

Rnf13 Cas9-KO Strategy

Designer:	Huan Wang
Reviewer:	Huan Fan
Design Date:	2020-4-22

Project Overview

Project Name

Rnf13

Project type

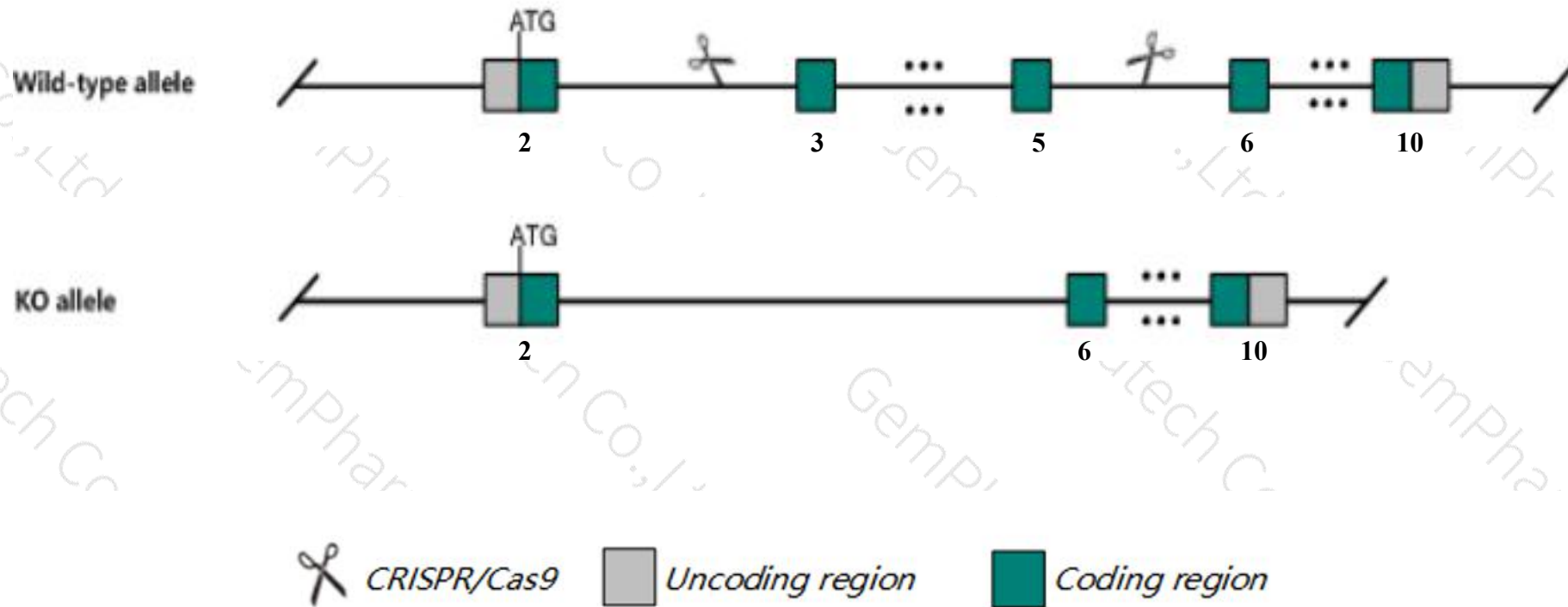
Cas9-KO

Strain background

C57BL/6JGpt

Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Rnf13* gene. The schematic diagram is as follows:



- The *Rnfl3* gene has 11 transcripts. According to the structure of *Rnfl3* gene, exon3-exon5 of *Rnfl3-201* (ENSMUST00000041826.13) transcript is recommended as the knockout region. The region contains 295bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Rnfl3* gene. The brief process is as follows: CRISPR/Cas9 system

- The *Rnf13* gene is located on the Chr3. 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)

Rnf13 ring finger protein 13 [Mus musculus (house mouse)]

