

# Sema5a Cas9-KO Strategy

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Reviewer: Rui Xiong

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# **Project Overview**



**Project Name** 

Sema5a

**Project type** 

Cas9-KO

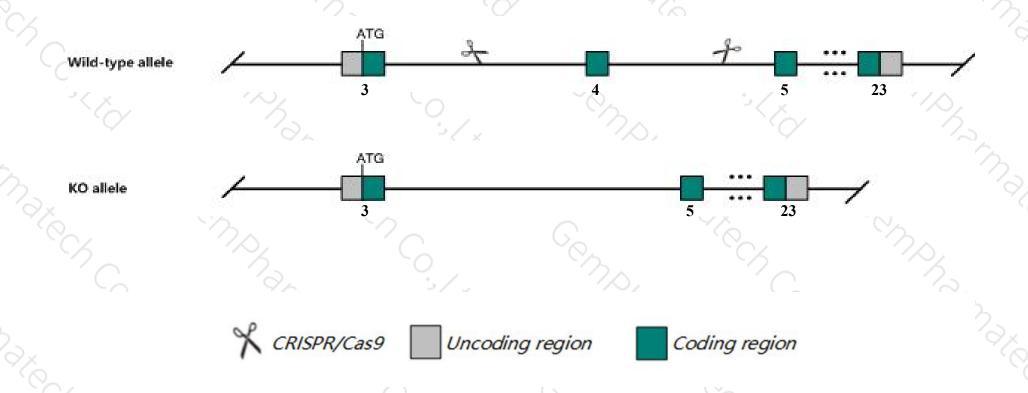
Strain background

C57BL/6JGpt

# **Knockout strategy**



This model will use CRISPR/Cas9 technology to edit the Sema5a gene. The schematic diagram is as follows:



### **Technical routes**



- ➤ The *Sema5a* gene has 9 transcripts. According to the structure of *Sema5a* gene, exon4 of *Sema5a-201* (ENSMUST00000067458.6) transcript is recommended as the knockout region. The region contains 100bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Sema5a* gene. The brief process is as follows: CRISPR/Cas9 syste

### **Notice**



- ➤ According to the existing MGI data, Mice homozygous for one null mutation die during organogenesis and display defects in branching of cranial vessels. Mice homozygous for another null mutation appear normal.
- The *Sema5a* gene is located on the Chr15. 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)



Sema5a sema domain, seven thrombospondin repeats (type 1 and type 1-like), transmembrane domain (TM) and short cytoplasmic domain, (semaphorin) 5A [Mus musculus (house mouse)]

Gene ID: 20356, updated on 12-Mar-2019

#### Summary

♠ ?

Official Symbol Sema5a provided by MGI

Official Full Name sema domain, seven thrombospondin repeats (type 1 and type 1-like), transmembrane domain (TM) and short cytoplasmic domain,

(semaphorin) 5A provided by MGI

Primary source MGI:MGI:107556

See related Ensembl:ENSMUSG00000022231

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 5930434A13, 9130201M22Rik, Al464145, Semaf, semF

Expression Broad expression in ovary adult (RPKM 13.8), limb E14.5 (RPKM 11.9) and 21 other tissuesSee more

Orthologs <u>human all</u>

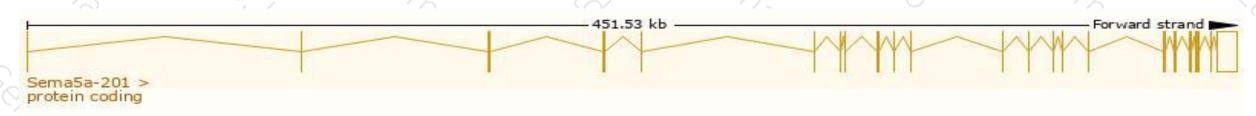
# Transcript information (Ensembl)



The gene has 9 transcripts, all transcripts are shown below:

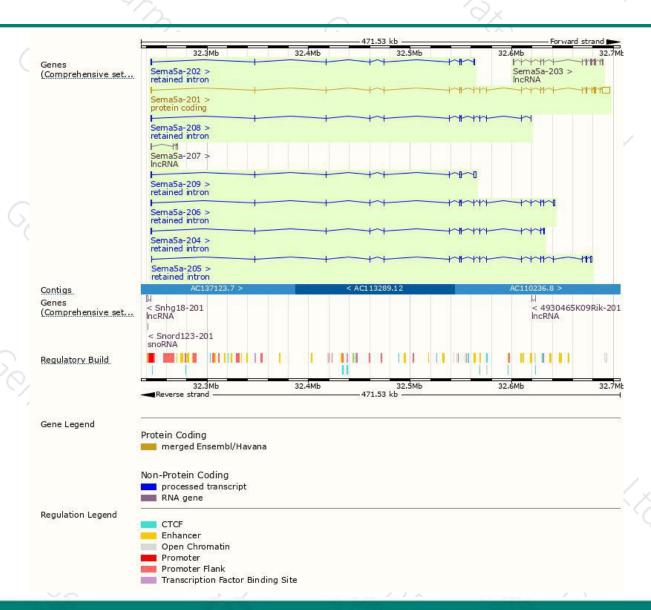
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Sema5a-201	ENSMUST00000067458.6	10809	<u>1074aa</u>	Protein coding	CCDS37055	Q3UPZ0	TSL:1 GENCODE basic APPRIS P1
Sema5a-209	ENSMUST00000228555.1	5442	No protein	Retained intron	-	-	
Sema5a-206	ENSMUST00000228015.1	3951	No protein	Retained intron		(4)	
Sema5a-205	ENSMUST00000227976.1	3725	No protein	Retained intron	2	1525	
Sema5a-204	ENSMUST00000227802.1	3158	No protein	Retained intron			
Sema5a-208	ENSMUST00000228442.1	2571	No protein	Retained intron	-	-	
Sema5a-202	ENSMUST00000226876.1	2375	No protein	Retained intron	9	(4)	
Sema5a-203	ENSMUST00000227429.1	2774	No protein	IncRNA	2	3528	
Sema5a-207	ENSMUST00000228103.1	902	No protein	IncRNA	5	121	

The strategy is based on the design of Sema5a-201 transcript, The transcription is shown below



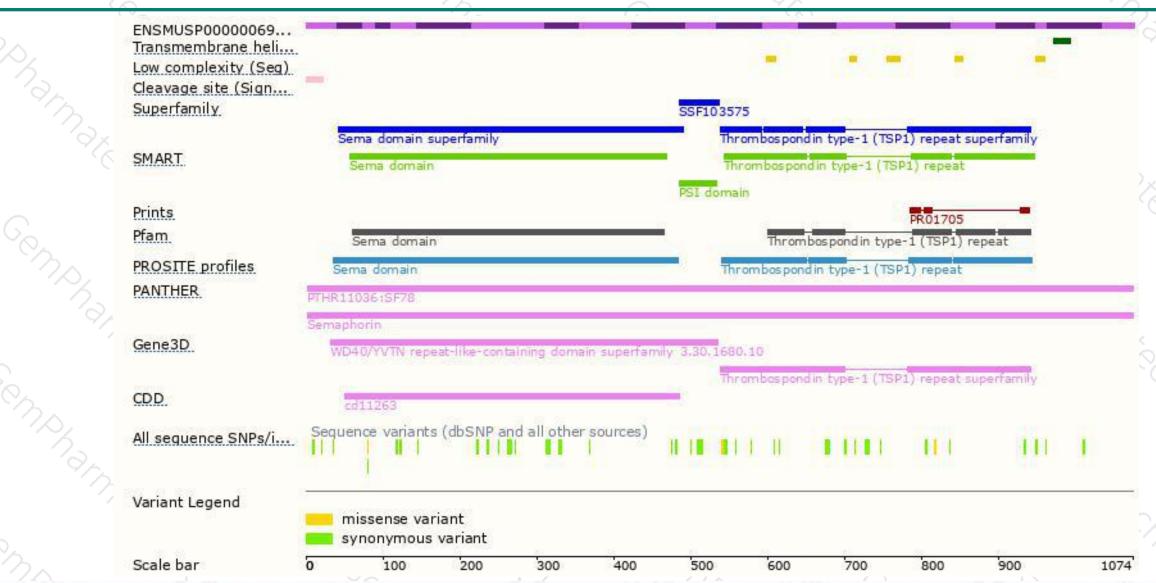
## Genomic location distribution





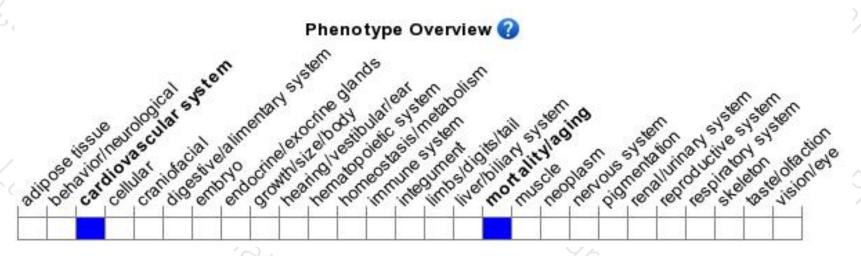
### Protein domain





# 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 one null mutation die during organogenesis and display defects in branching of cranial vessels. Mice homozygous for another null mutation appear normal.



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





