

Prmt2 Cas9-KO Strategy

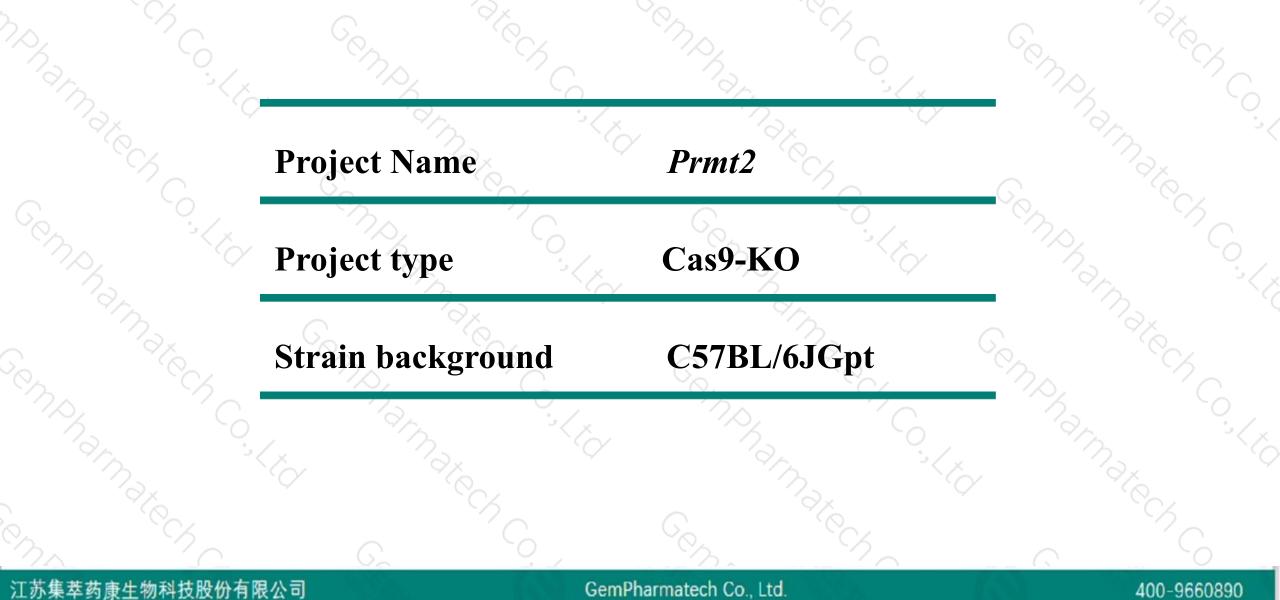
Designer: Reviewer:

Design Date:

Longyun Hu Yun Li 2020-2-18

Project Overview

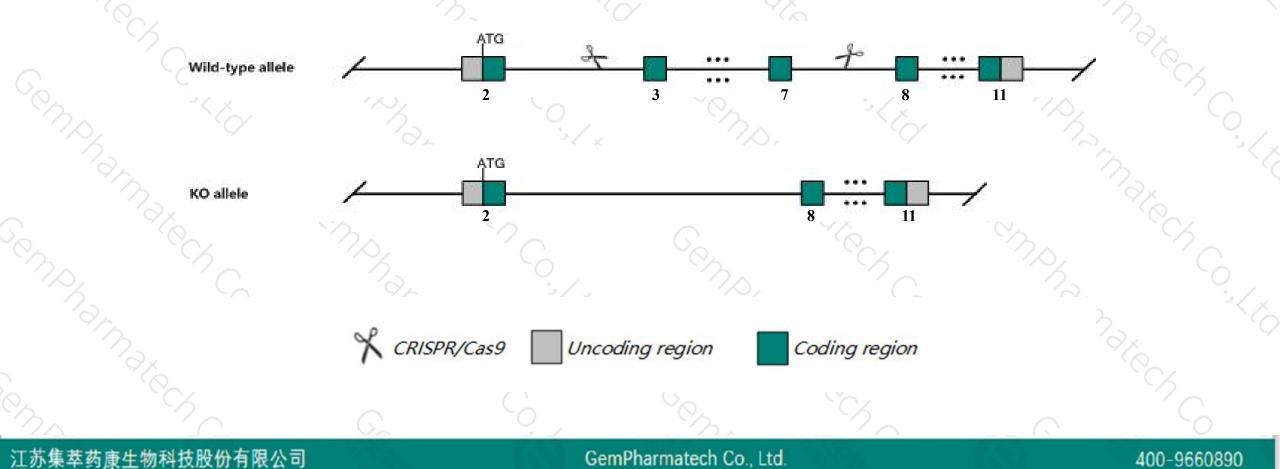




Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Prmt2* gene. The schematic diagram is as follows:





