

# Mxra8 Cas9-KO Strategy

Designer: Yanhua Shen

Reviewer: Jia Yu

Design Date: 2023-08-2

# Overview

## Target Gene Name

- Mxra8

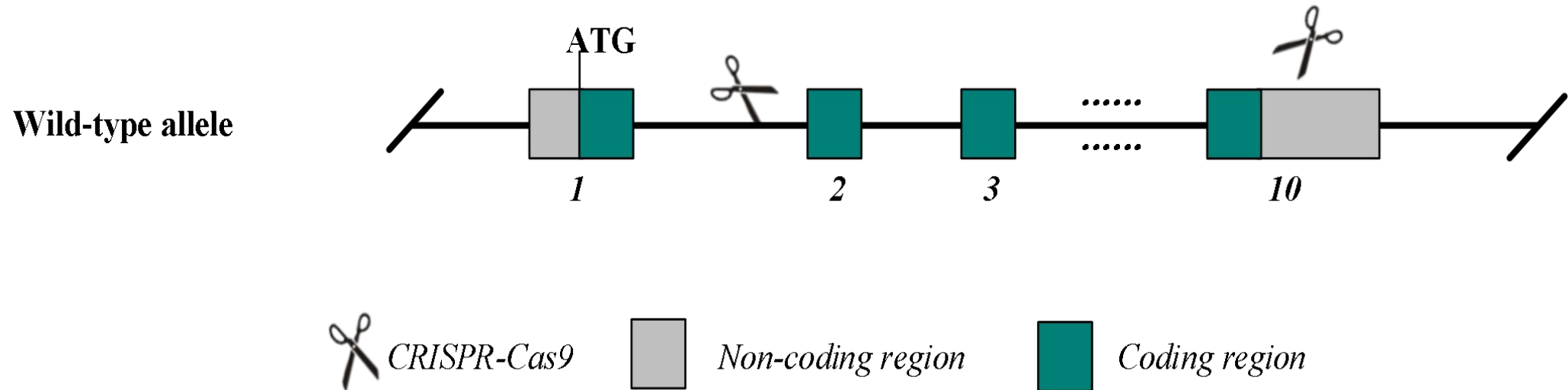
## Project Type

- Cas9-KO

## Genetic Background

- C57BL/6JGpt

# Strain Strategy



Schematic representation of CRISPR-Cas9 engineering used to edit the *Mxra8* gene.

# Technical Information

- The *Mxra8* gene has 7 transcripts. According to the structure of *Mxra8* gene, exon2-10 of *Mxra8*-201 (ENSMUST00000030947.4) transcript is recommended as the knockout region. The region contains most of coding sequences. Knocking out the region will result in disruption of protein function.
- In this project we use CRISPR-Cas9 technology to modify *Mxra8* gene. The brief process is as follows: gRNAs were transcribed in vitro. Cas9 and gRNAs 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 on-target amplicon sequencing. A stable F1-generation mouse strain was obtained by mating positive F0-generation mice with C57BL/6JGpt mice and confirmation of the desired mutant allele was carried out by PCR and on-target amplicon sequencing.

# Gene Information

## Mxra8 matrix-remodelling associated 8 [ *Mus musculus* (house mouse) ]

[Download Datasets](#)

Gene ID: 74761, updated on 21-Jun-2023

### Summary

<b>Official Symbol</b>	Mxra8 provided by <a href="#">MGI</a>
<b>Official Full Name</b>	matrix-remodelling associated 8 provided by <a href="#">MGI</a>
<b>Primary source</b>	<a href="#">MGI:MGI:1922011</a>
<b>See related</b>	<a href="#">Ensembl:ENSMUSG00000029070</a> <a href="#">AllianceGenome:MGI:1922011</a>
<b>Gene type</b>	protein coding
<b>RefSeq status</b>	VALIDATED
<b>Organism</b>	<a href="#">Mus musculus</a>
<b>Lineage</b>	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
<b>Also known as</b>	Asp3; Dicam; 1200013A08Rik; 1700095D18Rik
<b>Summary</b>	Acts upstream of or within establishment of glial blood-brain barrier. Located in cell surface. Is expressed in brain ventricle and choroid plexus; heart valve; and skeleton. Orthologous to human MXRA8 (matrix remodeling associated 8). [provided by Alliance of Genome Resources, Apr 2022]
<b>Expression</b>	Broad expression in lung adult (RPKM 383.7), ovary adult (RPKM 167.8) and 15 other tissues <a href="#">See more</a>
<b>Orthologs</b>	<a href="#">human</a> <a href="#">all</a>
<b>NEW</b>	Try the new <a href="#">Gene table</a> Try the new <a href="#">Transcript table</a>

