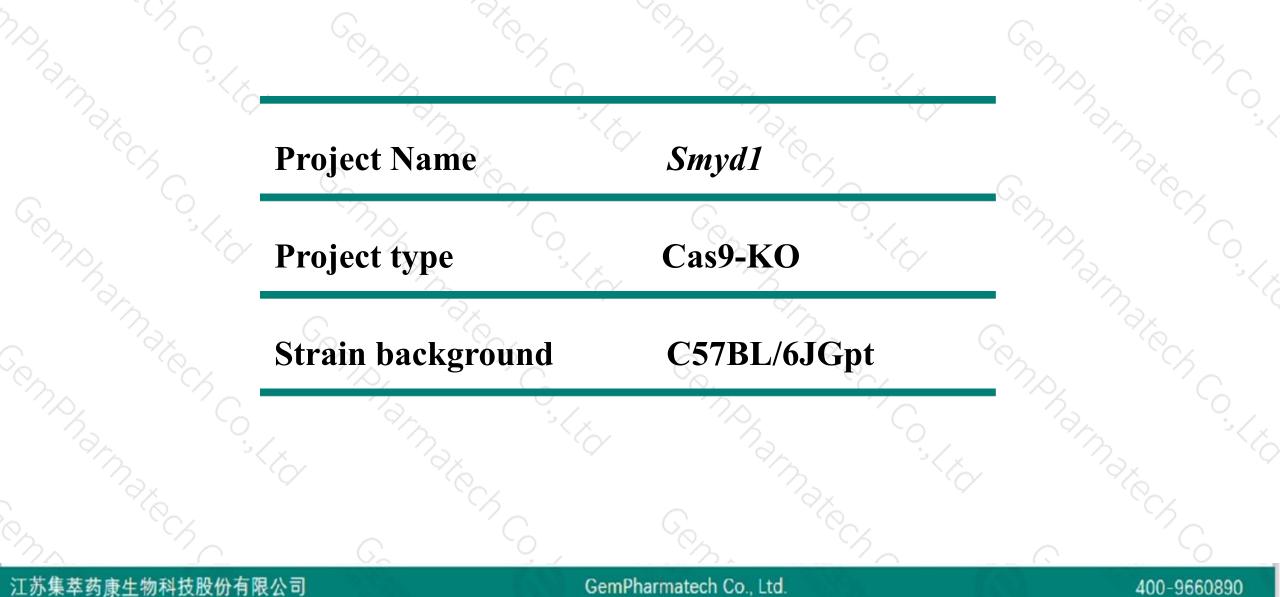


Smyd1 Cas9-KO Strategy

Designer: Design Date: Jinling Wang 2019-9-20

Project Overview

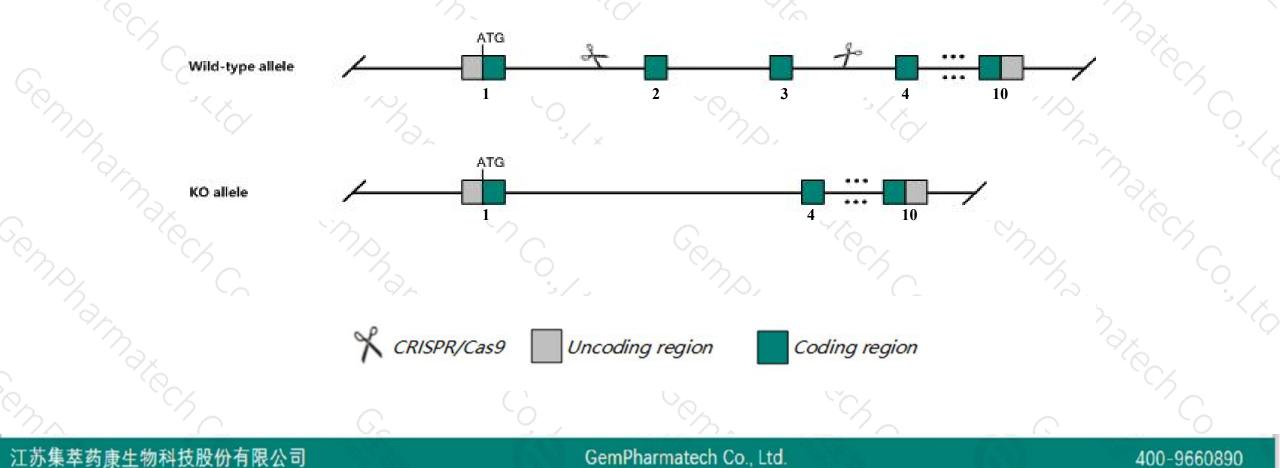




Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Smyd1 gene. The schematic diagram is as follows:





- The Smyd1 gene has 6 transcripts. According to the structure of Smyd1 gene, exon2-exon3 of Smyd1-201 (ENSMUST00000074301.9) transcript is recommended as the knockout region. The region contains 391bp coding sequence. Knock out the region will result in disruption of protein function.
- > In this project we use CRISPR/Cas9 technology to modify Smyd1 gene. The brief process is as follows: CRISPR/Cas9 system



- According to the existing MGI data, Homozygous mutation of this gene results in embryonic lethality at E10.5. Mutant embryos exhibit an enlarged heart and developmental abnormalities of the right ventricle.
- The Smyd1 gene is located on the Chr6. 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)



\$?

Smyd1 SET and MYND domain containing 1 [Mus musculus (house mouse)]

Gene ID: 12180, updated on 9-Apr-2019

Summary

Official Symbol	Smyd1 provided by MGI
Official Full Name	SET and MYND domain containing 1 provided by MGI