Gene ID: 24017, updated on 13-Mar-2020

Summary



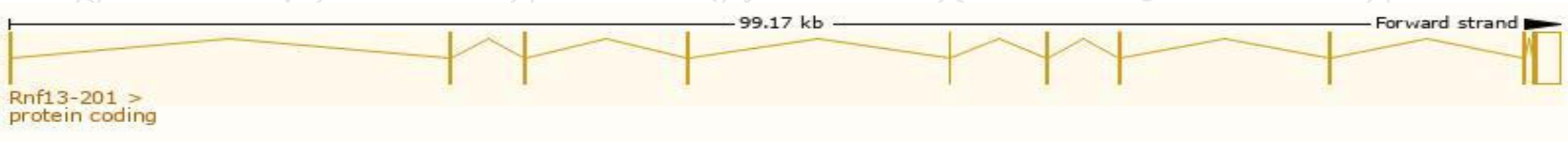
Official Symbol	Rnf13 provided by MGI
Official Full Name	ring finger protein 13 provided by MGI
Primary source	MGI:MGI:1346341
See related	Ensembl:ENSMUSG000000036503
Gene type	protein coding
RefSeq status	REVIEWED
Organism	Mus musculus
Lineage	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
Also known as	2010001H16Rik, Rzf
Summary	This gene encodes a member of the PA-TM-RING family of proteins that contain a protease associated (PA) domain and a RING finger domain separated by a transmembrane (TM) domain. The encoded protein is an E3 ubiquitin ligase localized to the endosomal-lysosomal vesicles and inner nuclear membrane. Mice lacking the encoded protein have impaired learning abilities associated with a decreased synaptic vesicle density and dysregulated SNARE complex assembly. Alternative splicing of this gene results in multiple transcript variants. A pseudogene for this gene has been identified on the X chromosome. [provided by RefSeq, Jan 2015]
Expression	Ubiquitous expression in cerebellum adult (RPKM 16.4), bladder adult (RPKM 14.8) and 28 other tissues See more
Orthologs	human all

Transcript information（Ensembl）

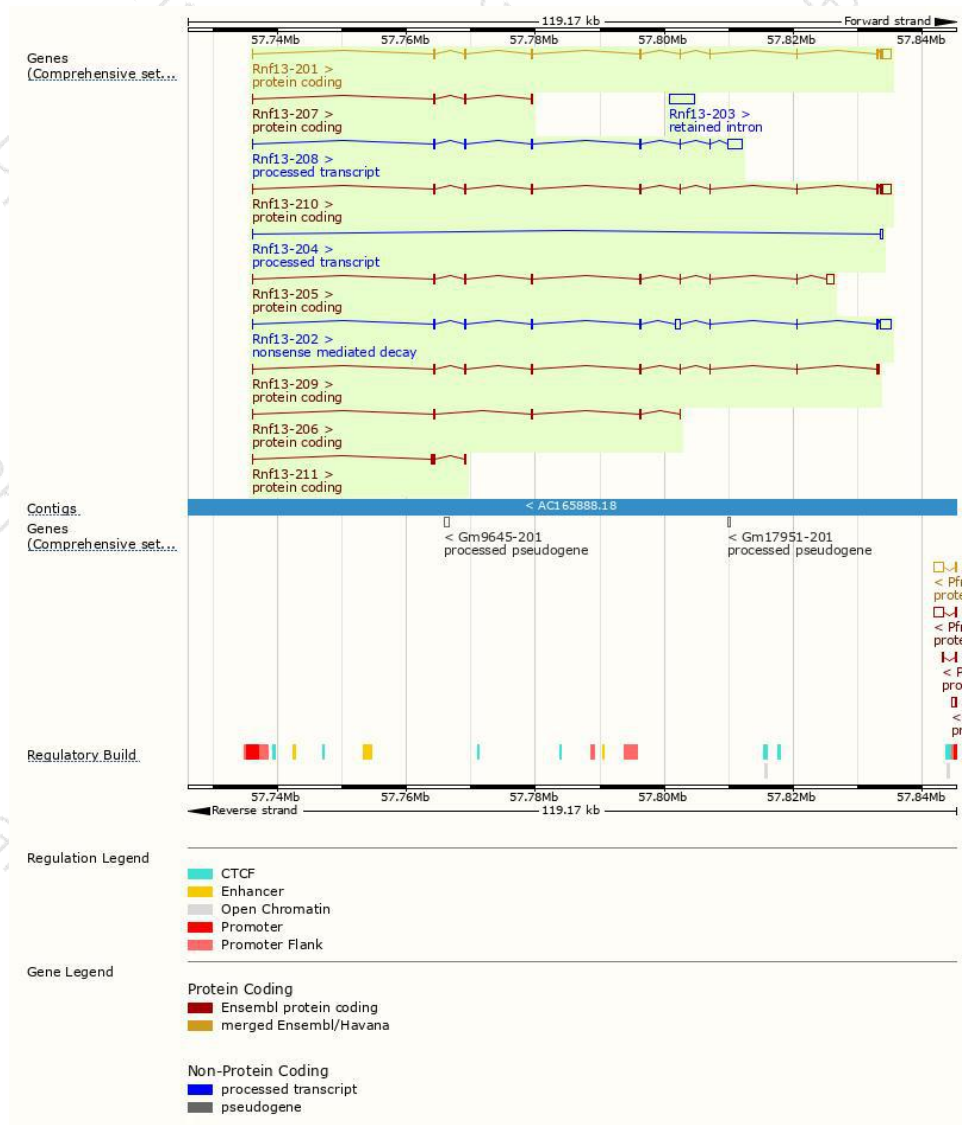
The gene has 11 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Rnf13-201	ENSMUST00000041826.13	2678	381aa	Protein coding	CCDS50913	Q54965	TSL:1 GENCODE basic APPRIS is a system to annotate alternatively spliced transcripts based on a range of computational methods to identify the most functionally important transcript(s) of a gene. APPRIS P1
Rnf13-210	ENSMUST000000200497.4	2573	352aa	Protein coding	CCDS84623	Q8CB78	TSL:1 GENCODE basic
Rnf13-205	ENSMUST000000198214.4	2027	244aa	Protein coding	CCDS84622	Q8C4F9	TSL:1 GENCODE basic
Rnf13-209	ENSMUST000000199041.1	1062	268aa	Protein coding	CCDS79910	Q54965_Q3UTG4	TSL:1 GENCODE basic
Rnf13-206	ENSMUST000000198249.4	486	131aa	Protein coding	-	A0A0G2JEM4	CDS 3' incomplete TSL:3
Rnf13-211	ENSMUST000000200600.4	352	64aa	Protein coding	-	A0A0G2JE17	CDS 3' incomplete TSL:3
Rnf13-207	ENSMUST000000198510.4	347	96aa	Protein coding	-	A0A0G2JDP6	CDS 3' incomplete TSL:2
Rnf13-202	ENSMUST000000197205.4	3244	166aa	Nonsense mediated decay	-	A0A0G2JGT7	TSL:1
Rnf13-208	ENSMUST000000198996.4	2848	No protein	Processed transcript	-	-	TSL:1
Rnf13-204	ENSMUST000000197920.4	435	No protein	Processed transcript	-	-	TSL:3
Rnf13-203	ENSMUST000000197783.1	3873	No protein	Retained intron	-	-	TSL:NA

