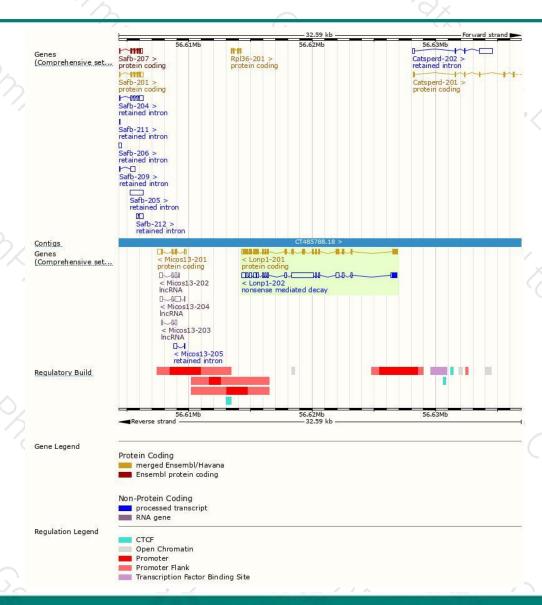
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Lonp1-201	ENSMUST00000047226.9	2951	949aa	Protein coding	CCDS28910	Q8CGK3	TSL:1 GENCODE basic APPRIS P1
Lonp1-202	ENSMUST00000233732.1	4362	<u>141aa</u>	Nonsense mediated decay	<del>.</del> 8	A0A3B2W834	

The strategy is based on the design of Lonp1-201 transcript, The transcription is shown below



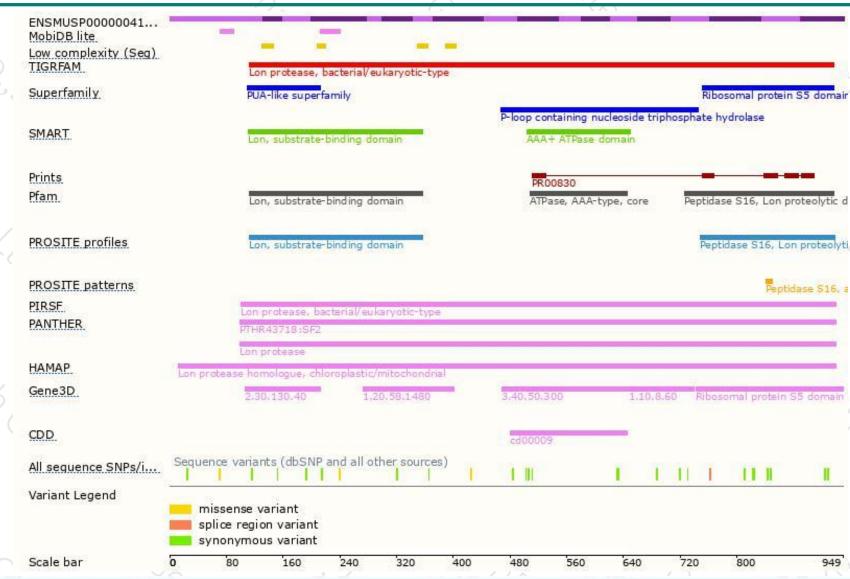
### Genomic location distribution





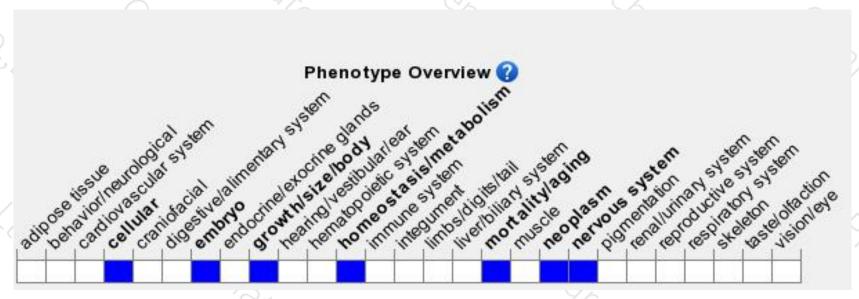
### Protein domain





# 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, Mice homozygous for a knock-out allele exhibit embryonic lethality with embryonic growth retardation, small size and decreased mitochondrial DNA content. Mice heterozygous for this allele exhibit reduced chemically-induced tumors.



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