- The Prmt2 gene has 9 transcripts. According to the structure of Prmt2 gene, exon3-exon7 of Prmt2-203 (ENSMUST00000099572.9) transcript is recommended as the knockout region. The region contains 812bp coding sequence. Knock out the region will result in disruption of protein function.
- > In this project we use CRISPR/Cas9 technology to modify *Prmt2* gene. The brief process is as follows: CRISPR/Cas9 system

- According to the existing MGI data, Mice homozygous for a knock-out allele display a hyperplastic response to vascular injury while mutant mouse embryonic fibroblasts show an earlier S phase entry following release of serum starvation.
- The Prmt2 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.

Notice

Gene information (NCBI)



\$?

Prmt2 protein arginine N-methyltransferase 2 [Mus musculus (house mouse)]

Gene ID: 15468, updated on 31-Jan-2019

Summary

12122200 212 10 10				
Official Symbol	Prmt2 provided by MGI			
Official Full Name	protein arginine N-methyltransferase 2 provided by MGI			
Primary source	MGI:MGI:1316652			
See related	Ensembl:ENSMUSG0000020230			
Gene type	protein coding			
RefSeq status	VALIDATED			
Organism	Mus musculus			
Lineage Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Roden				
	Muroidea; Muridae; Murinae; Mus; Mus			
Also known as	AI504737, Hrmt1I1			
Expression	Broad expression in CNS E18 (RPKM 82.6), whole brain E14.5 (RPKM 68.0) and 23 other tissues See more			
Orthologs	human all			

江苏集萃药康生物科技股份有限公司

GemPharmatech Co., Ltd.

400-9660890

Transcript information (Ensembl)



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

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Prmt2-203	ENSMUST00000099572.9	2100	<u>475aa</u>	Protein coding	CCDS78836	Q3UID4	TSL:1 GENCODE basic APPRIS ALT
Prmt2-202	ENSMUST0000099571.9	2052	<u>445aa</u>	Protein coding	CCDS35942	Q3UKX1	TSL:1 GENCODE basic APPRIS P3
Prmt2-201	ENSMUST00000020452.11	2016	<u>445aa</u>	Protein coding	CCDS35942	Q3UKX1	TSL:1 GENCODE basic APPRIS P3
Prmt2-205	ENSMUST00000128099.7	2066	<u>254aa</u>	Nonsense mediated decay	80 <u>0</u> 8	<u>M0QW88</u>	TSL:1
rmt2-206	ENSMUST00000137857.7	1967	<u>254aa</u>	Nonsense mediated decay	1251	<u>M0QW88</u>	TSL:2
rmt2-208	ENSMUST00000217726.1	725	<u>127aa</u>	Nonsense mediated decay	(H)	A0A1W2P6Z3	TSL:3
rmt2-207	ENSMUST00000144670.7	769	No protein	Retained intron	83 2 0	-	TSL:2
rmt2-209	ENSMUST00000220116.1	751	No protein	Retained intron	1020	12	TSL:5
rmt2-204	ENSMUST00000128048.1	608	No protein	IncRNA	(2 7 1)	7	TSL:3
		1 1 2			S. M. etc.	5 V	

The strategy is based on the design of Prmt2-203 transcript, The transcription is shown below

