

Dcaf17 Cas9-KO Strategy

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Design Date: 2020-9-30

Project Overview



Project Name Dcaf17

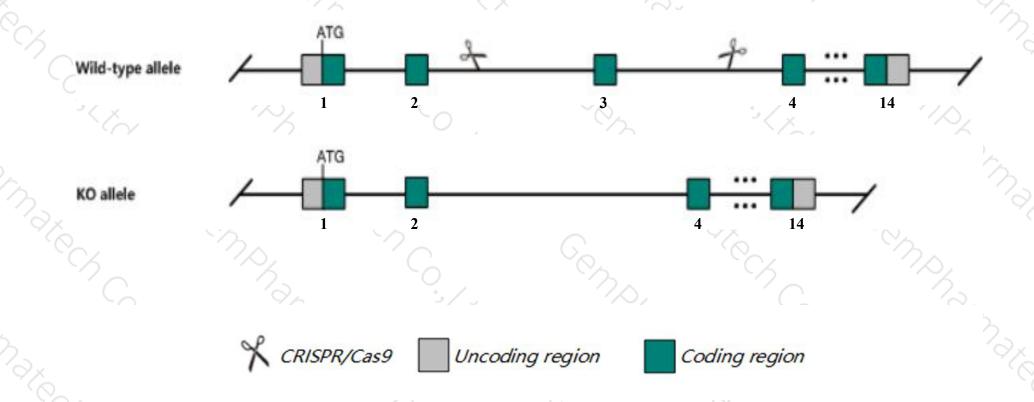
Project type Cas9-KO

Strain background C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Dcaf17* gene. The schematic diagram is as follows:



Technical routes



- > The Dcaf17 gene has 9 transcripts. According to the structure of Dcaf17 gene, exon3 of Dcaf17-209(ENSMUST00000154704.7) transcript is recommended as the knockout region. The region contains 91bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Dcaf17* gene. The brief process is as follows: CRISPR/Cas9 system 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.

Notice



- > According to the existing MGI data, mice homozygous for a knock-out allele exhibit male infertility associated with defects in spermatogenesis.
- > The KO region is close to *Mettl8* gene.Knockout the region may affect the function of *Mettl8* gene.
- > The *Dcaf17* gene is located on the Chr2. 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)



Dcaf17 DDB1 and CUL4 associated factor 17 [Mus musculus (house mouse)]

Gene ID: 75763, updated on 25-Sep-2020

Summary



Official Symbol Dcaf17 provided by MGI

Official Full Name DDB1 and CUL4 associated factor 17 provided by MGI

Primary source MGI:MGI:1923013

See related Ensembl: ENSMUSG00000041966

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 Al448937; 2810055O12Rik; 4833418A01Rik; A030004A10Rik; A930009G19Rik

Expression Ubiquitous expression in CNS E18 (RPKM 1.5), CNS E14 (RPKM 1.4) and 28 other tissues See more

Orthologs human all

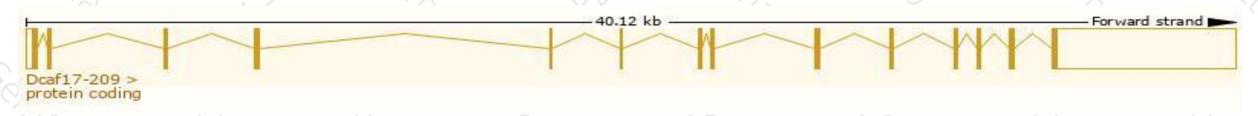
Transcript information (Ensembl)



