

# *Clstn1* Cas9-KO Strategy

**Designer:**

**Huimin Su**

**Reviewer:**

**Ruirui Zhang**

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# Project Overview

**Project Name**

*Clstn1*

**Project type**

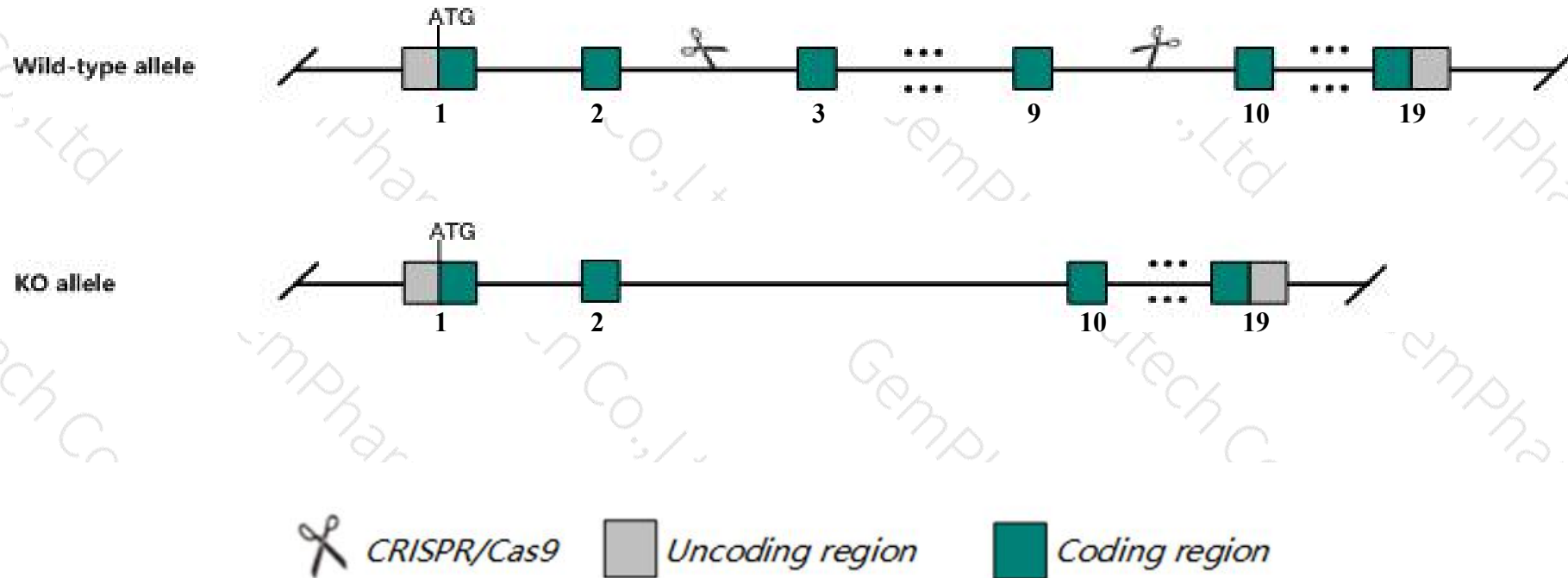
**Cas9-KO**

**Strain background**

**C57BL/6JGpt**

# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Clstn1* gene. The schematic diagram is as follows:



- The *Clstn1* gene has 4 transcripts. According to the structure of *Clstn1* gene, exon3-exon9 of *Clstn1-201* (ENSMUST00000039144.6) transcript is recommended as the knockout region. The region contains 1275bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Clstn1* gene. The brief process is as follows: CRISPR/Cas9 system w

- According to the existing MGI data, Juvenile mice homozygous for a null allele show reduced basal excitatory synaptic transmission, abnormal excitatory postsynaptic currents, enhanced NMDA receptor-dependent long term potentiation, and delayed dendritic spine maturation in CA1 hippocampal pyramidal cells.
- The *Clstn1* gene is located on the Chr4. 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)