< Prmt2-203 protein coding

Reverse strand

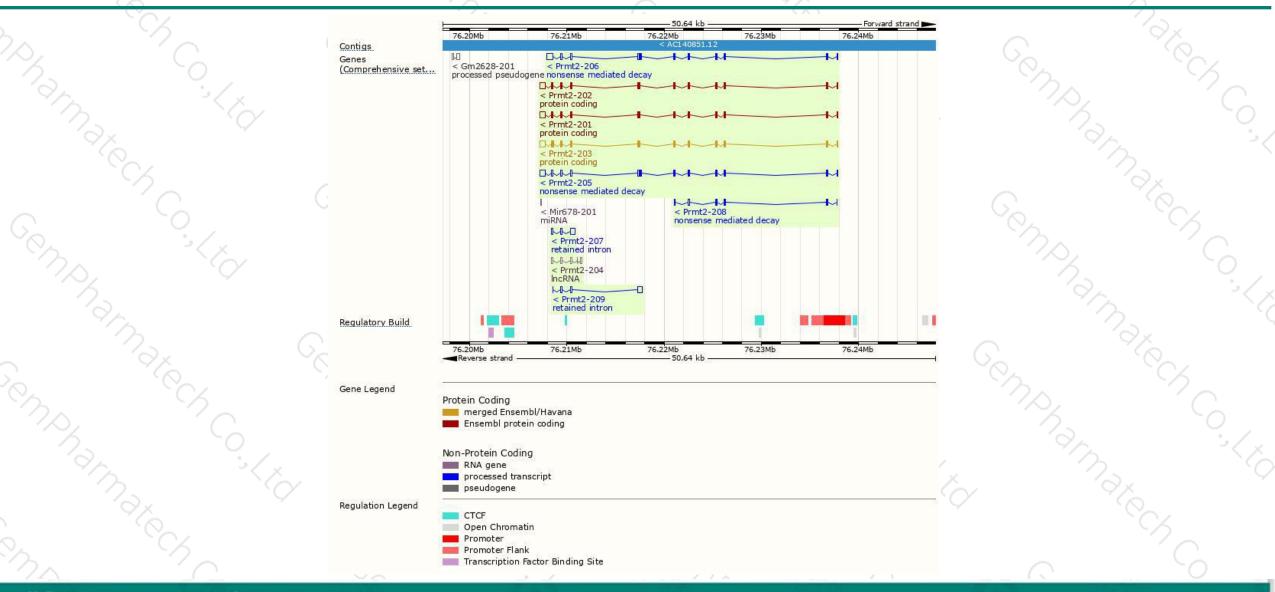
– 30.61 kb –

江苏集萃药康生物科技股份有限公司

GemPharmatech Co., Ltd.

400-9660890

Genomic location distribution



集萃药康 GemPharmatech

400-9660890

江苏集萃药康生物科技股份有限公司

GemPharmatech Co., Ltd.

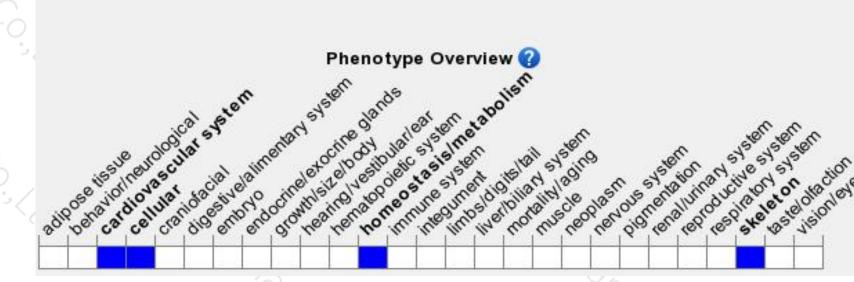
Protein domain



			, h
DD5	ENSMUSP00000097 Low complexity (Seg) Superfamily	S-adenosyl-L-methionine-dependent methyltransferase	ate ch
arnar.	SMART Pfam	SH3-like domain superfamily SH3 domain SH3 domain Methyltransferase small domain	
G	PROSITE profiles	SH3 domain Protein arginine N-methyltransferase	× CCC
C M D L	PANTHER	PTHR11006:SF92	6
~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~	Gene3D	PTHR11006 2,30.30.40 3.40.50.150	
	CDD	cd11806	
mohs	All sequence SNPs/i	Sequence variants (dbSNP and all other sources)	, ^C
	Variant Legend	stop gained missense variant splice region variant synonymous variant	
Sha .	Scale bar	<b>0</b> 40 80 120 160 200 240 280 320 360 400 475	
江苏集萃药康生	_{上物科技股份有限公司}	GemPharmatech Co., Ltd. 4	00-9660890

### 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 a knock-out allele display a hyperplastic response to vascular injury while mutant mouse embryonic fibroblasts show an earlier S phase entry following release of serum starvation.



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



