

# Actn4 Cas9-KO Strategy

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### **Project Overview**



**Project Name** 

Actn4

**Project type** 

Cas9-KO

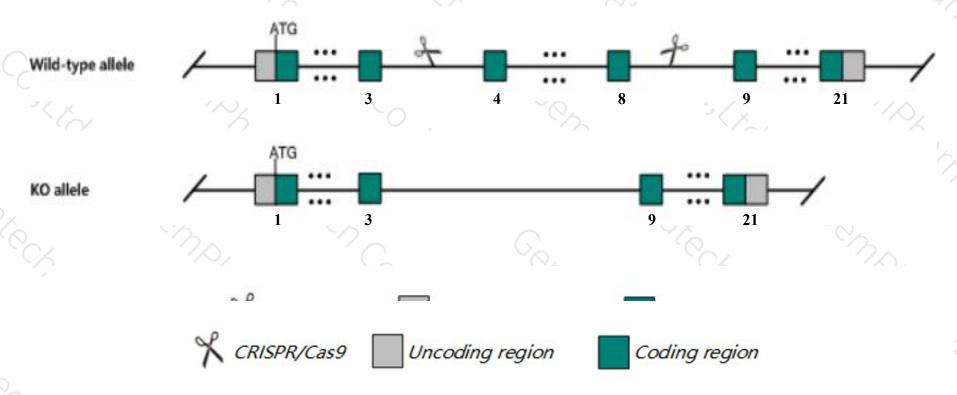
Strain background

C57BL/6JGpt

### **Knockout strategy**



This model will use CRISPR/Cas9 technology to edit the *Actn4* gene. The schematic diagram is as follows:



The KO region includes exon8(86bp) of Actn4-202 transcript.

### **Technical routes**



- The *Actn4* gene has 11 transcripts. According to the structure of *Actn4* gene, exon4-exon8 of *Actn4-201* (ENSMUST00000068045.13) transcript is recommended as the knockout region. The region contains 422bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Actn4* gene. The brief process is as follows: CRISPR/Cas9 system v

### **Notice**



- ➤ According to the existing MGI data, mice homozygous for a disruption in this gene die either around birth or within a few months of birth. those who do survive after birth show poor growth and kidney abnormalities including glomerulosclerosis. this is manifested functionally as proteinuria and abnormal blood urea nitrogen.
- > The *Actn4* gene is located on the Chr7. 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 gene knockout on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.

### Gene information (NCBI)



#### Actn4 actinin alpha 4 [Mus musculus (house mouse)]

Gene ID: 60595, updated on 13-Mar-2020

#### Summary

☆ ?

Official Symbol Actn4 provided by MGI

Official Full Name actinin alpha 4 provided by MGI

Primary source MGI:MGI:1890773

See related Ensembl:ENSMUSG00000054808

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 C77391

Expression Ubiquitous expression in small intestine adult (RPKM 213.0), large intestine adult (RPKM 207.4) and 28 other tissuesSee more

Orthologs human all

# Transcript information (Ensembl)



The gene has 11 transcripts, all transcripts are shown below:

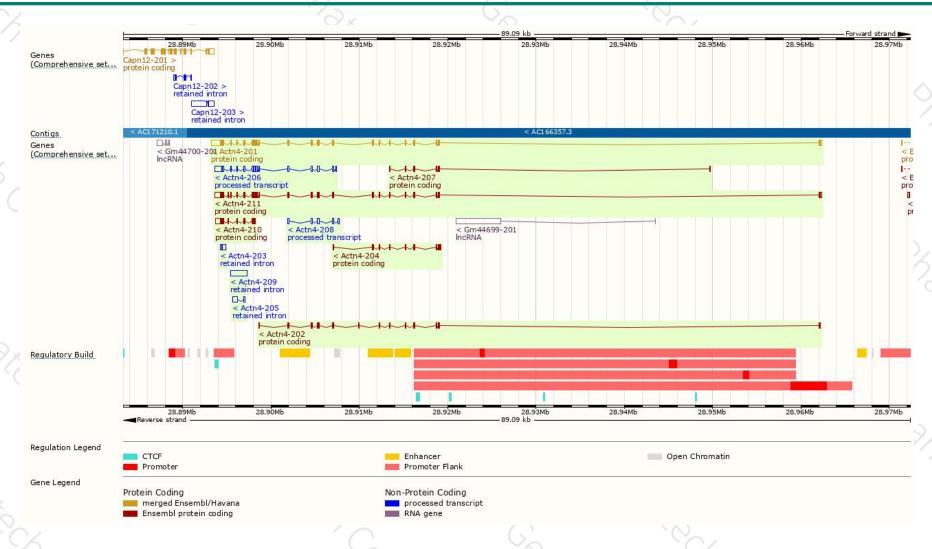
	* /						
Name 🍦	Transcript ID	bp 🍦	Protein 4	Biotype	CCDS 🍦	UniProt	Flags
Actn4-201	ENSMUST00000068045.13	3877	<u>912aa</u>	Protein coding	CCDS21061 ₽	P57780 ₺ Q3ULT2 ₺	TSL:1 GENCODE basic APPRIS P2
Actn4-211	ENSMUST00000217157.1	3530	<u>932aa</u>	Protein coding		A0A1L1SV25₽	TSL:5 GENCODE basic APPRIS ALT1
Actn4-202	ENSMUST00000127210.7	1594	<u>518aa</u>	Protein coding		E9Q2W9₽	CDS 3' incomplete TSL:5
Actn4-210	ENSMUST00000216863.1	1489	323aa	Protein coding	-	A0A1L1SVJ6굡	CDS 5' incomplete TSL:5
Actn4-204	ENSMUST00000140622.7	774	206aa	Protein coding	-	<u>D3Z0L8</u> ₽	CDS 3' incomplete TSL:5
Actn4-207	ENSMUST00000148196.2	617	<u>133aa</u>	Protein coding	-	<u>D3Z761</u> ₽	CDS 3' incomplete TSL:5
ctn4-206	ENSMUST00000144909.7	2581	No protein	Processed transcript	-	-	TSL:1
Actn4-208	ENSMUST00000150493.1	726	No protein	Processed transcript		-	TSL:3
Actn4-209	ENSMUST00000207765.1	1890	No protein	Retained intron		-	TSL:NA
Actn4-205	ENSMUST00000143584.1	750	No protein	Retained intron			TSL:1
Actn4-203	ENSMUST00000129338.1	508	No protein	Retained intron	-	-	TSL:1

The strategy is based on the design of Actn4-201 transcript, the transcription is shown below



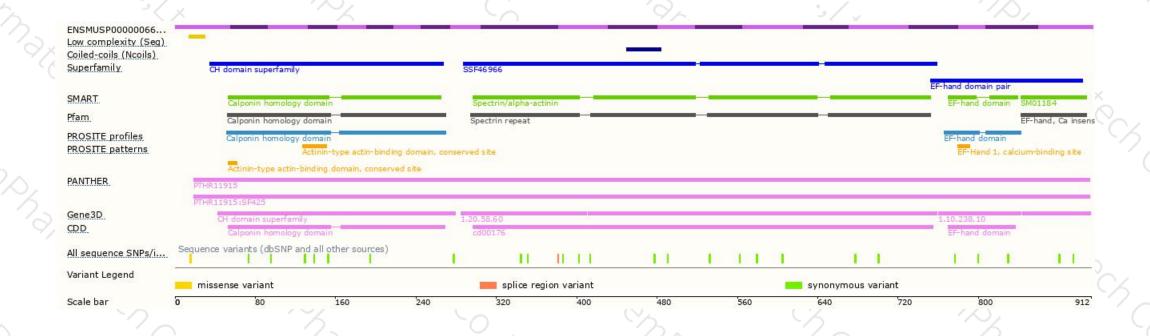
### Genomic location distribution





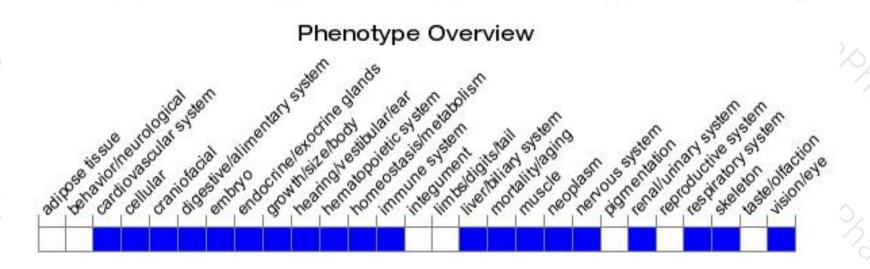
### Protein domain





## 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 disruption in this gene die either around birth or within a few months of birth. Those who do survive after birth show poor growth and kidney abnormalities including glomerulosclerosis. This is manifested functionally as proteinuria and abnormal blood urea nitrogen.



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





