

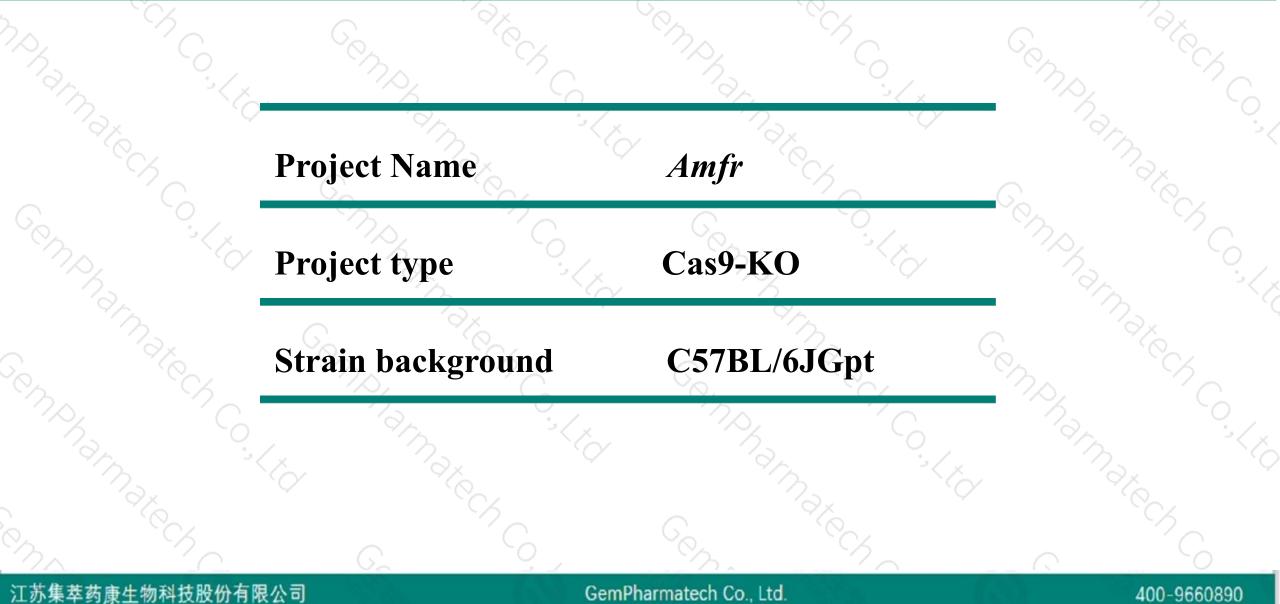
Amfr Cas9-KO Strategy

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Project Overview

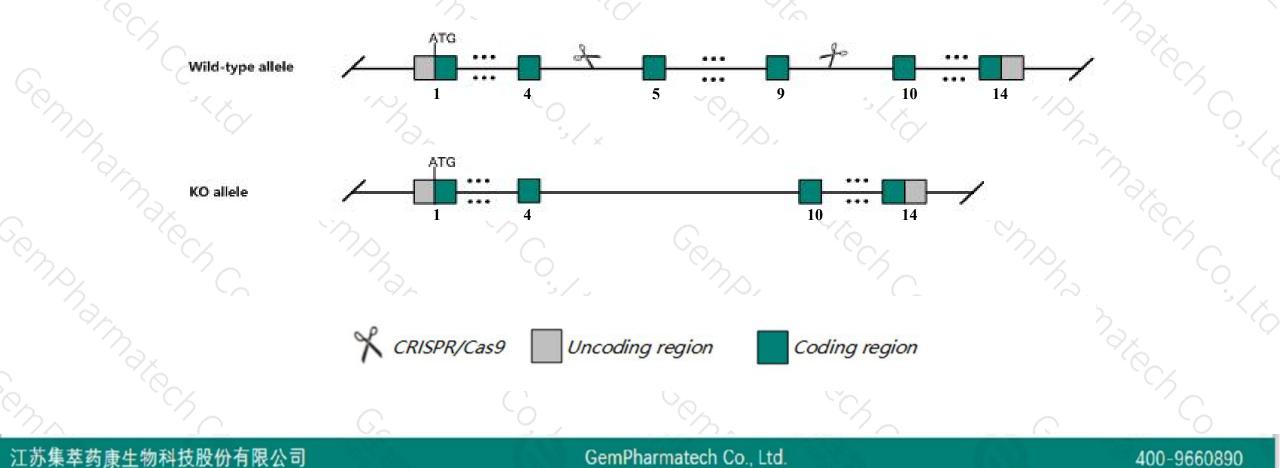




Knockout strategy



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





- The Amfr gene has 4 transcripts. According to the structure of Amfr gene, exon5-exon9 of Amfr-201 (ENSMUST00000053766.13) transcript is recommended as the knockout region. The region contains 610bp coding sequence. Knock out the region will result in disruption of protein function.
- > In this project we use CRISPR/Cas9 technology to modify Amfr gene. The brief process is as follows: CRISPR/Cas9 system v

