# Cul7 Cas9-KO Strategy

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Reviewer: Huan Fan

**Design Date:** 2020-4-16

# **Project Overview**



**Project Name** 

Cul7

**Project type** 

Cas9-KO

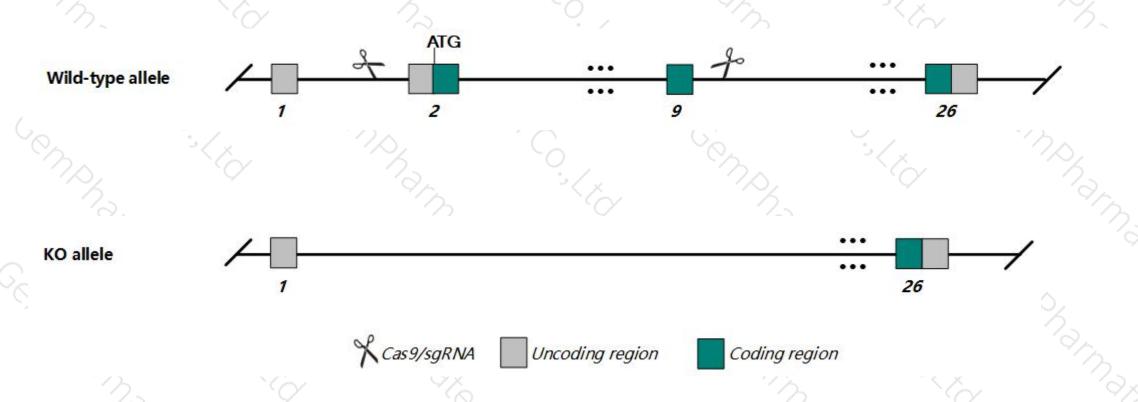
Strain background

C57BL/6JGpt

### **Knockout strategy**



This model will use CRISPR/Cas9 technology to edit the *Cul7* gene. The schematic diagram is as follows:



### **Technical routes**



- The *Cul7* gene has 7 transcripts. According to the structure of *Cul7* gene, exon2-9 of *Cul7*-201 (ENSMUST00000043464.13) transcript is recommended as the knockout region. The region contains start codon ATG. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Cul7* gene. The brief process is as follows: gRNA was transcribed in vitro.Cas9, gRNA 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.

### **Notice**



- According to the existing MGI data, during late gestation, homozygous null fetuses display reduced growth associated with abnormal placental development and hemorrhaging due to vascular defects. Mutant mice are born but die shortly after birth, succumbing to respiratory distress.
- The *Cul7* 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 the loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.

# Gene information (NCBI)



#### Cul7 cullin 7 [ Mus musculus (house mouse) ]

Gene ID: 66515, updated on 13-Mar-2020

#### Summary

☆ ?

Official Symbol Cul7 provided by MGI
Official Full Name cullin 7 provided by MGI
Primary source MGI:MGI:1913765

See related Ensembl: ENSMUSG00000038545

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 p185; p193; AA409809; 2510004L20Rik; C230011P08Rik

Expression Ubiquitous expression in limb E14.5 (RPKM 31.4), placenta adult (RPKM 28.7) and 27 other tissues See more

