

# Chn1 Cas9-KO Strategy

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**Reviewer:** 

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# **Project Overview**



**Project Name** 

Chn1

**Project type** 

Cas9-KO

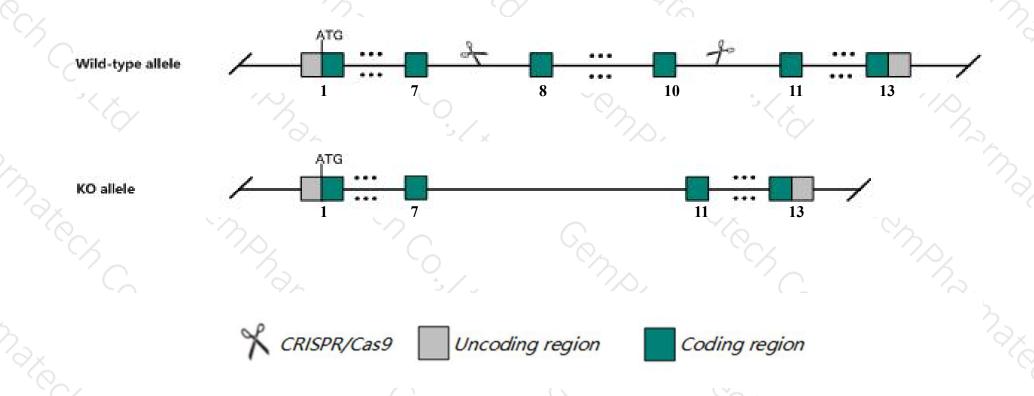
Strain background

C57BL/6JGpt

# **Knockout strategy**



This model will use CRISPR/Cas9 technology to edit the *Chn1* gene. The schematic diagram is as follows:



### **Technical routes**



- ➤ The *Chn1* gene has 15 transcripts. According to the structure of *Chn1* gene, exon8-exon10 of *Chn1-203*(ENSMUST00000112024.9) transcript is recommended as the knockout region. The region contains 337bp coding sequence.

  Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Chn1* gene. The brief process is as follows: CRISPR/Cas9 system

### **Notice**



- According to the existing MGI data, mice homologous for a null allele exhibit transient postnatal size reduction, abnormal gait and abnormal innervation of the spinal cord. Part of null homozygous show preweaning lethality.
- > Transcript *Chn1-205* may not be affected.
- > The *Chn1* gene is located on the Chr2. 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)



#### Chn1 chimerin 1 [ Mus musculus (house mouse) ]

Gene ID: 108699, updated on 19-Nov-2019

#### Summary

△ ?

Official Symbol Chn1 provided by MGI

Official Full Name chimerin 1 provided by MGI

Primary source MGI:MGI:1915674

See related Ensembl: ENSMUSG00000056486

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 ARHGAP2; Al413815; 0610007l19Rik; 0710001E19Rik; 1700112L09Rik; 2900046J01Rik

Expression Biased expression in cortex adult (RPKM 122.5), frontal lobe adult (RPKM 81.5) and 6 other tissues See more

Orthologs human all

# Transcript information (Ensembl)



#### The gene has 15 transcripts, all transcripts are shown below:

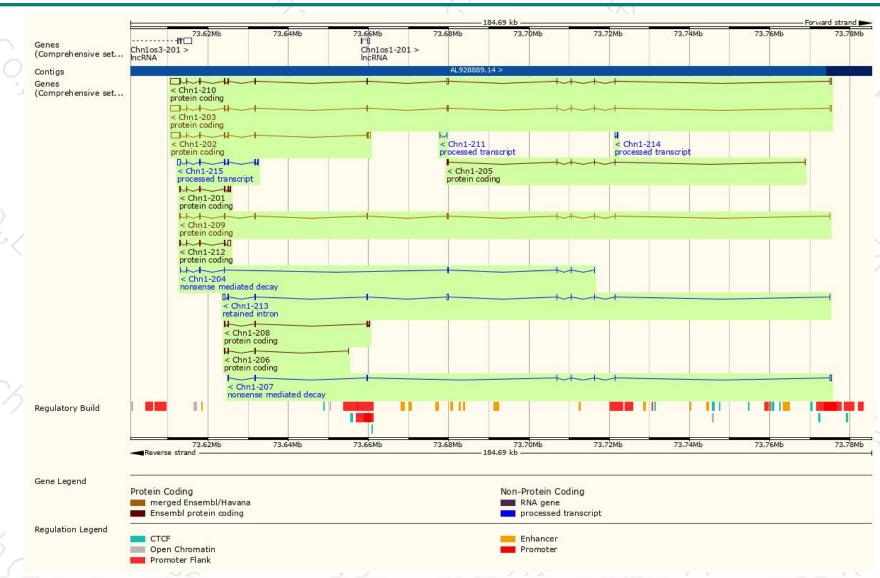
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Name	Transcript ID 👙	bp 🌲	Protein	Biotype	CCDS 🍦	UniProt	Flags
Chn1-203	ENSMUST00000112024.9	4050	459aa	Protein coding	CCDS50608 ₽	Q91V57 ₽	TSL:1 GENCODE basic APPRIS P1
Chn1-210	ENSMUST00000180045.7	4050	210aa	Protein coding	CCDS38145 ₽	A7VK14 & Q91V57 &	TSL:1 GENCODE basic
Chn1-202	ENSMUST00000102677.10	3760	<u>334aa</u>	Protein coding	CCDS16133 ₽	<u>Q91V57</u> ₽	TSL:1 GENCODE basic
Chn1-212	ENSMUST00000229731.1	1403	210aa	Protein coding	CCDS38145 ₽	A7VK14@Q91V57@	GENCODE basic
Chn1-209	ENSMUST00000166199.8	1324	<u>401aa</u>	Protein coding	CCDS50607 ₽	<u>A7VK13</u> ₽	TSL:1 GENCODE basic
Chn1-201	ENSMUST00000070579.6	1023	210aa	Protein coding	CCDS38145 ₽	A7VK14 @ Q91V57 @	TSL:1 GENCODE basic
Chn1-208	ENSMUST00000154258.7	712	<u>95aa</u>	Protein coding	-	B2FDI0 ₺	CDS 3' incomplete   TSL:5
Chn1-205	ENSMUST00000135904.1	631	<u>128aa</u>	Protein coding	121	B2FDI2 €	CDS 3' incomplete   TSL:2
Chn1-206	ENSMUST00000136953.1	413	81aa	Protein coding		B2FDI1 €	CDS 3' incomplete   TSL:3
Chn1-207	ENSMUST00000139252.1	881	<u>87aa</u>	Nonsense mediated decay	121	D6RCX8 ₽	TSL:3
Chn1-204	ENSMUST00000124450.1	643	<u>29aa</u>	Nonsense mediated decay	-	F7C3N6 ₽	CDS 5' incomplete   TSL:3
Chn1-215	ENSMUST00000231013.1	1544	No protein	Processed transcript	121	2	(22)
Chn1-214	ENSMUST00000230959.1	205	No protein	Processed transcript	-	1.70	lie!
Chn1-211	ENSMUST00000229312.1	20	No protein	Processed transcript	-	2	(22)
Chn1-213	ENSMUST00000229987.1	1431	No protein	Retained intron	-	-	(lei

