

# Erp29 Cas9-CKO Strategy

Designer:Xiaojing Li

Reviewer:JiaYu

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## **Project Overview**



**Project Name** 

Erp29

**Project type** 

Cas9-CKO

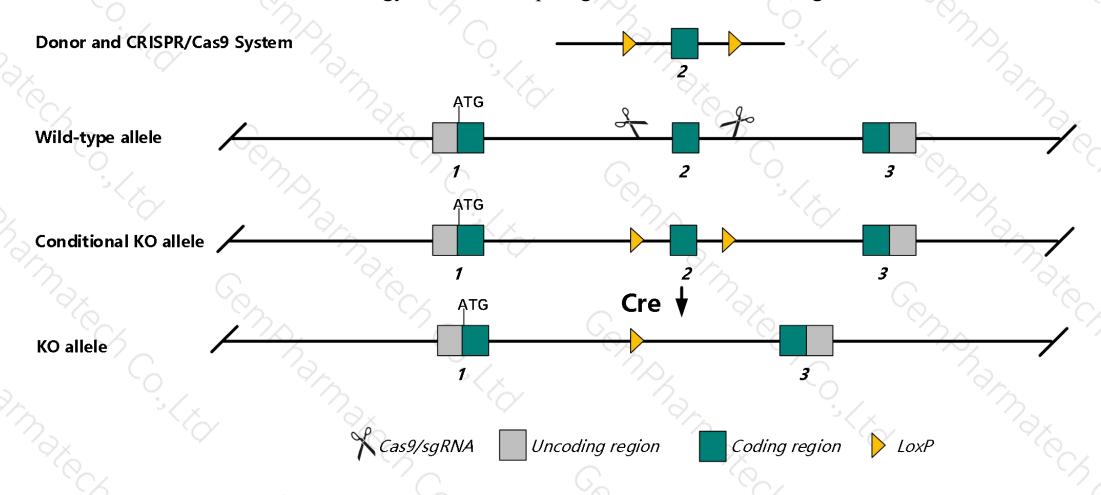
Strain background

C57BL/6JGpt

## Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Erp29 gene. The schematic diagram is as follows:



### Technical routes



- The *Erp29* gene has 4 transcripts. According to the structure of *Erp29* gene, exon2 of *Erp29-203*(ENSMUST00000130451.1) transcript is recommended as the knockout region. The region contains 139bp coding sequence.

  Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Erp29* 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 null allele exhibit decreased thyrocyte and dermal fibroblast apoptosis sensitivities to induced ER stress and slightly reduced litter sizes.
- ➤ The knockout region is located at the 5-terminal of Tmem116 about 4kb, which may affect its 5-terminal regulation after knockout.
- ➤ This strategy has no impact on Erp29-201.
- > The Erp29 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)



#### Erp29 endoplasmic reticulum protein 29 [ Mus musculus (house mouse) ]

Gene ID: 67397, updated on 20-Aug-2019

#### Summary

Official Symbol Erp29 provided by MGI

Official Full Name endoplasmic reticulum protein 29 provided by MGI

Primary source MGI:MGI:1914647

See related Ensembl: ENSMUSG00000029616

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 Erp28; Erp31; PDI-Db; AW209030; 1200015M03Rik; 2810446M09Rik

Expression Ubiquitous expression in placenta adult (RPKM 91.8), ovary adult (RPKM 52.5) and 28 other tissues See more

Orthologs human all

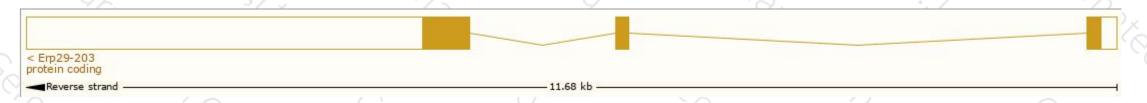
## Transcript information (Ensembl)



The gene has 4 transcripts, all transcripts are shown below:

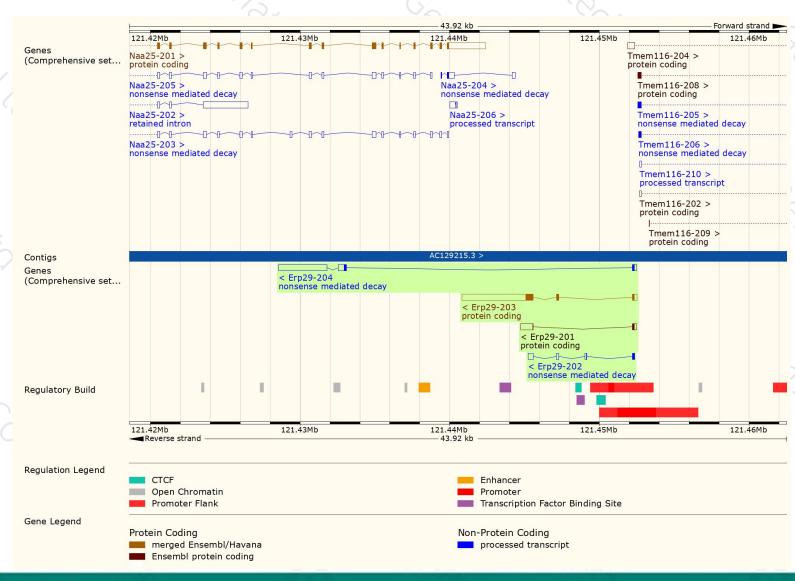
Name A	Transcript ID 🍦	bp 🍦	Protein 4	Biotype	CCDS 🍦	UniProt 4		Flags	4
Erp29-201	ENSMUST00000052590.7	1123	<u>55aa</u>	Protein coding	3.50	F8WJI4 €	TSL:1	GENCODE basic	
Erp29-202	ENSMUST00000111802.3	748	<u>57aa</u>	Nonsense mediated decay	S=3	F8WIM7 €	TSL:2		
Erp29-203	ENSMUST00000130451.1	5204	<u>262aa</u>	Protein coding	CCDS19634₽	<u>P57759</u> ₺	TSL:1 GENO	ODE basic APPRIS	S P1
Erp29-204	ENSMUST00000153758.1	4034	<u>95aa</u>	Nonsense mediated decay	5.E3	<u>D6RG87</u> ₽		TSL:1	

The strategy is based on the design of Erp29-203 transcript, The transcription is shown below



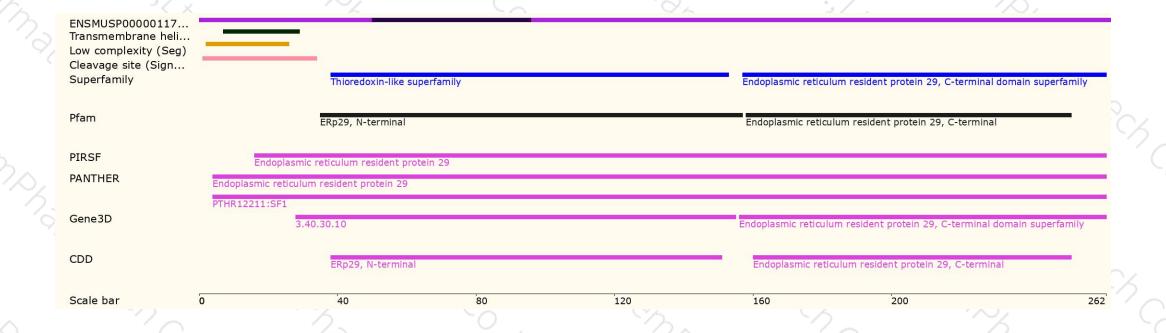
### Genomic location distribution





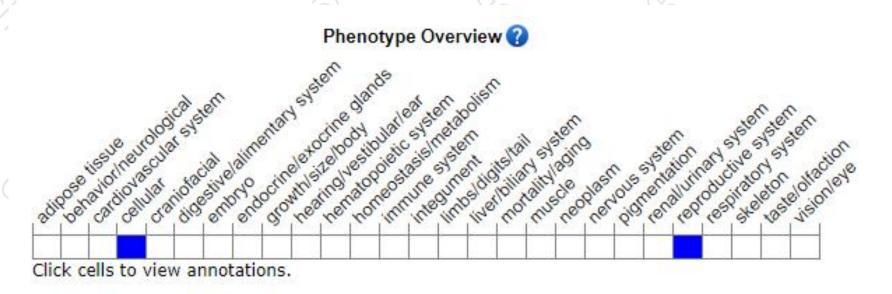
### Protein domain





## Mouse phenotype description(MGI)





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

Mice homozygous for a null allele exhibit decreased thyrocyte and dermal fibroblast apoptosis sensitivities to induced ER stress and slightly reduced litter sizes.



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