- According to the existing MGI data, Mice for a gene-trapped null allele are obese and develop liver steatosis and/or hepatic inflammation resembling nonalcoholic steatohepatitis. Some mice develop liver tumors. Mice homozygous for another knock-out allele exhibit normal HMGCR turnover in mouse embryonic fibroblasts.
- The Amfr gene is located on the Chr8. 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)



Amfr autocrine motility factor receptor [Mus musculus (house mouse)]

Gene ID: 23802, updated on 10-Oct-2019

Summary

Official Symbol	Amfr provided by MGI	2
Official Full Name	autocrine motility factor receptor provided by MGI	
Primary source	MGI:MGI:1345634	
See related	Ensembl:ENSMUSG00000031751	
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	gp78	
Expression	Ubiquitous expression in adrenal adult (RPKM 95.7), liver adult (RPKM 63.4) and 28 other tissues See more	
Orthologs	human all	
UT X		

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Transcript information (Ensembl)



The gene has 4 transcripts, all transcripts are shown below:

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Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
ENSMUST0000053766.13	3880	<u>639aa</u>	Protein coding	CCDS22533	Q3TCI2 Q9R049	TSL:1 GENCODE basic APPRIS P1
ENSMUST00000143265.1	414	<u>105aa</u>	Protein coding	5 .)	H3BJC0	CDS 3' incomplete TSL:5
ENSMUST00000139702.1	779	No protein	Retained intron	8 2 0	-	TSL:2
ENSMUST00000137475.1	392	No protein	IncRNA	120	-	TSL:5
	ENSMUST0000053766.13 ENSMUST00000143265.1 ENSMUST00000139702.1	ENSMUST0000053766.13 3880 ENSMUST00000143265.1 414 ENSMUST00000139702.1 779	ENSMUST0000053766.13 3880 639aa ENSMUST0000143265.1 414 105aa ENSMUST0000139702.1 779 No protein	ENSMUST0000053766.133880639aaProtein codingENSMUST00000143265.1414105aaProtein codingENSMUST00000139702.1779No proteinRetained intron	ENSMUST0000053766.133880639aaProtein codingCCDS22533ENSMUST0000143265.1414105aaProtein coding-ENSMUST0000139702.1779No proteinRetained intron-	ENSMUST0000053766.133880639aaProtein codingCCDS22533Q3TCl2 Q9R049ENSMUST0000143265.1414105aaProtein coding-H3BJC0ENSMUST0000139702.1779No proteinRetained intron

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

< Amfr-201 protein coding

Reverse strand

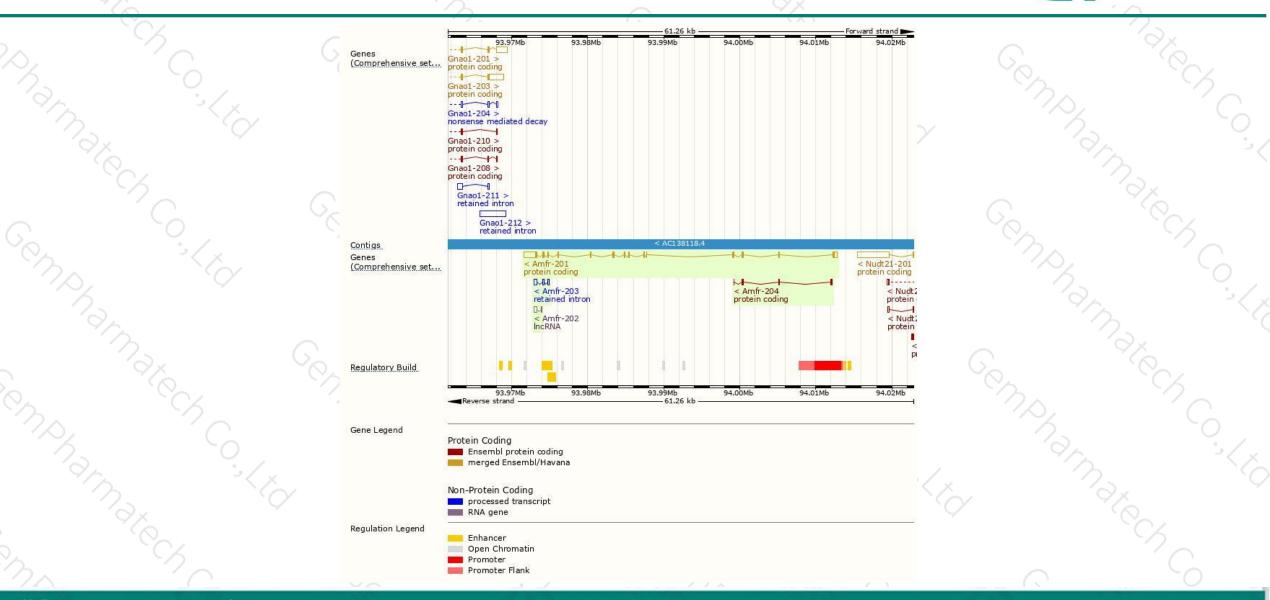
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41.26 kb

