F2r Cas9-KO Strategy Romphamater Contraction

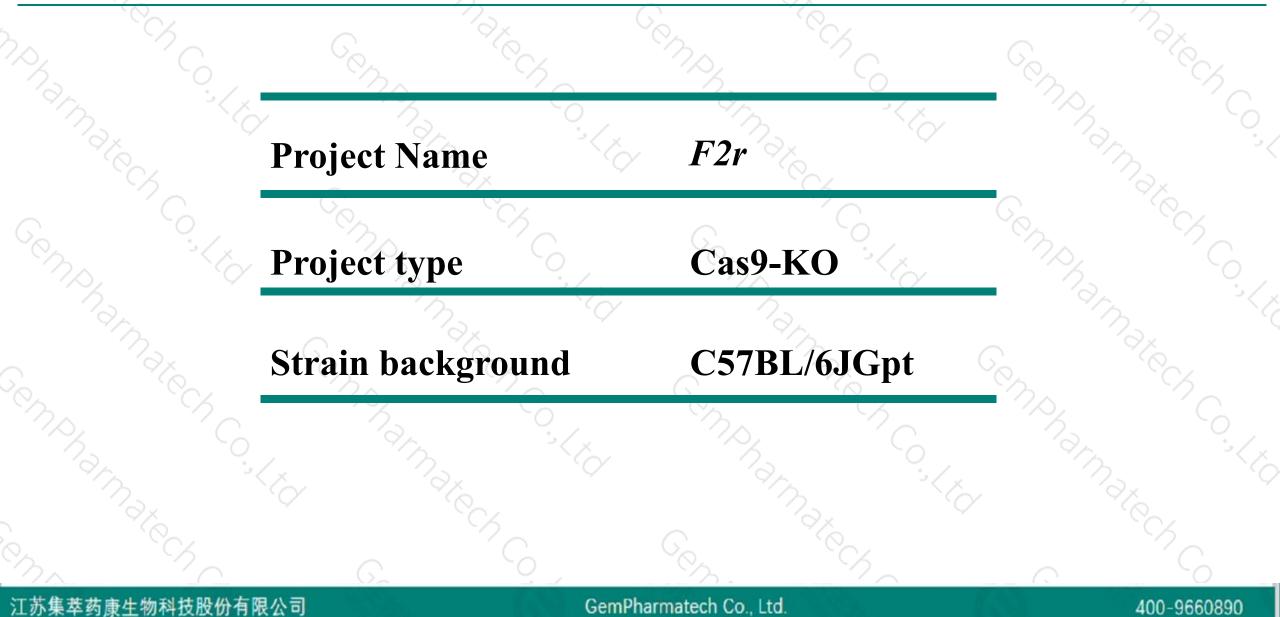
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Cemphamaten Co. Cenphamaten Co. Designer:Yun Li Reviewer:Longyun Hu Design Date: 2019-12-24

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Project Overview



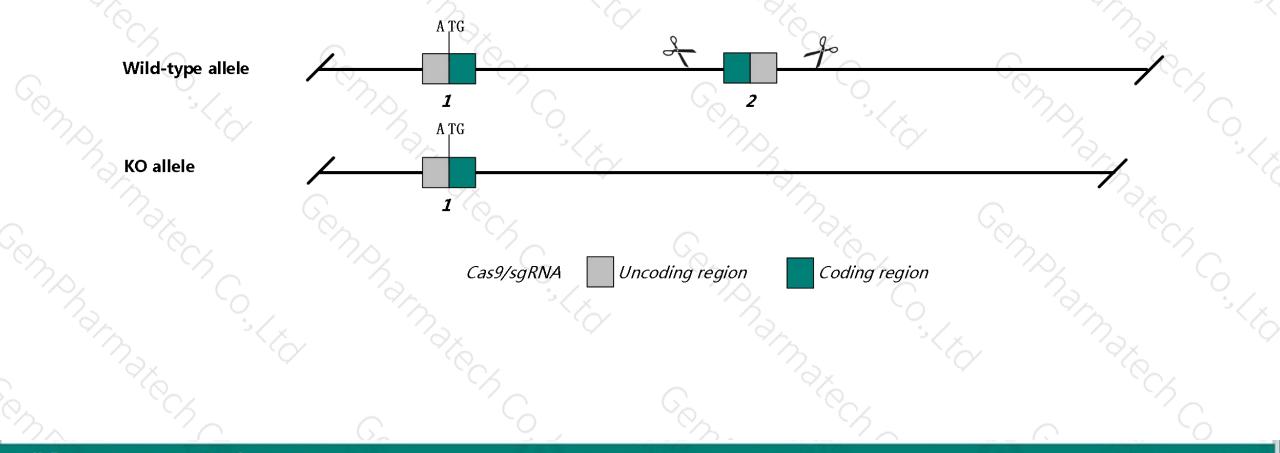


Knockout strategy



400-9660890

This model will use CRISPR/Cas9 technology to edit the F2r gene. The schematic diagram is as follows:



Technical routes



The *F2r* gene has 1 transcript.According to the structure of *F2r* gene, exon2 of *F2r*-201 (ENSMUST00000059193.6)transcript is recommended as the knockout region.The region contains 1205bp coding sequence.Knock out the region will result in disruption of protein function.