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

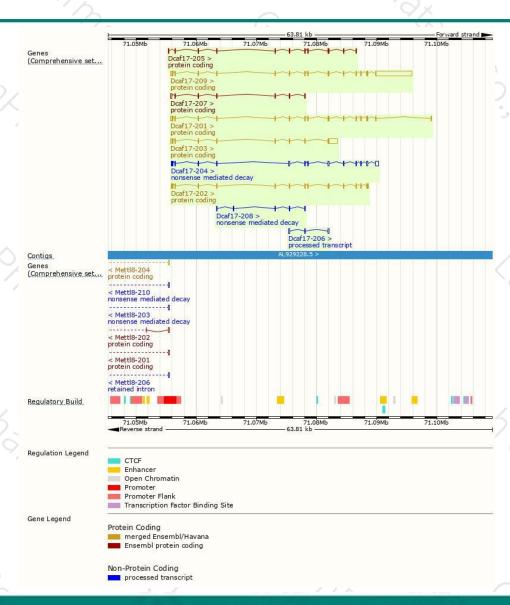
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Dcaf17-209	ENSMUST00000154704.7	7739	519aa	Protein coding	CCDS50601	Q3TUL7	TSL:1 GENCODE basic APPRIS P1
Dcaf17-203	ENSMUST00000112159.8	2490	286aa	Protein coding	CCDS50602	Q3TUL7	TSL:1 GENCODE basic
Dcaf17-201	ENSMUST00000064141.11	1905	519aa	Protein coding	CCDS50601	Q3TUL7	TSL:1 GENCODE basic APPRIS P
Dcaf17-202	ENSMUST00000102701.4	1703	505aa	Protein coding	CCDS16111	Z4YK64	TSL:1 GENCODE basic
Dcaf17-205	ENSMUST00000130292.7	1214	<u>405aa</u>	Protein coding	-	F6YAD1	CDS 5' and 3' incomplete TSL:1
Dcaf17-207	ENSMUST00000135357.7	765	<u>157aa</u>	Protein coding	678	B0R0U2	CDS 3' incomplete TSL:3
Dcaf17-204	ENSMUST00000112167.8	2061	<u>156aa</u>	Nonsense mediated decay	5 - 2	F8WIJ2	TSL:5
Dcaf17-208	ENSMUST00000136299.1	441	<u>58aa</u>	Nonsense mediated decay	122	<u>F6XZ79</u>	CDS 5' incomplete TSL:3
Dcaf17-206	ENSMUST00000132619.1	443	No protein	Processed transcript	Asta		TSL:2
		///				1 V.	

The strategy is based on the design of *Dcaf17-209* transcript, the transcription is shown below:



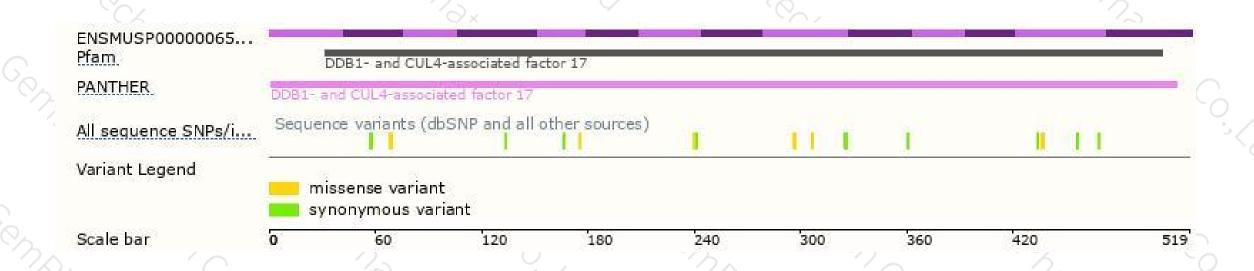
Genomic location distribution





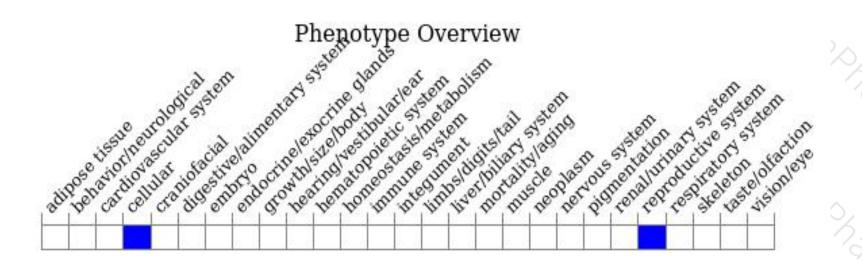
Protein domain





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, mice homozygous for a knock-out allele exhibit male infertility associated with defects in spermatogenesis.



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