The strategy is based on the design of *Chn1-203* transcript, the transcription is shown below:



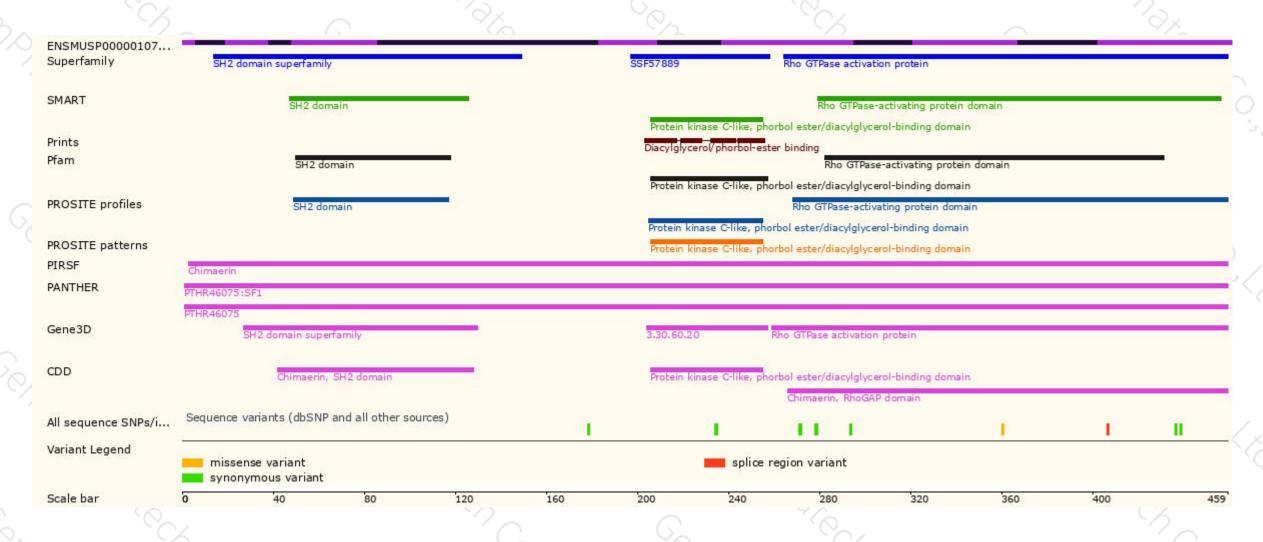
### Genomic location distribution





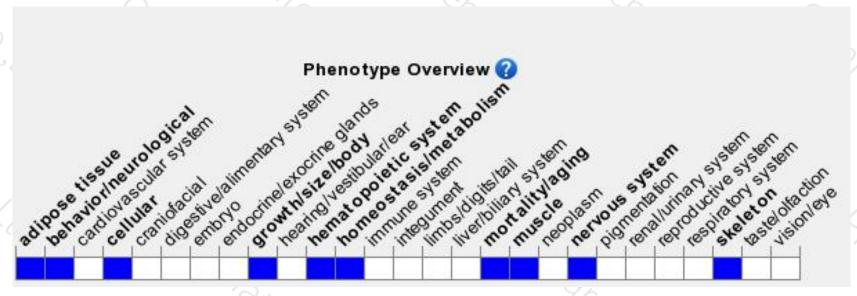
### Protein domain





## Mouse phenotype description(MGI)





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

According to the existing MGI data, mice homologous for a null allele exhibit transient postnatal size reduction, abnormal gait and abnormal innervation of the spinal cord. Part of null homozygous show preweaning lethality.



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