Primary source	MGI:MGI:104790
See related	Ensembl:ENSMUSG00000055027
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	4632404M21Rik, Bop, C78565, Zmynd18
Expression	Biased expression in heart adult (RPKM 89.8), mammary gland adult (RPKM 9.3) and 1 other tissueSee more
Orthologs	human all

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Transcript information (Ensembl)



The gene has 6 transcripts, all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags	
Smyd1-201	ENSMUST0000074301.9	3361	<u>490aa</u>	Protein coding	CCDS51805	P97443	TSL:1 GENCODE basic APPRIS ALT1	
Smyd1-202	ENSMUST00000114186.8	3292	<u>477aa</u>	Protein coding	CCDS20227	P97443 Q3UQT9	TSL:1 GENCODE basic APPRIS P3	
Smyd1-203	ENSMUST00000114188.2	2064	<u>456aa</u>	Protein coding	040	<u>G5E8R7</u>	TSL:5 GENCODE basic	
Smyd1-205	ENSMUST00000173297.1	947	<u>21aa</u>	Protein coding	8 <u>4</u> 8	G3UXZ0	TSL:5 GENCODE basic	
Smyd1-206	ENSMUST00000173730.1	428	<u>58aa</u>	Protein coding	151	<u>G3UZ94</u>	CDS 5' incomplete TSL:5	
Smyd1-204	ENSMUST00000129630.2	376	<u>69aa</u>	Protein coding	240	Q8BMV4	TSL:1 GENCODE basic	

The strategy is based on the design of Smyd1-201 transcript, The transcription is shown below