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Genomic location distribution



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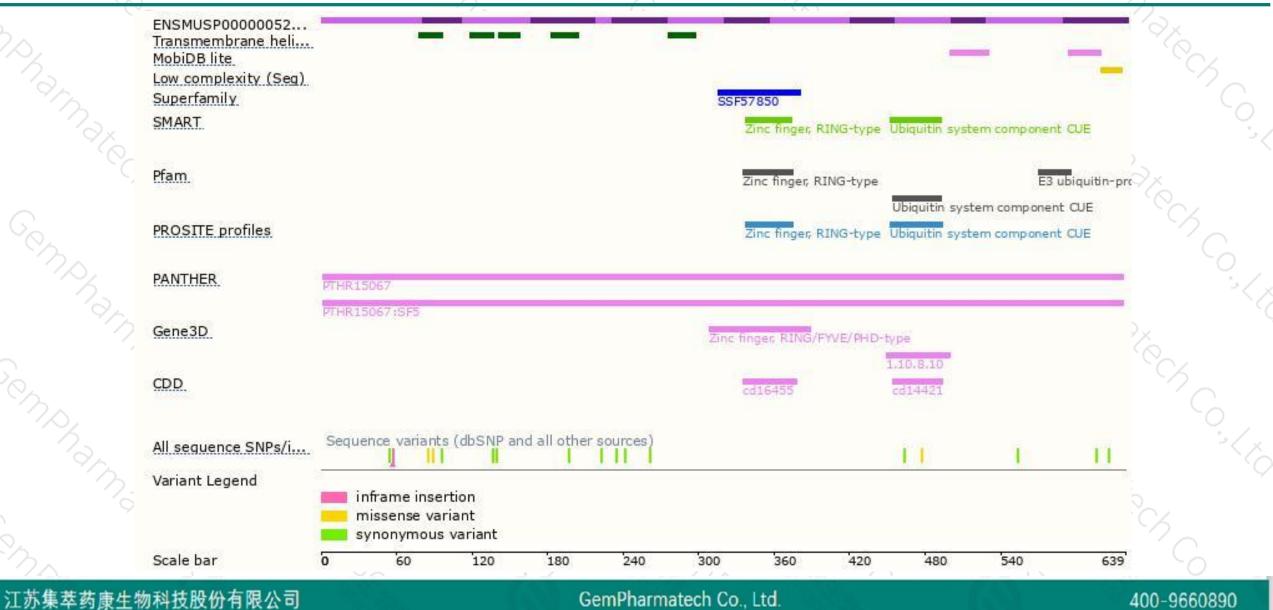
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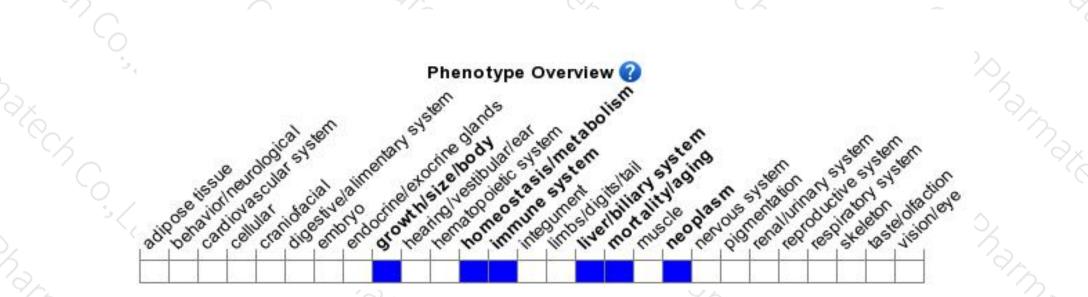
Protein domain





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 for a gene-trapped null allele are obese and develop liver steatosis and/or hepatic inflammation resembling nonalcoholic steatohepatitis. Some mice develop liver tumors. Mice homozygous for another knock-out allele exhibit normal HMGCR turnover in mouse embryonic fibroblasts.

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



