

# Map2k7 Cas9-KO Strategy

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# **Project Overview**



**Project Name** 

Map2k7

**Project type** 

Cas9-KO

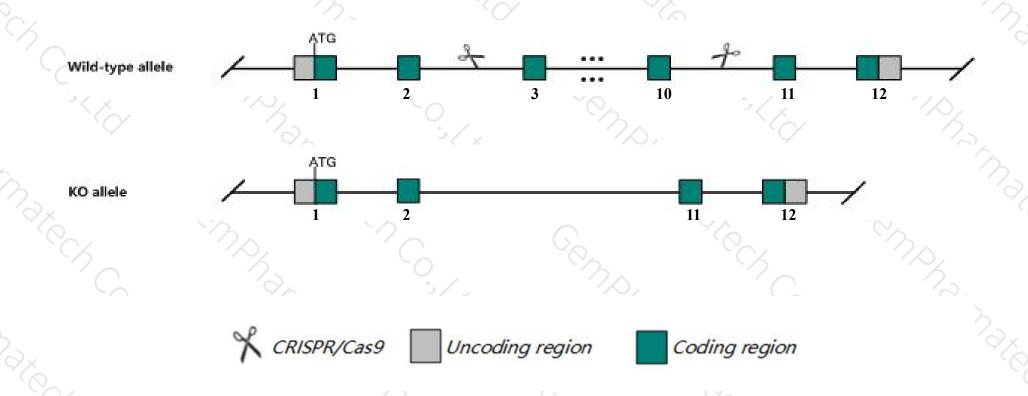
Strain background

C57BL/6JGpt

# **Knockout strategy**



This model will use CRISPR/Cas9 technology to edit the Map2k7 gene. The schematic diagram is as follows:



### **Technical routes**



- ➤ The *Map2k7* gene has 10 transcripts. According to the structure of *Map2k7* gene, exon3-exon10 of *Map2k7-201* (ENSMUST0000003027.13) transcript is recommended as the knockout region. The region contains 955bp coding sequence Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Map2k7* gene. The brief process is as follows: CRISPR/Cas9 syste

### **Notice**



- > According to the existing MGI data, Mice homozygous for disruptions in this gene die during embryogenesis.
- ➤ The konckout region near to the 5'UTR of *Tgfbr3l* gene. Knockout the region may affect the regulatory function of the 5'UTR of *Tgfbr3l* gene.
- ightharpoonup The Gm49320 gene will be deleted together in this strategy.
- The *Map2k7* gene is located on the Chr8. 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)



#### Map2k7 mitogen-activated protein kinase kinase 7 [Mus musculus (house mouse)]

Gene ID: 26400, updated on 23-Feb-2019

#### Summary

☆ ?

Official Symbol Map2k7 provided by MGI

Official Full Name mitogen-activated protein kinase kinase 7 provided by MGI

Primary source MGI:MGI:1346871

See related Ensembl:ENSMUSG00000002948

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 5930412N11Rik, JNKK 2, Jnkk2, MAPKK 7, MEK 7, Mapkk7, Mek7, Mkk7, Prkmk7, sek2

Expression Ubiquitous expression in testis adult (RPKM 34.8), adrenal adult (RPKM 26.2) and 28 other tissuesSee more

Orthologs <u>human</u> all

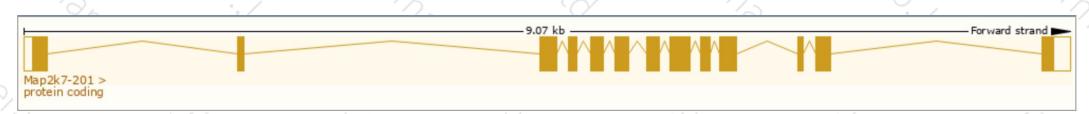
# Transcript information (Ensembl)



The gene has 10 transcripts, all transcripts are shown below:

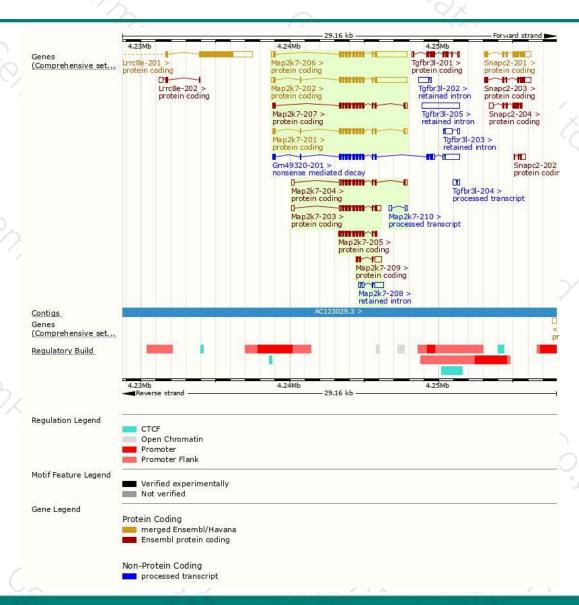
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Map2k7-202	ENSMUST00000062686.10	3525	<u>435aa</u>	Protein coding	CCDS52474	Q8CE90	TSL:1 GENCODE basic APPRIS ALT1
Map2k7-206	ENSMUST00000110998.8	3497	419aa	Protein coding	CCDS40209	Q8CE90	TSL:1 GENCODE basic APPRIS P3
Map2k7-201	ENSMUST00000003027.13	1626	<u>468aa</u>	Protein coding	CCDS40208	Q8CE90	TSL:1 GENCODE basic
Map2k7-203	ENSMUST00000110994.8	1600	346aa	Protein coding	CCDS80854	Q8CE90	TSL:1 GENCODE basic
/lap2k7-207	ENSMUST00000110999.7	1578	452aa	Protein coding	CCDS80852	Q8CE90	TSL:1 GENCODE basic
Map2k7-204	ENSMUST00000110995.7	1558	<u>379aa</u>	Protein coding	CCDS80853	Q8CE90	TSL:1 GENCODE basic
Map2k7-205	ENSMUST00000110996.1	1188	<u>391aa</u>	Protein coding	-	Q8CE90	TSL:1 GENCODE basic
Map2k7-209	ENSMUST00000129866.7	760	<u>79aa</u>	Protein coding	-	A0A140LHN8	CDS 5' incomplete TSL:3
Map2k7-210	ENSMUST00000207247.1	324	No protein	Processed transcript	5	150	TSL:2
Map2k7-208	ENSMUST00000129537.1	934	No protein	Retained intron			TSL:2

The strategy is based on the design of Map2k7-201 transcript, The transcription is shown below



### Genomic location distribution





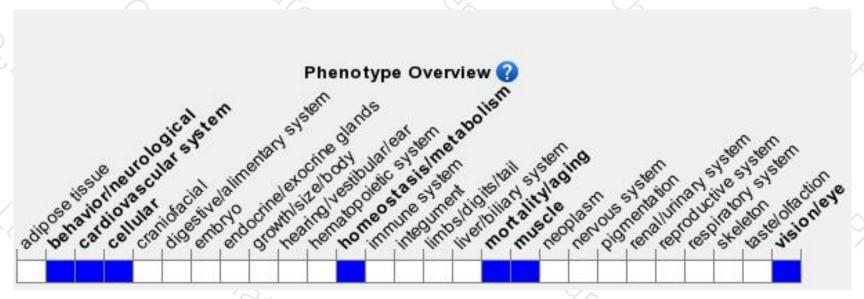
### Protein domain





## 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, Mice homozygous for disruptions in this gene die during embryogenesis.



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