In this project we use CRISPR/Cas9 technology to modify *F2r* gene. The brief process is as follows: gRNA was transcribed in vitro.Cas9 and gRNA 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, Targeted mutations of this locus result in increased midgestational lethality, with up to ~50% of mutants surviving to adulthood. Gene deficiency does not affect thrombin signaling in mouse platelets but markedly attenuates thrombin signaling in mouse microvascular endothelial cells.
- > The KO region deletes most of the coding sequence and result in frameshift.
- The F2r gene is located on the Chr13. 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)

F2r coagulation factor II (thrombin) receptor [Mus musculus (house mouse)]

Gene ID: 14062, updated on 6-Nov-2018

Summary

Official Symbol F2r provided by MGI Official Full Name coagulation factor II (thrombin) receptor provided by MGI Primary source MGI:MGI:101802 See related Ensembl:ENSMUSG0000048376 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 Cf2r; Par1; ThrR; Al482343 Expression Broad expression in lung adult (RPKM 53.2), adrenal adult (RPKM 42.2) and 24 other tissues See more human all Orthologs

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Transcript information (Ensembl)



The gene has 1 transcripts, and the transcript is shown below:

| Show/ | hide columns (1 hidden) | | | | | | | Filter |
|---------|-------------------------|------|--------------|----------------|------------|-----------|----------------------------------|-------------------------------|
| Name 👙 | Transcript ID 👙 | bp 🖕 | Protein 👙 | Biotype 👌 | CCDS 👌 | UniProt 🛊 | RefSeq 🖕 | Flags |
| F2r-201 | ENSMUST0000059193.6 | 3336 | <u>430aa</u> | Protein coding | CCDS26701@ | P30558@ | <u>NM 010169</u> 교 NP 034299교 | TSL:1 GENCODE basic APPRIS P1 |

The strategy is based on the design of F2r-201 transcript, The transcription is shown below

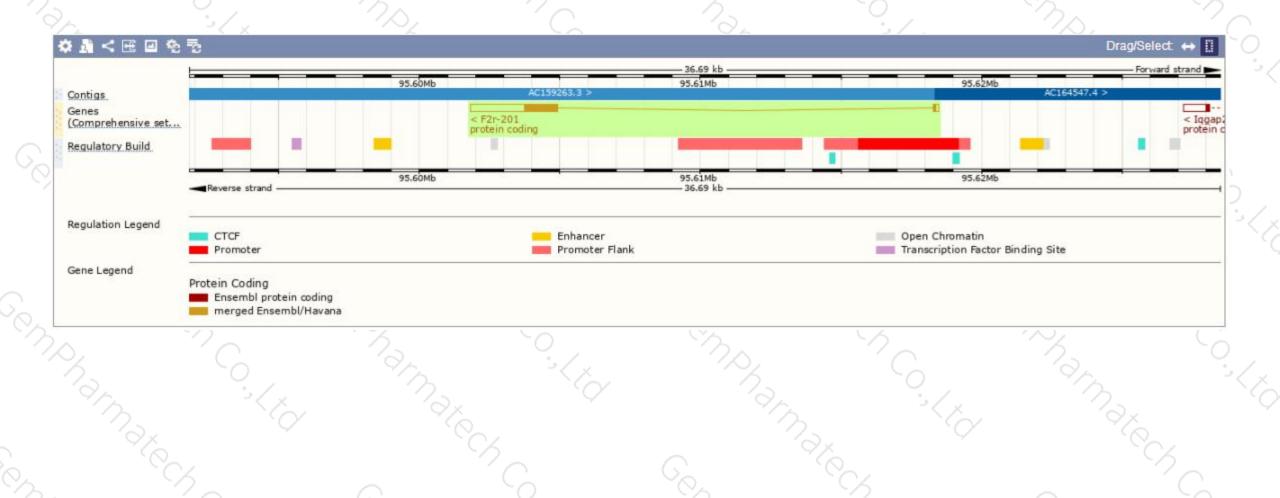
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Genomic location (Ensembl)





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Protein domain (Ensembl)

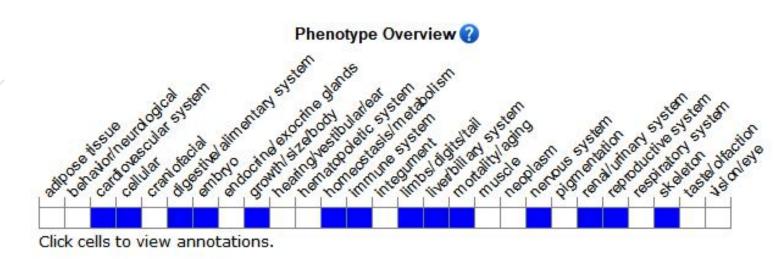


| Protease-activated receptor Pfam. domain. PROSITE profiles PROSITE patterns PROSITE patterns Gene3D. All sequence SNPs/L Sequence variants (dbSNP and all other sources) | |
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Mouse phenotype description(MGI)



Phenotypes affected by the gene are marked in blue. Data quoted from MGI database(http://www.informatics.jax.org/).

Targeted mutations of this locus result in increased midgestational lethality, with up to \sim 50% of mutants surviving to adulthood. Gene deficiency does not affect thrombin signaling in mouse platelets but markedly attenuates thrombin signaling in mouse microvascular endothelial cells.

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



