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Project Overview



Project Name Cfh

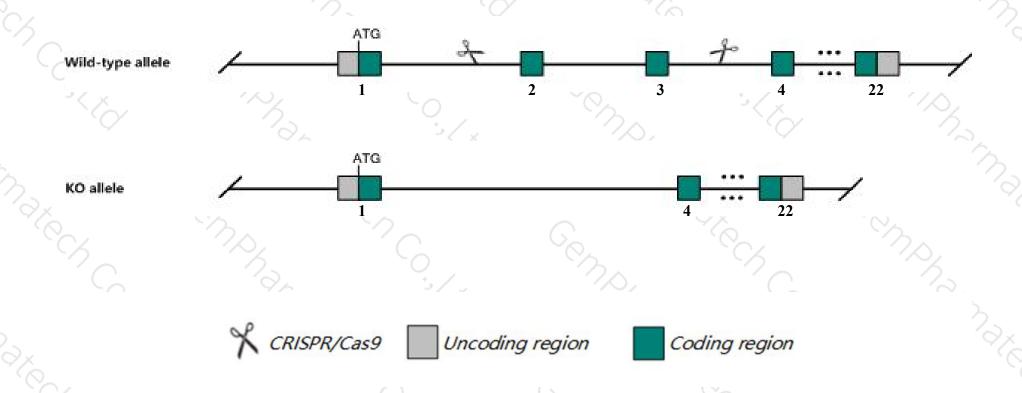
Project type Cas9-KO

Strain background C57BL/6JGpt

Knockout strategy



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



Technical routes



- ➤ The *Cfh* gene has 8 transcripts. According to the structure of *Cfh* gene, exon2-exon3 of *Cfh-202*(ENSMUST00000111976.8) transcript is recommended as the knockout region. The region contains 292bp coding sequence.

 Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Cfh* gene. The brief process is as follows: CRISPR/Cas9 system w

Notice



- ➤ According to the existing MGI data, Homozygous mutation of this gene results in markedly reduced serum C3, abnormal renal histology, spontaneous membranoproliferative glomerulonephritis (MPGN), hematuria, proteinuria, and increased mortality at 8 months of age.
- > The *Cfh* gene is located on the Chr1. 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)



Cfh complement component factor h [Mus musculus (house mouse)]

Gene ID: 12628, updated on 19-Mar-2019

Summary

☆ ?

Official Symbol Cfh provided by MGI

Official Full Name complement component factor h provided by MGI

Primary source MGI:MGI:88385

See related Ensembl:ENSMUSG00000026365

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 Mud-1, NOM, Sas-1, Sas1

Expression Biased expression in liver E18 (RPKM 75.9), liver adult (RPKM 32.7) and 10 other tissuesSee more

Orthologs <u>human</u> all

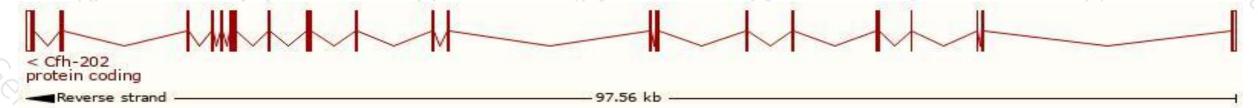
Transcript information (Ensembl)



The gene has 8 transcripts, all transcripts are shown below:

