

# Mcm9 Cas9-KO Strategy

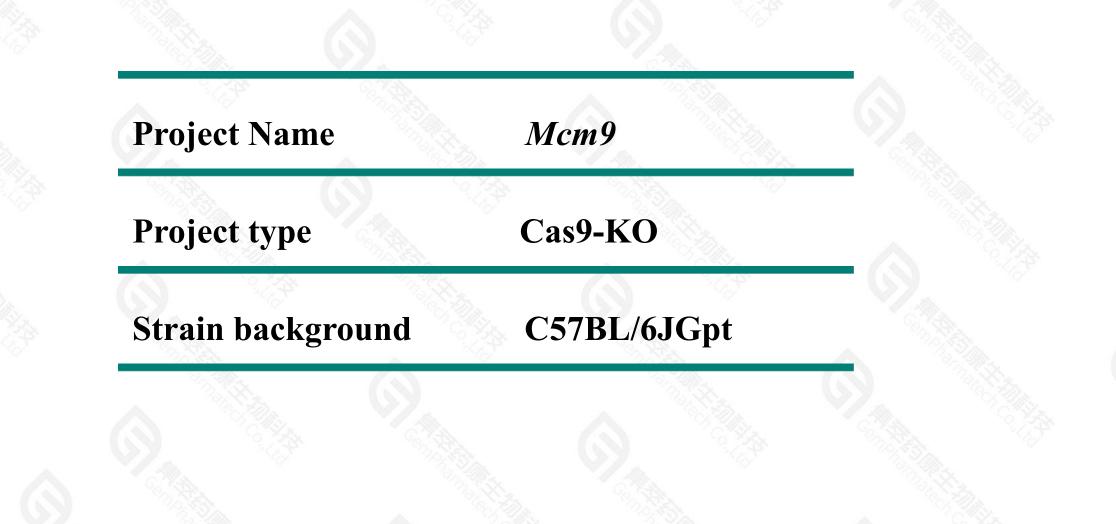
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**Reviewer: Miaomiao Cui** 

**Design Date: 2021-5-26** 

# **Project Overview**





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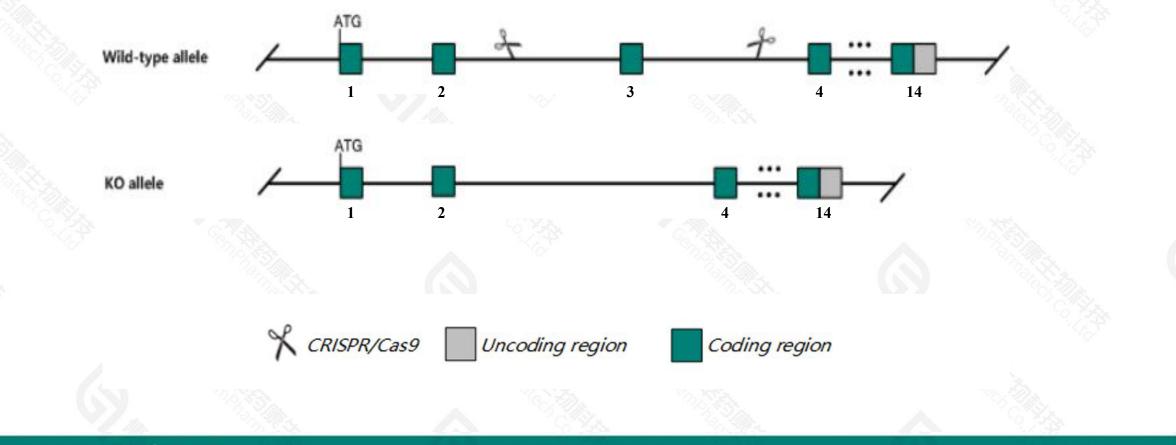
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# **Knockout strategy**



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This model will use CRISPR/Cas9 technology to edit the *Mcm9* gene. The schematic diagram is as follows:



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> The *Mcm9* gene has 8 transcripts. According to the structure of *Mcm9* gene, exon3 of *Mcm9*-201(ENSMUST00000075540.8) transcript is recommended as the knockout region. The region contains 319bp coding sequence. Knock out the region will result in disruption of protein function.

