

# Lrig1 Cas9-CKO Strategy

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**Design Date:** 2019-12-23

## **Project Overview**



**Project Name** 

Lrig1

**Project type** 

Cas9-CKO

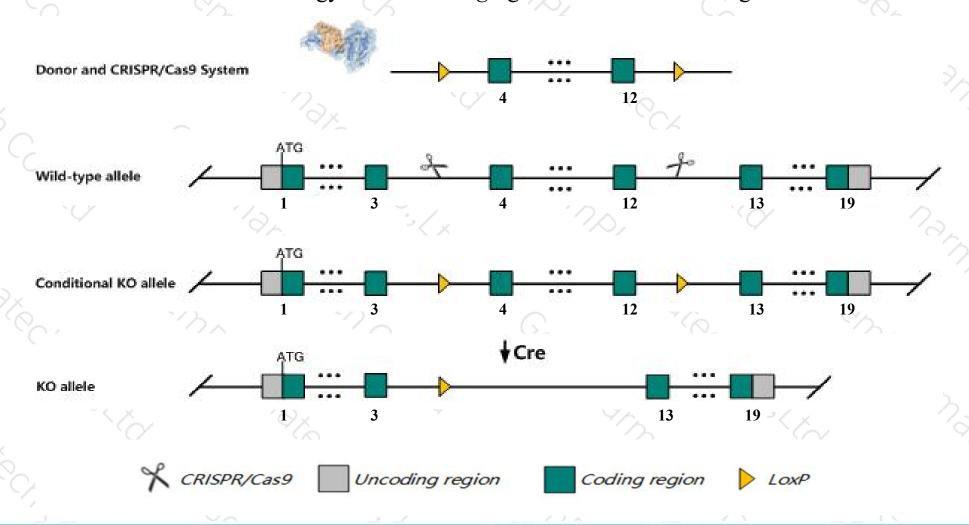
Strain background

C57BL/6JGpt

## Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Lrig1* gene. The schematic diagram is as follows:



### Technical routes



- ➤ The *Lrig1* gene has 7 transcripts. According to the structure of *Lrig1* gene, exon4-exon12 of *Lrig1-202*(ENSMUST00000101126.2) transcript is recommended as the knockout region. The region contains 1103bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Lrig1* 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, Homozygous null mice developed psoriasiform epidermal hyperplasia. Homozygotes exhibit hair follicle, epidermis, vertebral, eye and hearing abnormalities, decreased body size and fat amount, and increased susceptibility to bacterial infection.
- The *Lrig1* gene is located on the Chr6. 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)



#### Lrig1 leucine-rich repeats and immunoglobulin-like domains 1 [ Mus musculus (house mouse) ]

Gene ID: 16206, updated on 16-Dec-2019

#### Summary



Official Symbol Lrig1 provided by MGI

Official Full Name leucine-rich repeats and immunoglobulin-like domains 1 provided by MGI

Primary source MGI:MGI:107935

See related Ensembl: ENSMUSG00000030029

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 Img; LIG-1; D6Bwg0781e

Expression Ubiquitous expression in colon adult (RPKM 30.2), cerebellum adult (RPKM 20.8) and 26 other tissues See more

Orthologs <u>human</u> all

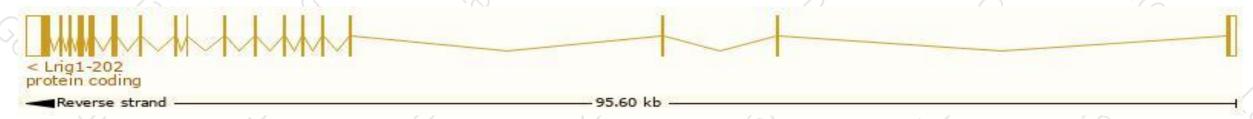
## Transcript information (Ensembl)



The gene has 7 transcripts, all transcripts are shown below:

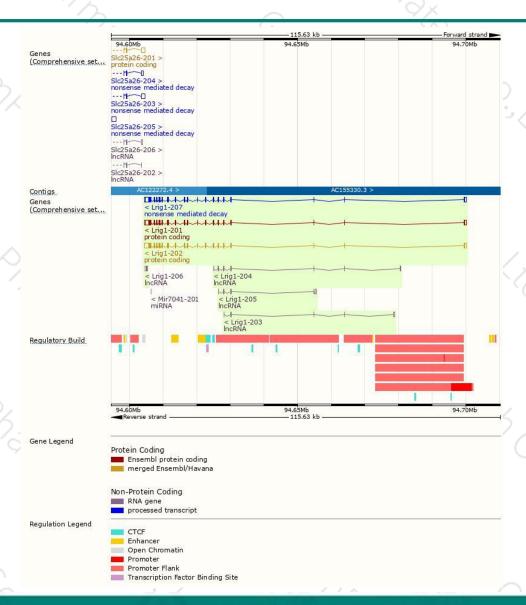
Name 🍦	Transcript ID 🍦	bp 🌲	Protein 4	Translation ID 👙	Biotype	CCDS 🍦	UniProt 🌲	Flags
Lrig1-202	ENSMUST00000101126.2	4951	<u>1091aa</u>	ENSMUSP00000098686.2	Protein coding	CCDS20380 ₽	P70193₺	TSL:1 GENCODE basic APPRIS P1
Lrig1-201	ENSMUST00000032105.10	4831	<u>1091aa</u>	ENSMUSP00000032105.4	Protein coding	CCDS20380 ₽	P70193₽	TSL:1 GENCODE basic APPRIS P1
Lrig1-207	ENSMUST00000204645.2	4849	1091aa	ENSMUSP00000144963.1	Nonsense mediated decay	æ 1	P70193₽	TSL:1
Lrig1-205	ENSMUST00000150811.7	700	No protein	(3-)	<b>I</b> IncRNA	9 <del>-</del> 0.	-	TSL:3
Lrig1-204	ENSMUST00000141014.7	666	No protein	(3-)	I IncRNA	9 <del>-</del> 0.	-	TSL:3
Lrig1-203	ENSMUST00000124165.1	587	No protein	(3-)	<b>I</b> IncRNA	9 <del>-</del> 0.	-	TSL:3
Lrig1-206	ENSMUST00000203876.1	438	No protein	(ce)	IncRNA	890.	-	TSL:3

The strategy is based on the design of Lrig1-202 transcript, The transcription is shown below



## Genomic location distribution





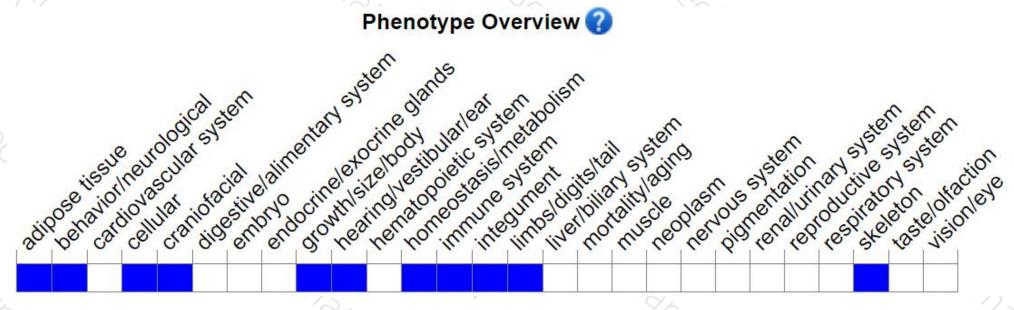
### Protein domain





## Mouse phenotype description(MGI)





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

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