The strategy is based on the design of *Rnf13-201* transcript,the transcription is shown below:



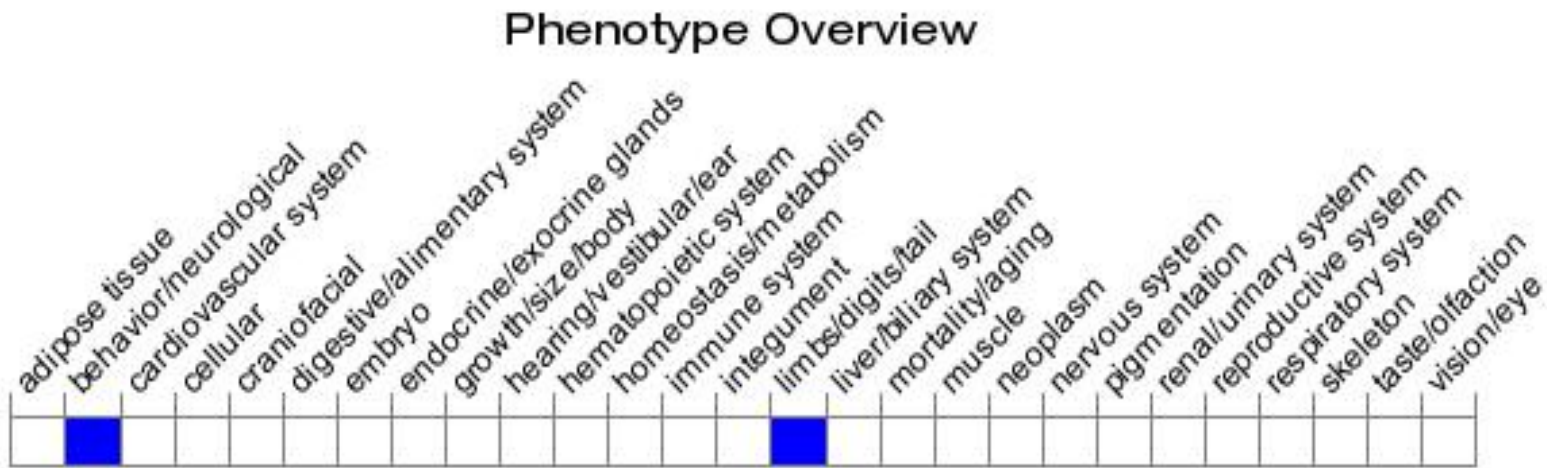
Genomic location distribution



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/>).

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

Tel: 400-9660890

