

Cmtm7 Cas9-KO Strategy

Designer:

Reviewer:

Design Date:

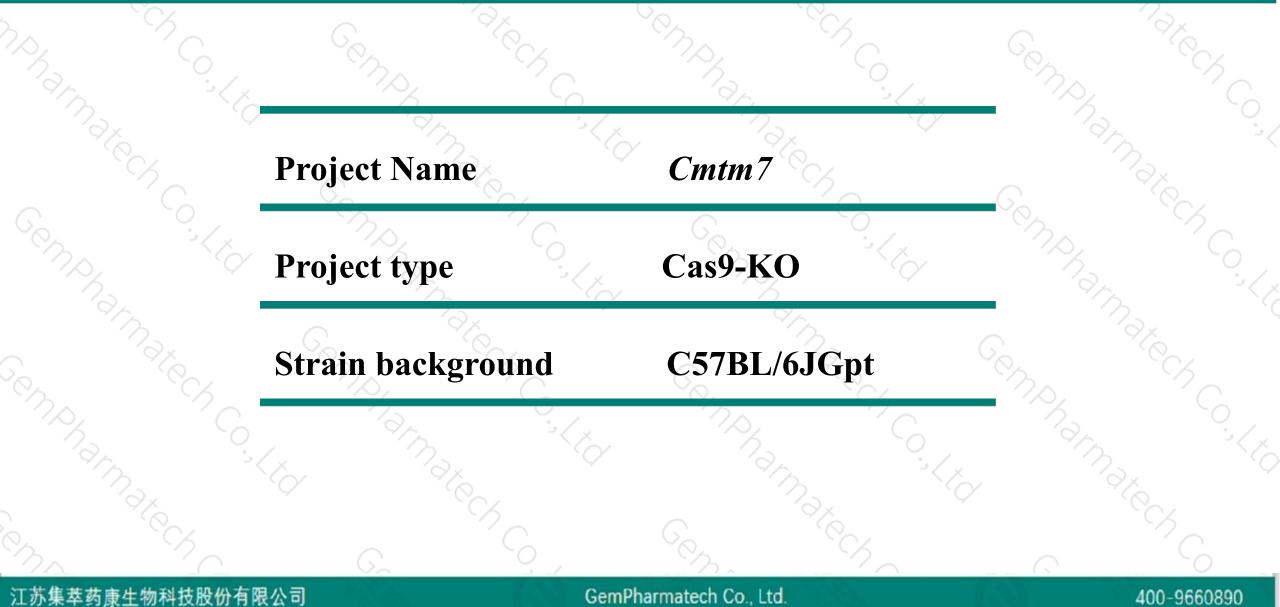
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Project Overview





Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Cmtm7 gene. The schematic diagram is as follows:

ATG Wild-type allele KO allele CRISPR/Cas9 Uncoding region Coding region

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- The Cmtm7 gene has 5 transcripts. According to the structure of Cmtm7 gene, exon1-exon5 of Cmtm7-201 (ENSMUST00000035009.15) transcript is recommended as the knockout region. The region contains all of the coding sequence. Knock out the region will result in disruption of protein function.
- > In this project we use CRISPR/Cas9 technology to modify Cmtm7 gene. The brief process is as follows: CRISPR/Cas9 system

- According to the existing MGI data, heterozygous mice die shortly after birth. Null mutations in this gene are haploinsufficient. a chimeric mouse showed defects in b-1a cell numbers and physiology.
- > Gm9888-201 gene may be destroyed.