< Smyd1-201 protein coding

Reverse strand

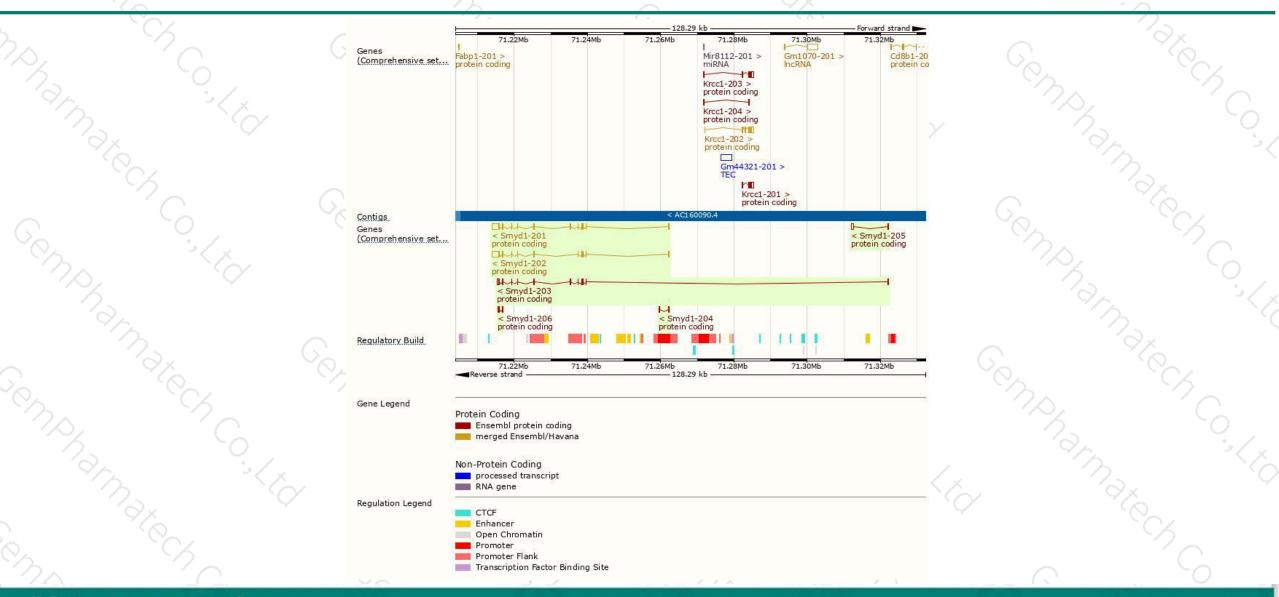
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48.34 kb

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Genomic location distribution



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Protein domain



ENSMUSP00000073							
SIFTS import Superfamily	SSF82199						
SMART	SET domain						
Pfam.	SET domain						
	Zinc finger, MYND-t	уре					
PROSITE profiles	SET domain						
PROSITE patterns	Zinc finger, MYND-t	he na -					6
PANTHER	Zinc finger, MYND-t	уре					
	FTHR12197:SF184						
Gene3D	2.170.270.10			1.25.40	970	Tetratricopeptic	de-like helical do
	3.30.60.180 3.30.7	0.3410					2
All sequence SNPs/i	Sequence variants (dbSNP an	d all other sources)	0 0	1	11	0.000	9
Variant Legend							
	missense variant synonymous variant						
Scale bar		20 180	240	300	360	420	490
\sim	G.	0	Sh.	-7-		0	0

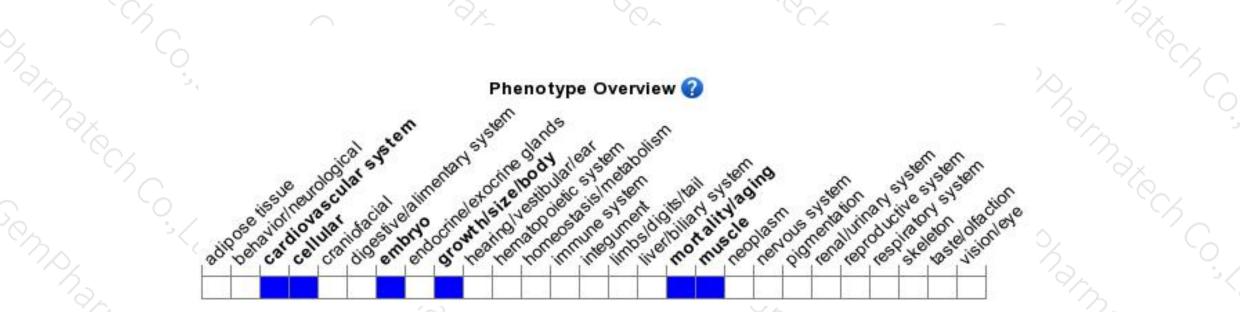
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Mouse phenotype description(MGI)





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, Homozygous mutation of this gene results in embryonic lethality at E10.5. Mutant embryos exhibit an enlarged heart and developmental abnormalities of the right ventricle.



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