Orthologs human all

# Transcript information (Ensembl)



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

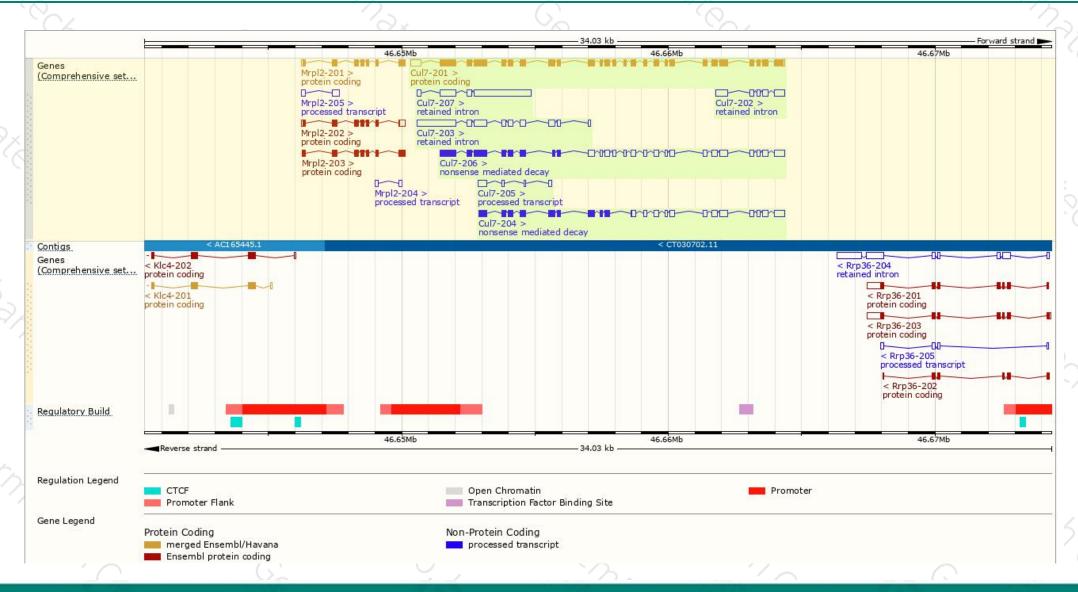
Name 🌲	Transcript ID .	bp 🌲	Protein	Biotype	CCDS 🍦	UniProt	Flags
Cul7-201	ENSMUST00000043464.13	5530	<u>1689aa</u>	Protein coding	CCDS28833₽	A9C491 & Q8VE73 &	TSL:1 GENCODE basic APPRIS P1
Cul7-206	ENSMUST00000145567.7	5016	649aa	Nonsense mediated decay	-	E9Q6H4₽	TSL:1
Cul7-204	ENSMUST00000133393.7	4047	<u>577aa</u>	Nonsense mediated decay	194	F6Q1R4函	CDS 5' incomplete TSL:5
Cul7-205	ENSMUST00000144966.1	628	No protein	Processed transcript	:-:	€:	TSL:5
Cul7-203	ENSMUST00000132790.1	3056	No protein	Retained intron	-	€:	TSL:1
Cul7-207	ENSMUST00000151002.7	3010	No protein	Retained intron	-	€:	TSL:1
Cul7-202	ENSMUST00000125949.1	1321	No protein	Retained intron	6.40		TSL:1

The strategy is based on the design of Cul7-201 transcript, the transcription is shown below



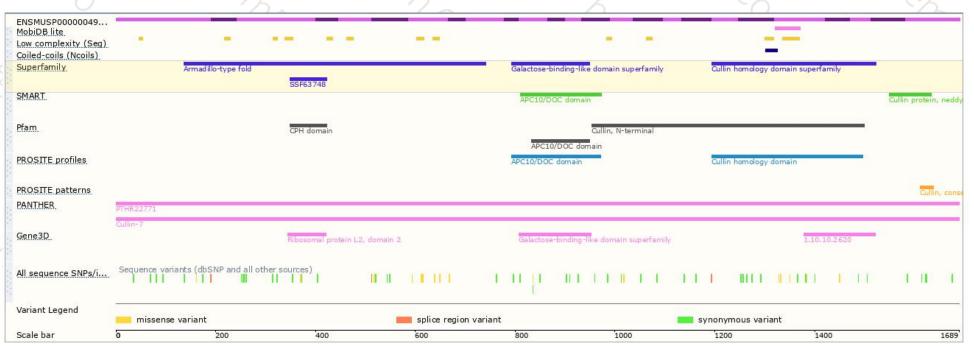
### Genomic location distribution





### Protein domain





Caratter at a

Ave. residue weight: 113.850 g/mol

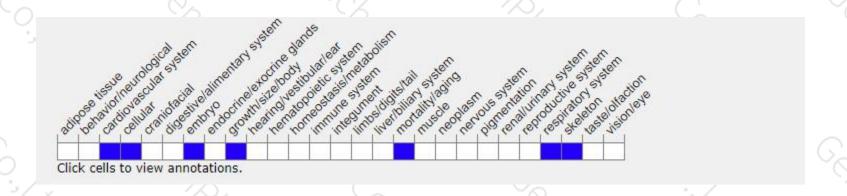
Charge: -8.5

Isoelectric point: 6.2429

Molecular weight: 192,292.17 g/mol Number of residues: 1,689 aa

## Mouse phenotype description(MGI)





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

During late gestation, homozygous null fetuses display reduced growth associated with abnormal placental development and hemorrhaging due to vascular defects. Mutant mice are born but die shortly after birth, succumbing to respiratory distress.

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