## Clstn1 calsynenin 1 [ *Mus musculus* (house mouse) ]

Gene ID: 65945, updated on 5-Jan-2020

### Summary

**Official Symbol** Clstn1 provided by [MGI](#)  
**Official Full Name** calsynenin 1 provided by [MGI](#)  
**Primary source** [MGI:MGI:1929895](#)  
**See related** [Ensembl:ENSMUSG00000039953](#)  
**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** Cst-1; Cstn1; 1810034E21Rik  
**Expression** Broad expression in cortex adult (RPKM 93.0), frontal lobe adult (RPKM 80.3) and 23 other tissues [See more](#)  
**Orthologs** [human](#) [all](#)

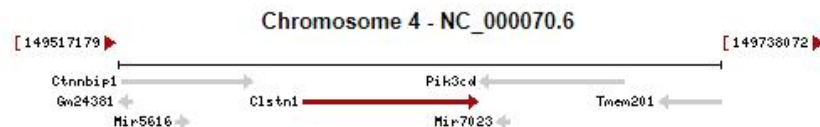
### Genomic context

Location: 4; 4 E2

See Clstn1 in [Genome Data Viewer](#)

Exon count: 21

Annotation release	Status	Assembly	Chr	Location
<a href="#">108</a>	current	GRCh38.p6 ( <a href="#">GCF_000001635.26</a> )	4	NC_000070.6 (149585111..149648899)
Build 37.2	previous assembly	MGSCv37 ( <a href="#">GCF_000001635.18</a> )	4	NC_000070.5 (148960747..149022008)