Notice

- The Cmtm7 gene is located on the Chr9. 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)



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Cmtm7 CKLF-like MARVEL transmembrane domain containing 7 [Mus musculus (house mouse)]

Gene ID: 102545, updated on 13-Mar-2020

Summary

Official Symbol	Cmtm7 provided by MGI
Official Full Name	CKLF-like MARVEL transmembrane domain containing 7 provided by MGI
Primary source	MGI:MGI:2447166
See related	Ensembl:ENSMUSG0000032436
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	AI481279, Cklfsf7, LNV
Expression	Broad expression in thymus adult (RPKM 190.5), stomach adult (RPKM 118.3) and 20 other tissuesSee more
Orthologs	human all

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Transcript information (Ensembl)



The gene has 5 transcripts, all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags	
Cmtm7-201	ENSMUST0000035009.15	993	<u>167aa</u>	Protein coding	CCDS57710	Q9ESD6	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 F	21
Cmtm7-202	ENSMUST0000084867.8	874	<u>134aa</u>	Protein coding	CCDS57709	Q9ESD6	TSL:1 GENCODE basic	
Cmtm7-204	ENSMUST00000216760.1	665	<u>63aa</u>	Protein coding		A0A1L1SRR1	TSL:1 GENCODE basic	
Cmtm7-205	ENSMUST00000217056.1	345	No protein	Processed transcript	-	1923	TSL:3	
Cmtm7-203	ENSMUST00000214604.1	2183	No protein	Retained intron		173	TSL:1	
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The strategy is based on the design of *Cmtm7-201* transcript, the transcription is shown below:

< Cmtm7-201 protein coding

Reverse strand

- 25.01 kb -

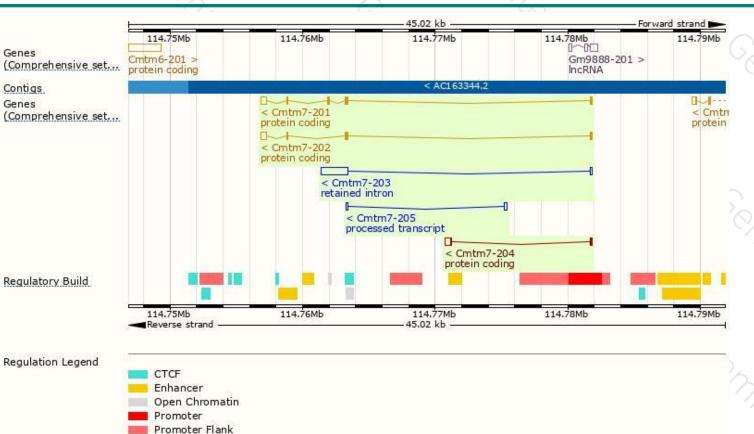
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Genomic location distribution







Gene Legend

Protein Coding Ensembl protein coding

merged Ensembl/Havana

Non-Protein Coding processed transcript RNA gene

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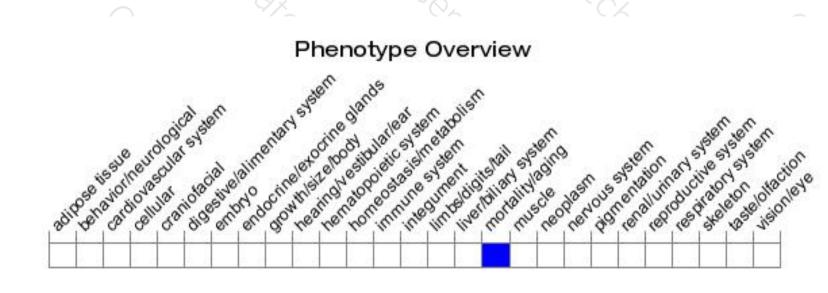
Protein domain



ENSMUSP0000035 Transmembrane heli Low complexity (Seq) Pfam. PROSITE profiles PANTHER All sequence SNPs/i	PTHR22776:SF31 PTHR22776	Marvel domain Marvel domain					
PANTHER.	PTHR22776	Marvel domain					
All sequence SNPs/i							
		dbSNP and all oth	er sources)	CL L	() ()		•
Variant Legend	missense varia synonymous va						2
Scale bar o	20	40	60	80	100	120 140	167 2 2 2 3 2 3 3 3 3 3 3 3 3 3 3 3 3 3 3

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, heterozygous mice die shortly after birth. Null mutations in this gene are haploinsufficient. A chimeric mouse showed defects in B-1a cell numbers and physiology.



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