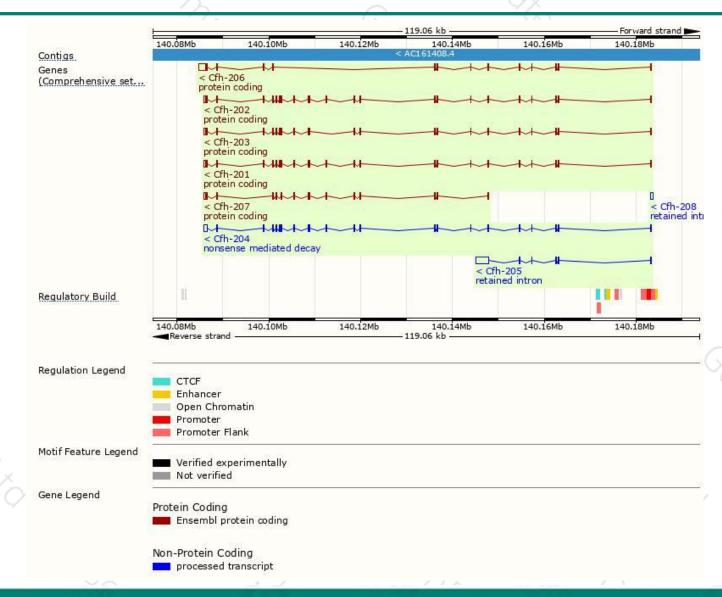
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Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
ENSMUST00000111976.8	4365	1252aa	Protein coding	CCDS48388	E9Q810	TSL:1 GENCODE basic APPRIS P2
ENSMUST00000066859.12	4361	<u>1234aa</u>	Protein coding	677	P06909	TSL:1 GENCODE basic APPRIS ALT2
ENSMUST00000111977.7	4100	<u>1195aa</u>	Protein coding	140	E9Q8H9	TSL:1 GENCODE basic APPRIS ALT2
ENSMUST00000192880.5	3746	707aa	Protein coding	100	A0A0A6YVP8	TSL:5 GENCODE basic
ENSMUST00000192919.5	2976	<u>837aa</u>	Protein coding	1753	A0A0A6YWP4	CDS 5' incomplete TSL:5
ENSMUST00000123238.1	4300	<u>1110aa</u>	Nonsense mediated decay	684	D6RGQ0	TSL:1
ENSMUST00000148225.1	3415	No protein	Retained intron	940	-	TSL:1
ENSMUST00000194688.1	631	No protein	Retained intron	100	2	TSL:NA
	ENSMUST00000111976.8 ENSMUST00000066859.12 ENSMUST00000111977.7 ENSMUST00000192880.5 ENSMUST00000192919.5 ENSMUST00000123238.1 ENSMUST00000148225.1	ENSMUST00000111976.8 4365 ENSMUST00000066859.12 4361 ENSMUST00000111977.7 4100 ENSMUST00000192880.5 3746 ENSMUST00000192919.5 2976 ENSMUST00000123238.1 4300 ENSMUST00000148225.1 3415	ENSMUST00000111976.8 4365 1252aa ENSMUST00000066859.12 4361 1234aa ENSMUST00000111977.7 4100 1195aa ENSMUST00000192880.5 3746 707aa ENSMUST00000192919.5 2976 837aa ENSMUST00000123238.1 4300 1110aa ENSMUST00000148225.1 3415 No protein	ENSMUST00000111976.8 4365 1252aa Protein coding ENSMUST00000066859.12 4361 1234aa Protein coding ENSMUST00000111977.7 4100 1195aa Protein coding ENSMUST00000192880.5 3746 707aa Protein coding ENSMUST00000192919.5 2976 837aa Protein coding ENSMUST00000123238.1 4300 1110aa Nonsense mediated decay ENSMUST00000148225.1 3415 No protein Retained intron	ENSMUST00000111976.8 4365 1252aa Protein coding CCDS48388 ENSMUST00000066859.12 4361 1234aa Protein coding - ENSMUST00000111977.7 4100 1195aa Protein coding - ENSMUST00000192880.5 3746 707aa Protein coding - ENSMUST00000192919.5 2976 837aa Protein coding - ENSMUST00000123238.1 4300 1110aa Nonsense mediated decay - ENSMUST00000148225.1 3415 No protein Retained intron -	ENSMUST00000111976.8 4365 1252aa Protein coding CCDS48388 E9Q8I0 ENSMUST00000066859.12 4361 1234aa Protein coding - P06909 ENSMUST00000111977.7 4100 1195aa Protein coding - E9Q8H9 ENSMUST00000192880.5 3746 707aa Protein coding - A0A0A6YVP8 ENSMUST00000192919.5 2976 837aa Protein coding - A0A0A6YWP4 ENSMUST00000123238.1 4300 1110aa Nonsense mediated decay - D6RGQ0 ENSMUST00000148225.1 3415 No protein Retained intron - -