# Transcript information (Ensembl)

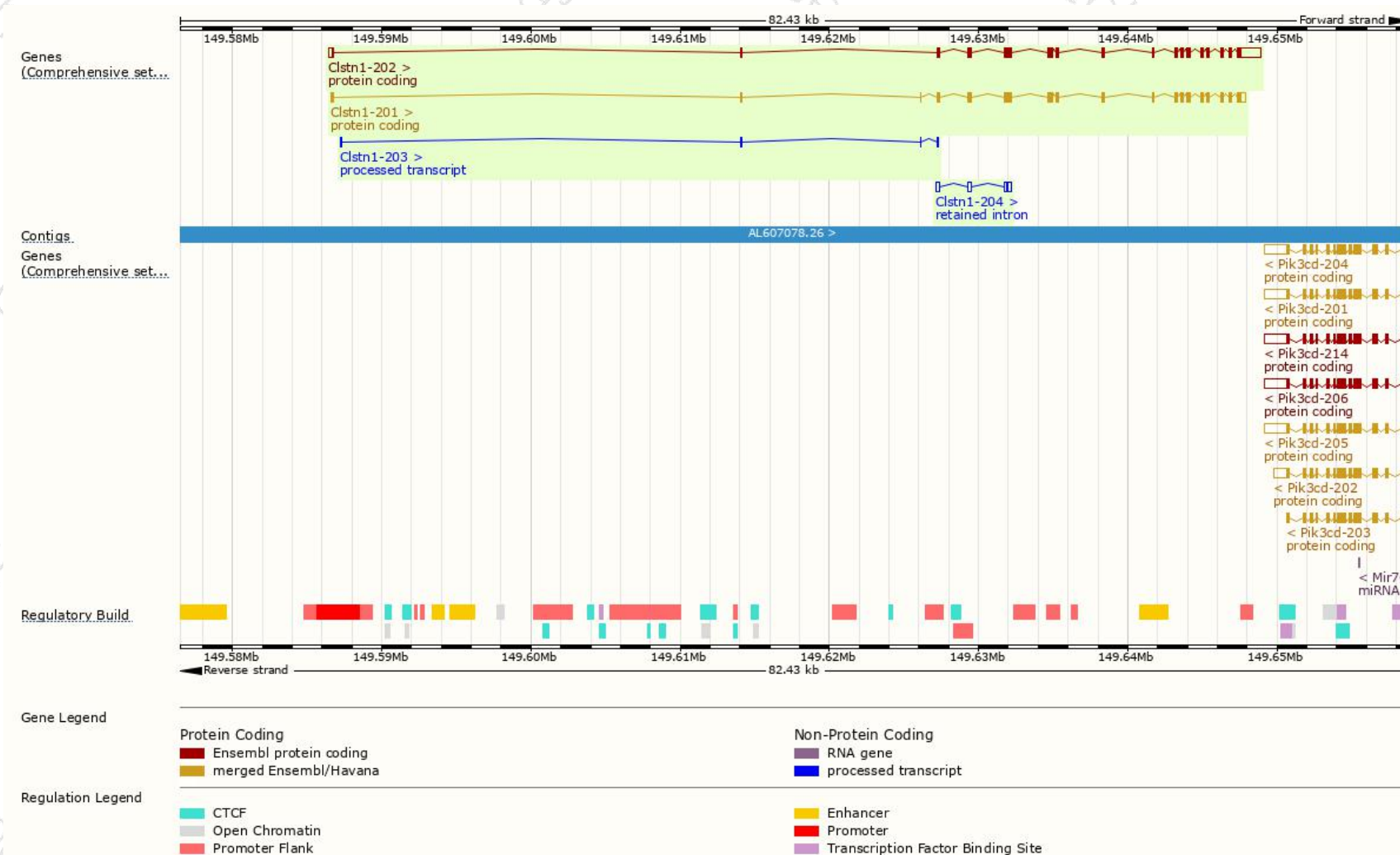
The gene has 4 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Clstn1-202	<a href="#">ENSMUST00000105691.7</a>	4459	<a href="#">969aa</a>	Protein coding	<a href="#">CCDS71522</a>	<a href="#">Q9EPL2</a>	TSL:1 GENCODE basic APPRIS ALT2
Clstn1-201	<a href="#">ENSMUST00000039144.6</a>	3319	<a href="#">979aa</a>	Protein coding	<a href="#">CCDS18963</a>	<a href="#">Q9EPL2</a>	TSL:1 GENCODE basic APPRIS P3
Clstn1-203	<a href="#">ENSMUST00000137232.1</a>	358	No protein	Processed transcript	-	-	TSL:3
Clstn1-204	<a href="#">ENSMUST00000151895.1</a>	822	No protein	Retained intron	-	-	TSL:2

The strategy is based on the design of *Clstn1-201* transcript,The transcription is shown below