➤ In this project we use CRISPR/Cas9 technology to modify *Mcm9* gene. The brief process is as follows: CRISPR/Cas9 system 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 sequencing. A stable F1 generation mouse model was obtained by mating positive F0 generation mice with C57BL/6JGpt mice.



- > According to the existing MGI data,mice homozygous for gene trap alleles display germ cell loss with reduced fertility or infertility and increased tumor incidence, particularly of hepatocellular carcinomas.
- ➤ Transcript *Mcm9*-203&205&206 may not be affected.
- > The *Mcm9* gene is located on the Chr10. 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)**

### Mcm9 minichromosome maintenance 9 homologous recombination repair factor [Mus musculus (house mouse)]

Gene ID: 71567, updated on 17-Feb-2021

#### Summary

Official Symbol	Mcm9 provided by MGI
<b>Official Full Name</b>	minichromosome maintenance 9 homologous recombination repair factor provided by MGI
Primary source	MGI:MGI:1918817
See related	Ensembl:ENSMUSG0000058298
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	9030408017Rik, Gm235, Mcmdc1
Expression	Ubiquitous expression in CNS E11.5 (RPKM 3.4), liver E14 (RPKM 2.7) and 28 other tissuesSee more
Orthologs	human all

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# **Transcript information (Ensembl)**



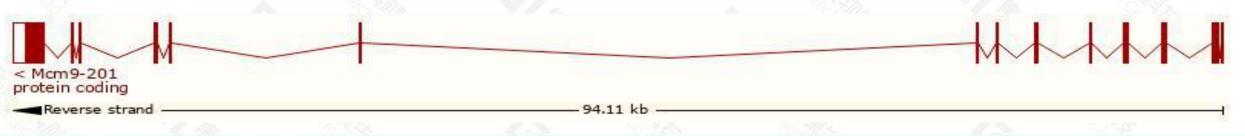
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### The gene has 8 transcripts, all transcripts are shown below:

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Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags	
Mcm9-201	ENSMUST0000075540.8	4868	<u>1290aa</u>	Protein coding	CCD523846		TSL:2, GENCODE basic, APPRIS P2,	
Mcm9-208	ENSMUST00000237608.2	4882	<u>1134aa</u>	Protein coding	-		GENCODE basic , APPRIS ALT2 ,	
Mcm9-204	ENSMUST00000219838.3	4132	<u>386aa</u>	Protein coding	2		TSL:5 , GENCODE basic ,	
Mcm9-206	ENSMUST00000220007.2	3513	<u>552aa</u>	Protein coding			TSL:1 , GENCODE basic ,	
Mcm9-203	ENSMUST00000219547.2	3168	<u>552aa</u>	Protein coding	¥		TSL:1 , GENCODE basic ,	
Mcm9-202	ENSMUST00000218549.3	2421	<u>414aa</u>	Protein coding	8		TSL:1 , GENCODE basic ,	
Mcm9-205	ENSMUST00000219841.3	307	<u>98aa</u>	Protein coding	-		CDS 5' incomplete , TSL:5 ,	
Mcm9-207	ENSMUST00000220260.2	464	No protein	Processed transcript	2		TSL:5,	
S	VX XX ALA	1			1			

