# Psme3 Cas9-CKO Strategy

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**Design Date:** 

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## **Project Overview**



**Project Name** 

Psme3

**Project type** 

Cas9-CKO

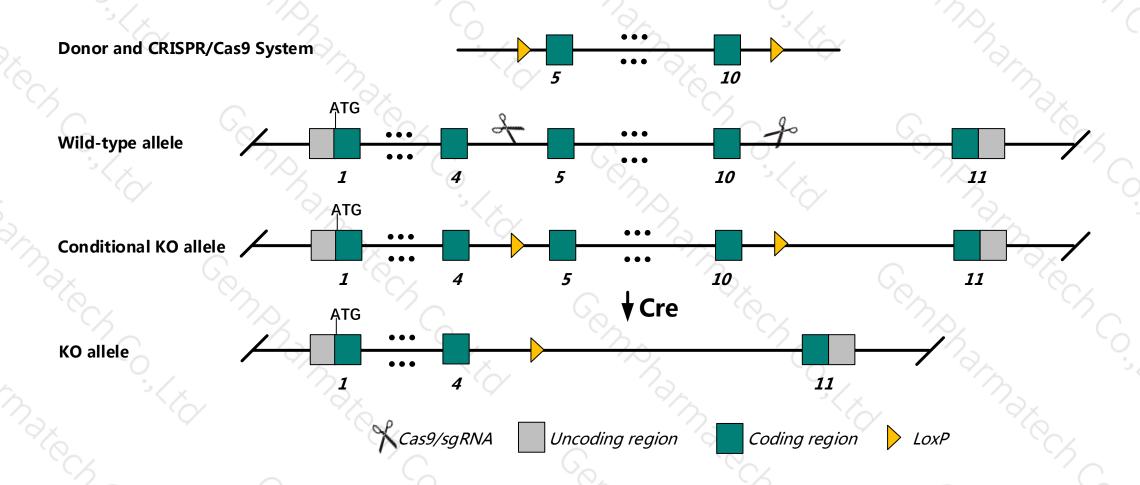
Strain background

C57BL/6JGpt

## **Conditional Knockout strategy**



This model will use CRISPR/Cas9 technology to edit the *Psme3* gene. The schematic diagram is as follows:



### **Technical routes**



- ➤ The *Psme3* gene has 5 transcript. According to the structure of *Psme3* gene, exon5~10 of *Psme3*-201 (ENSMUST00000019470.13) transcript is recommended as the knockout region. The region contains 441bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Psme3* gene. The brief process is as follows: sgRNA was transcribed in vitro, donor vector was constructed.Cas9, sgRNA 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 was knocked out after mating with mice expressing Cre recombinase, resulting in the loss of function of the target gene in specific tissues or cell types.

### **Notice**



- According to the existing MGI data, Homozygous null mutants are smaller than normal with a defect in cell proliferation and increased susceptibility to fungal infection.
- $\triangleright$  This strategy may affect the 5 terminal regulatory function of the Aoc2 gene.
- ➤ The *Psme3* gene is located on the Chr11. 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 loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.

## Gene information (NCBI)



Psme3 proteaseome (prosome, macropain) activator subunit 3 (PA28 gamma, Ki) [ Mus musculus (house mouse) ]

Gene ID: 19192, updated on 12-Aug-2019

#### Summary



Official Symbol Psme3 provided by MGI

Official Full Name proteaseome (prosome, macropain) activator subunit 3 (PA28 gamma, Ki) provided by MGI

Primary source MGI:MGI:1096366

See related Ensembl: ENSMUSG00000078652

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 Ki; pa28g; AA410043; AU020960; REGgamma; PA28gamma

Expression Ubiquitous expression in liver E14.5 (RPKM 81.8), liver E14 (RPKM 74.6) and 28 other tissues See more

Orthologs human all

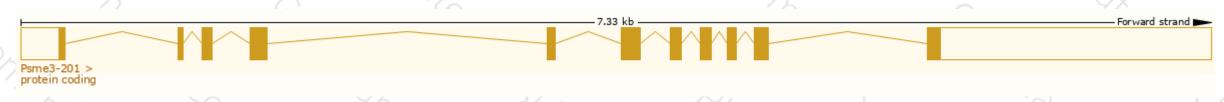
## Transcript information (Ensembl)



The gene has 5 transcripts, and all transcripts are shown below:

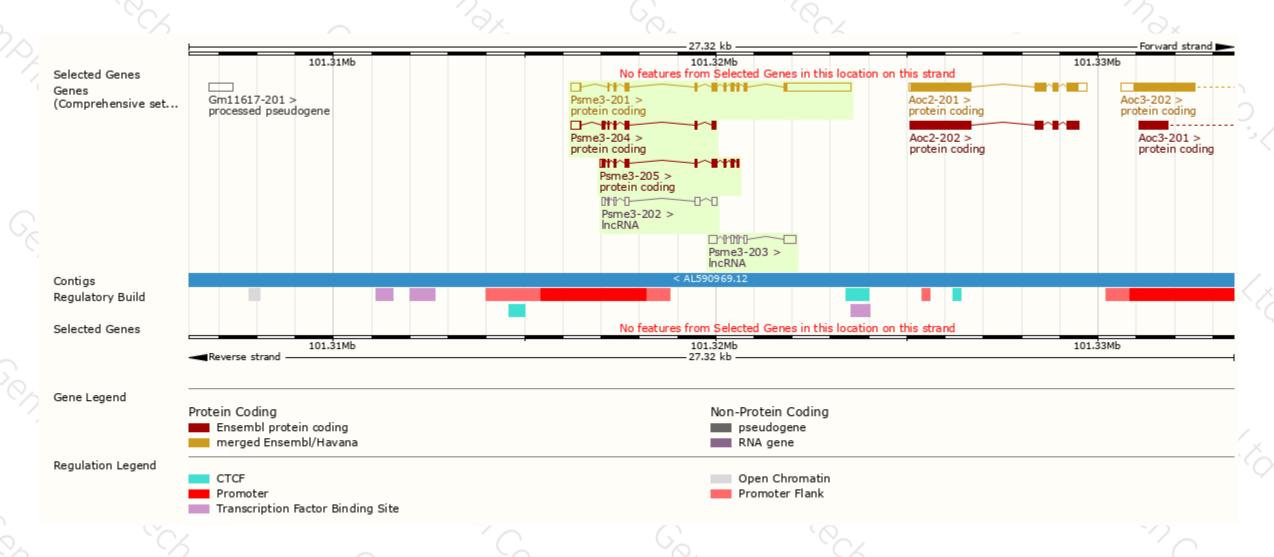
Name 🍦	Transcript ID 🗼	bp 🍦	Protein	Biotype	CCDS	UniProt ▼	Flags
Psme3-201	ENSMUST00000019470.13	2664	<u>254aa</u>	Protein coding	CCDS25463₺	<u>P61290</u> & Q4FK54&	TSL:1 GENCODE basic APPRIS P1
Psme3-204	ENSMUST00000142640.7	698	<u>154aa</u>	Protein coding	-	<u>A2A4J3</u> ₺	CDS 3' incomplete TSL:3
Psme3-205	ENSMUST00000151385.1	653	<u>203aa</u>	Protein coding	-	<u>A2A4J1</u> ₺	CDS 3' incomplete TSL:5
Psme3-203	ENSMUST00000131170.1	778	No protein	IncRNA	-	-	TSL:2
Psme3-202	ENSMUST00000127998.1	524	No protein	IncRNA	-	-	TSL:5

The strategy is based on the design of *Psme3*-201 transcript, The transcription is shown below:



### Genomic location distribution





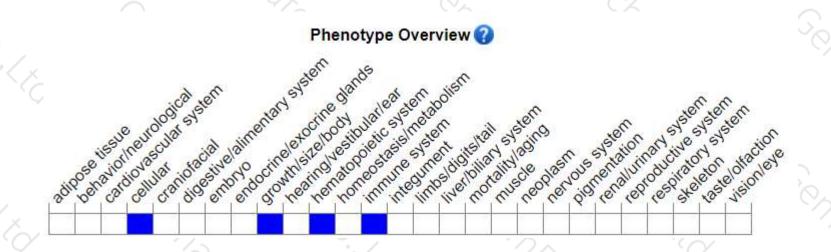
### Protein domain



	ENSMUSP00000019 Superfamily					· · · ·		
	Pfam	Proteasome activator superfamily Proteasome activator PA28, N-terminal domain			Proteasome activator PA28, C-terminal domain			
	PANTHER	Proteasome activator PA28						
O	Gene3D	PTHR10660:SF4 Proteasome activator PA	A28, N-terminal domain supe	rfamily	Proteasome activator PA28, C-t	erminal domain superfamily		
	All sequence SNPs/i	Sequence variants (dbSI	NP and all other sources)		1			,
	Variant Legend	synonymous variar	nt					
	Scale bar	0	40	80	120	160	200	254
	Dydyndxer	-X	Donal Dale	2% Co.3(x)	CONDAN.	373× (SCY Co. < )	X JOHN S	77 Co.34

### 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 null mutants are smaller than normal with a defect in cell proliferation and increased susceptibility to fungal infection.

If you have any questions, you are welcome to inquire. Tel: 025-5864 1534