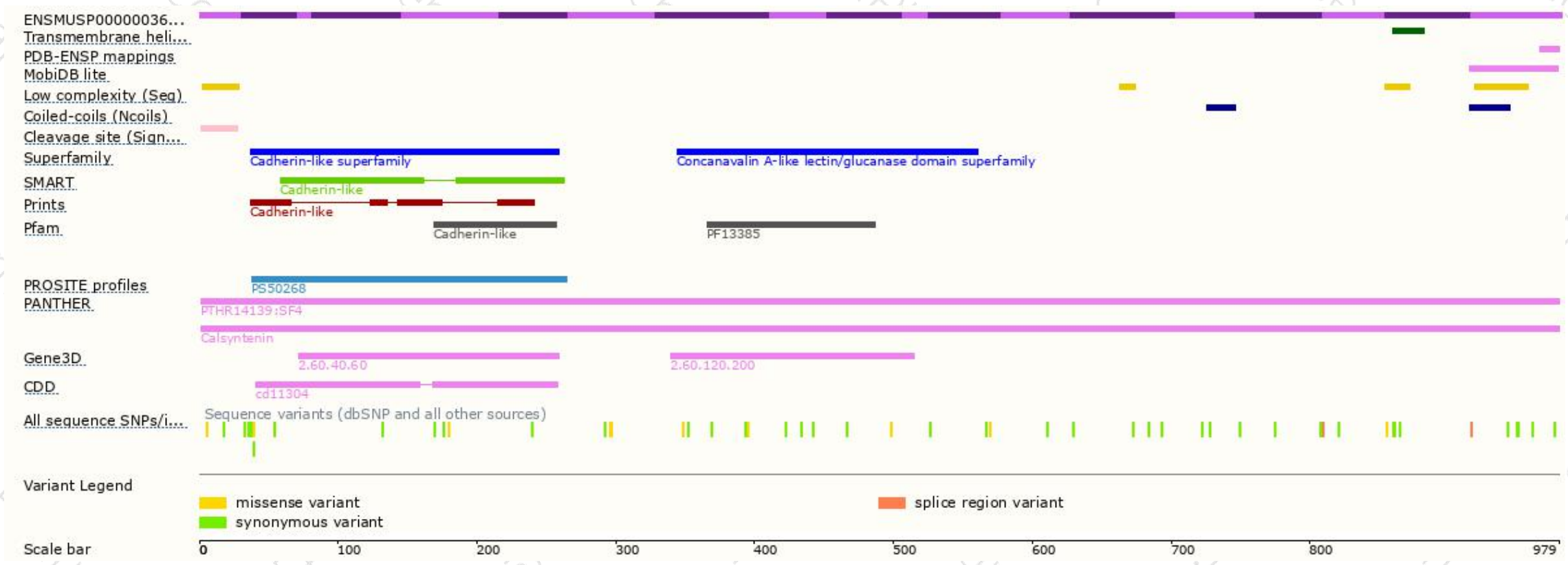


# Genomic location distribution

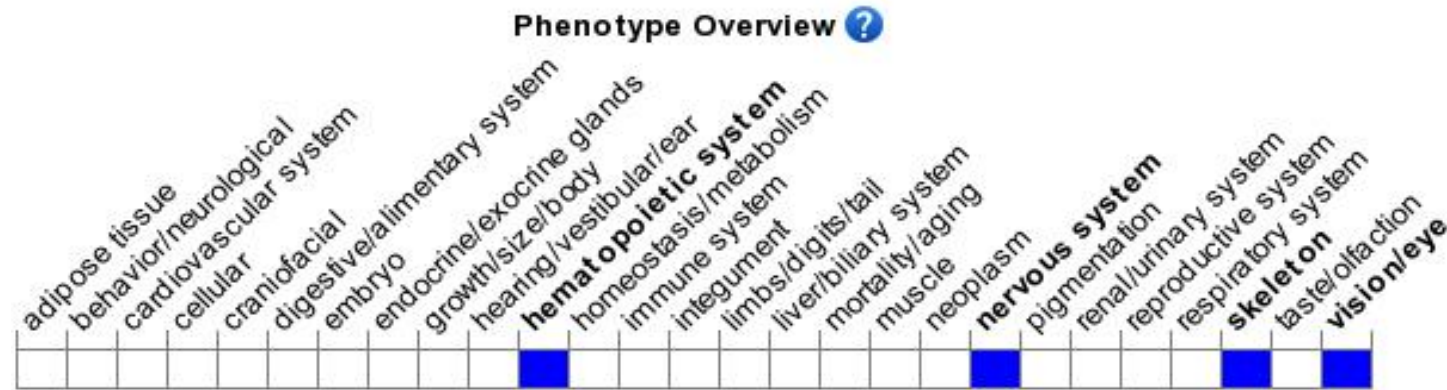




# Protein domain



# 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, Juvenile mice homozygous for a null allele show reduced basal excitatory synaptic transmission, abnormal excitatory postsynaptic currents, enhanced NMDA receptor-dependent long term potentiation, and delayed dendritic spine maturation in CA1 hippocampal pyramidal cells.

If you have any questions, you are welcome to inquire.

Tel: 400-9660890