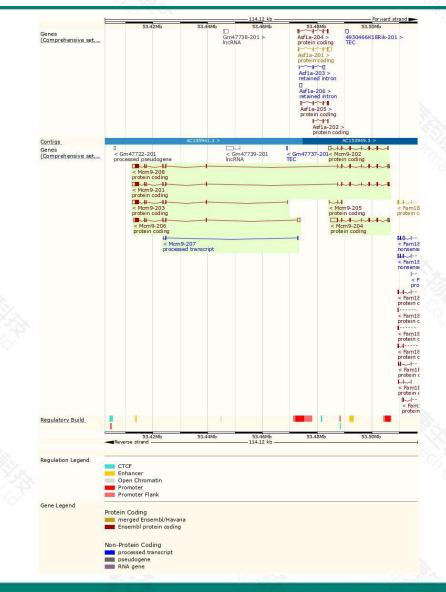
The strategy is based on the design of Mcm9-201 transcript, the transcription is shown below:



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### **Genomic location distribution**





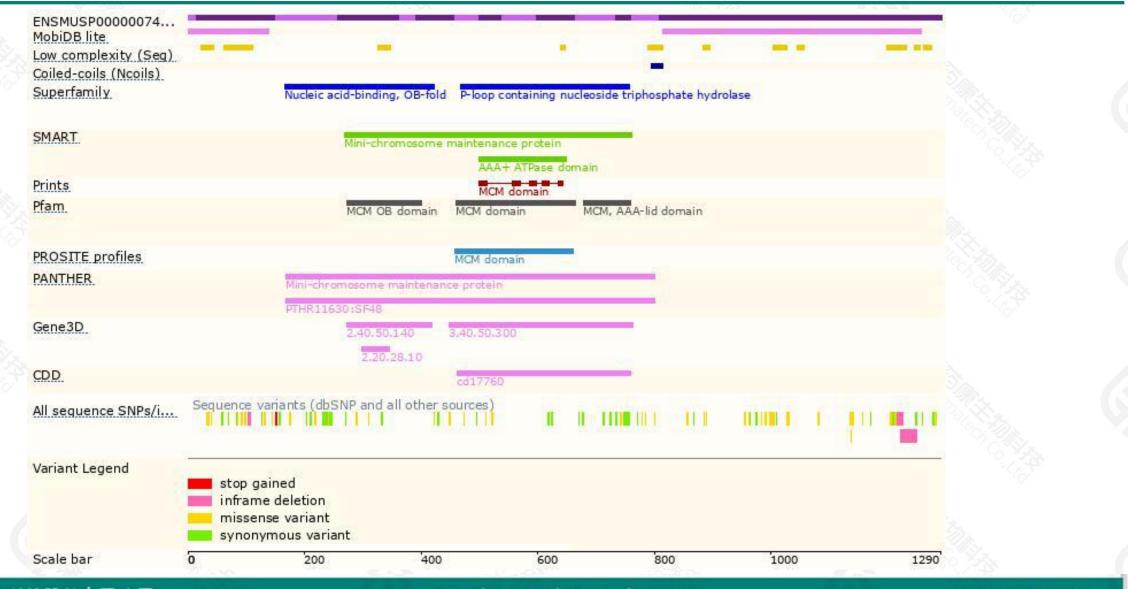
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### **Protein domain**



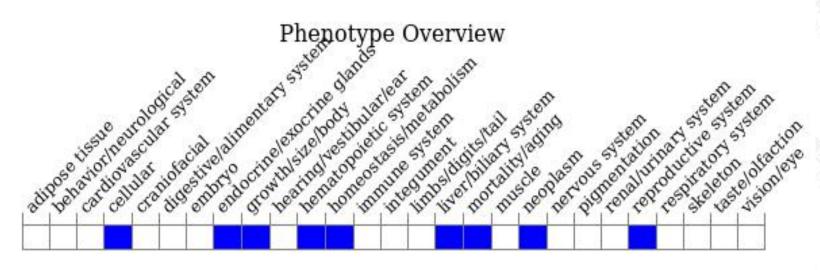


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### Mouse phenotype description(MGI)



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 gene trap alleles display germ cell loss with reduced fertility or infertility and increased tumor incidence, particulary of hepatocellular carcinomas.



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