### Genomic context

**Location:** 4 E2; 4 87.58 cM

**Exon count:** 10

See Mxra8 in [Genome Data Viewer](#)

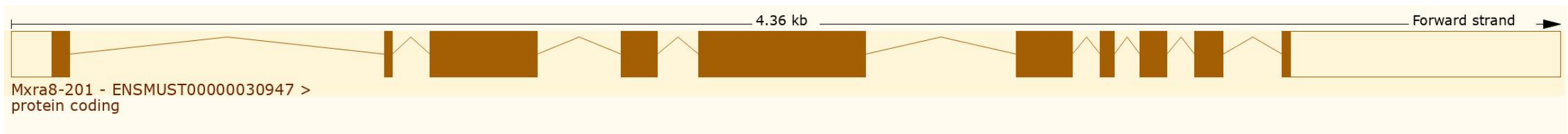
Source: <https://www.ncbi.nlm.nih.gov/>

# Transcript Information

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

Transcript ID	Name	bp	Protein	Biotype	CCDS	UniProt Match	Flags
<a href="#">ENSMUST00000030947.4</a>	Mxra8-201	2203	<a href="#">442aa</a>	Protein coding	<a href="#">CCDS19044</a>	<a href="#">Q9DBV4</a>	Ensembl Canonical GENCODE basic APPRIS P1 TSL:1
<a href="#">ENSMUST00000141883.8</a>	Mxra8-206	1097	<a href="#">311aa</a>	Protein coding		<a href="#">A2AD97</a>	TSL:2 CDS 3' incomplete
<a href="#">ENSMUST00000132142.2</a>	Mxra8-203	449	No protein	Protein coding CDS not defined		-	TSL:5
<a href="#">ENSMUST00000141766.8</a>	Mxra8-205	788	No protein	Retained intron		-	TSL:3
<a href="#">ENSMUST00000126487.2</a>	Mxra8-202	715	No protein	Retained intron		-	TSL:1
<a href="#">ENSMUST00000143886.2</a>	Mxra8-207	594	No protein	Retained intron		-	TSL:3
<a href="#">ENSMUST00000133592.2</a>	Mxra8-204	564	No protein	Retained intron		-	TSL:3

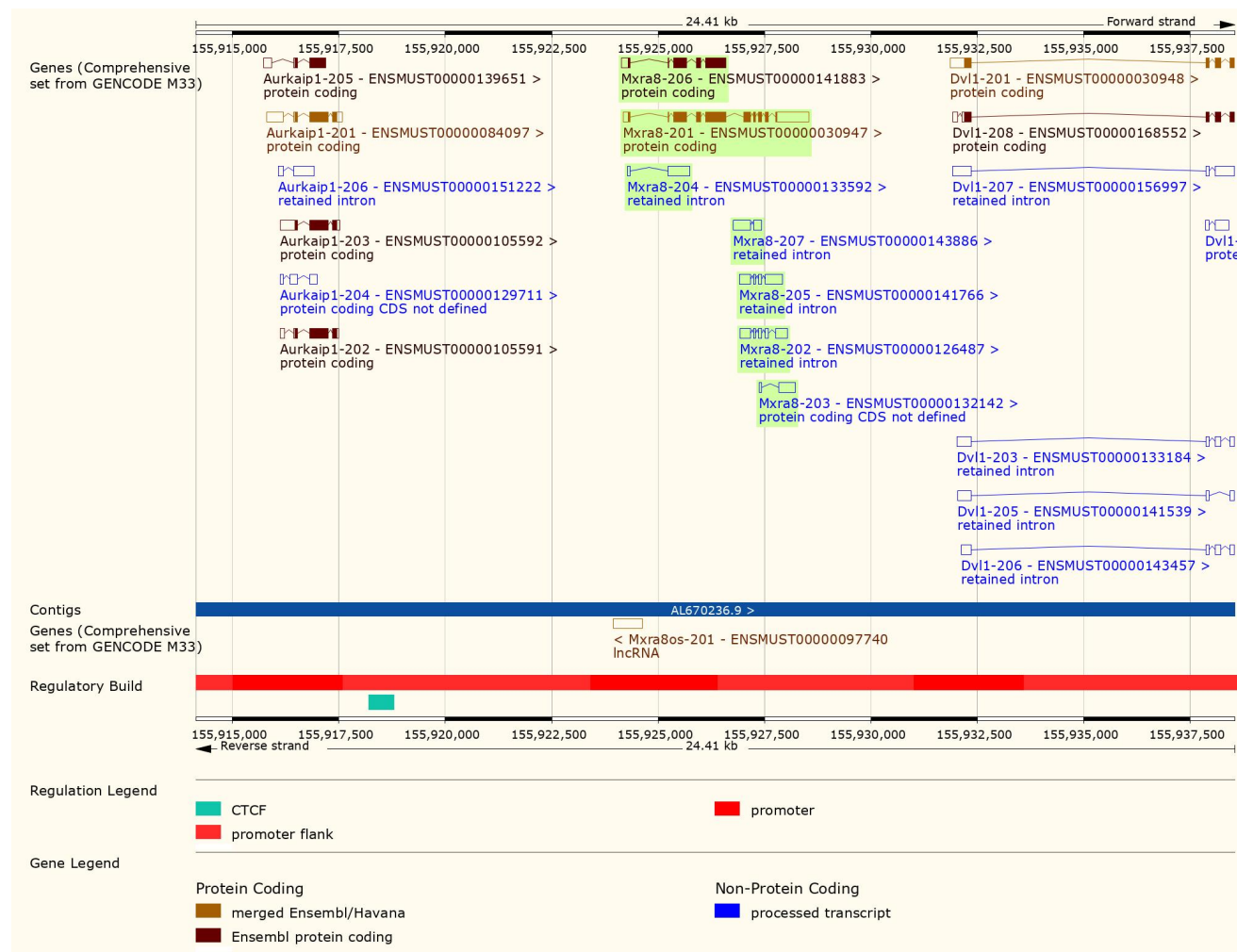
The strategy is based on the design of *Mxra8*-201 transcript, the transcription is shown below:



Source: <https://www.ensembl.org>



# Genomic Information

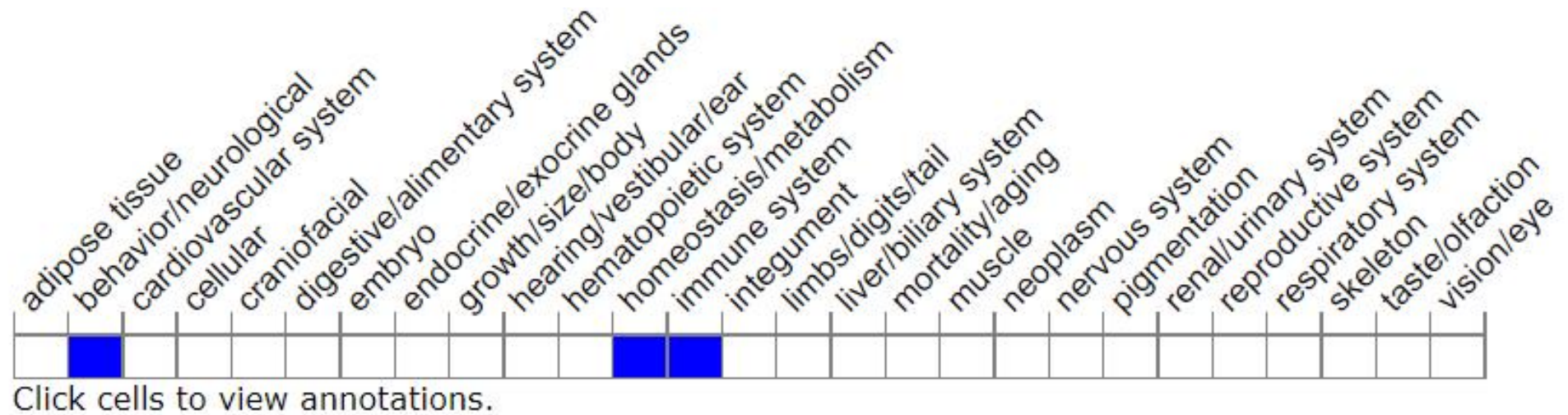


# Protein Information





# Mouse Phenotype Information (MGI)



- Phenotypic analysis of mice homozygous for a gene trap allele indicates this mutation has no notable phenotype in any parameter tested in a high-throughput screen. Mice homozygous for two alleles with intragenic deletions in the Ig-like domain 2 show decreased infection of musculoskeletal tissues with various alphaviruses.

# Important Information

- According theMGI data, phenotypic analysis of mice homozygous for a gene trap allele indicates this mutation has no notable phenotype in any parameter tested in a high-throughput screen. Mice homozygous for two alleles with intragenic deletions in the Ig-like domain 2 show decreased infection of musculoskeletal tissues with various alphaviruses.
- The effect of *Mxra8os*-201 and *Dvll*-201 is unknown.
- *Mxra8* is located on Chr4. If the knockout mice are crossed with other mouse strains to obtain double homozygous mutant offspring, please avoid the situation that the second gene is on the same chromosome.
- This Strategy is designed based on genetic information in existing databases. Due to the complexity of biological processes, all risks of the mutation on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.