The strategy is based on the design of Cfh-202 transcript, The transcription is shown below



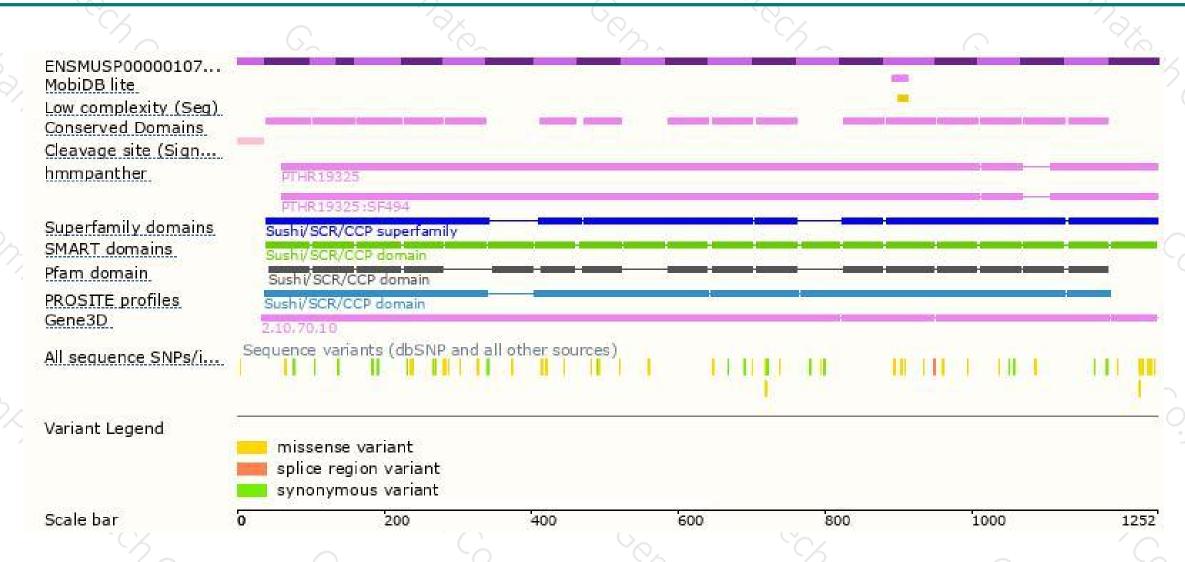
Genomic location distribution





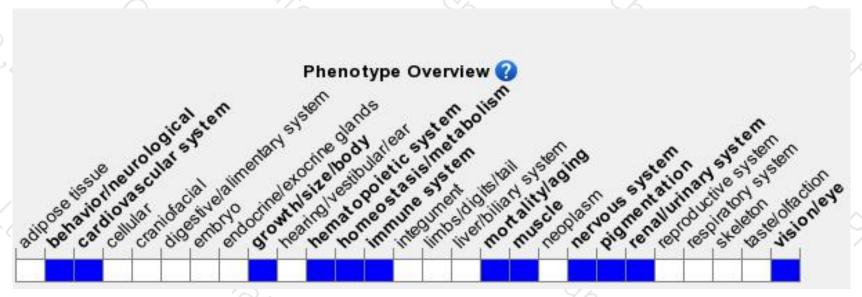
Protein domain





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, Homozygous mutation of this gene results in markedly reduced serum C3, abnormal renal histology, spontaneous membranoproliferative glomerulonephritis (MPGN), hematuria, proteinuria, and increased mortality at 8 months of age.



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





