

# Dcaf17 Cas9-CKO Strategy

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## **Project Overview**



Project Name Dcaf17

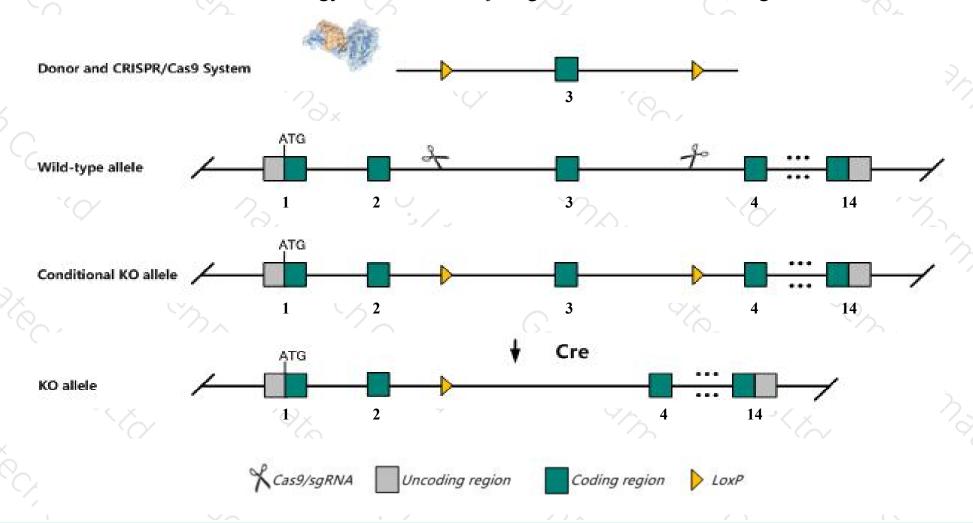
Project type Cas9-CKO

Strain background C57BL/6JGpt

## Conditional 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:sgRNA was transcribed in vitro, donor vector was constructed.Cas9, sgRNA and Donor 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.
- > The flox mice was knocked out after mating with mice expressing Cre recombinase, resulting in the loss of function of the target gene in specific tissues and cell types.

### **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 loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at existing technological 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

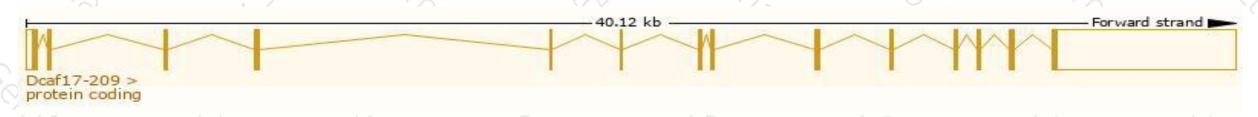
## Transcript information (Ensembl)



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

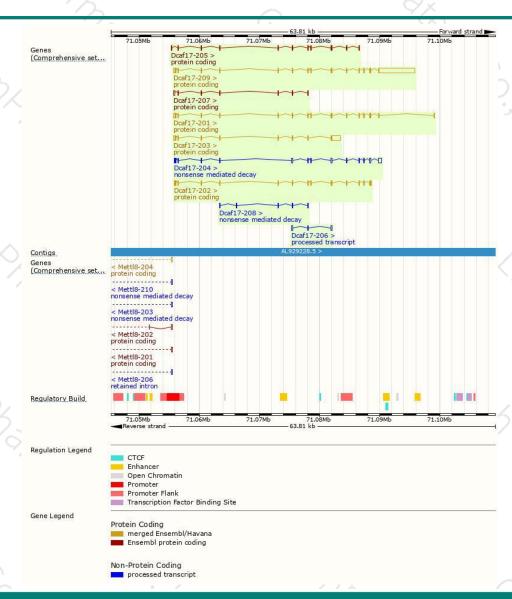
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Dcaf17-209	ENSMUST00000154704.7	7739	519aa	Protein coding	CCDS50601	Q3TUL7	TSL:1 GENCODE basic APPRIS P
Dcaf17-203	ENSMUST00000112159.8	2490	286aa	Protein coding	CCDS50602	Q3TUL7	TSL:1 GENCODE basic
Dcaf17-201	ENSMUST00000064141.11	1905	<u>519aa</u>	Protein coding	CCDS50601	Q3TUL7	TSL:1 GENCODE basic APPRIS P
Dcaf17-202	ENSMUST00000102701.4	1703	<u>505aa</u>	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
Ocaf17-207	ENSMUST00000135357.7	765	<u>157aa</u>	Protein coding	678	BOROU2	CDS 3' incomplete TSL:3
Dcaf17-204	ENSMUST00000112167.8	2061	<u>156aa</u>	Nonsense mediated decay		F8WIJ2	TSL:5
Ocaf17-208	ENSMUST00000136299.1	441	<u>58aa</u>	Nonsense mediated decay	12	F6XZ79	CDS 5' incomplete TSL:3
Ocaf17-206	ENSMUST00000132619.1	443	No protein	Processed transcript	878		TSL:2

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



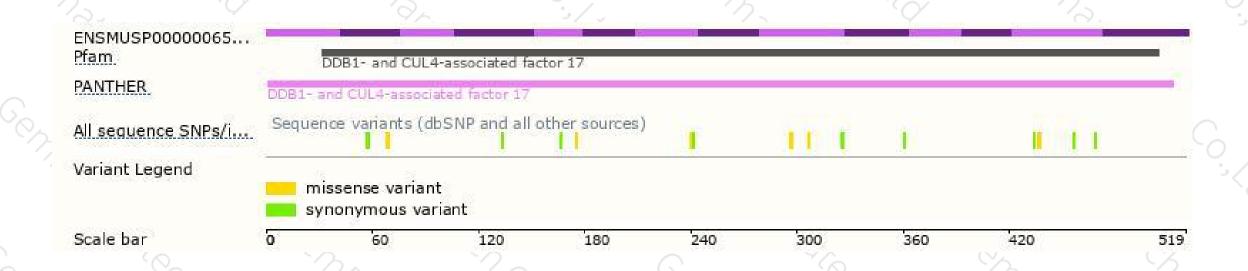
### Genomic location distribution





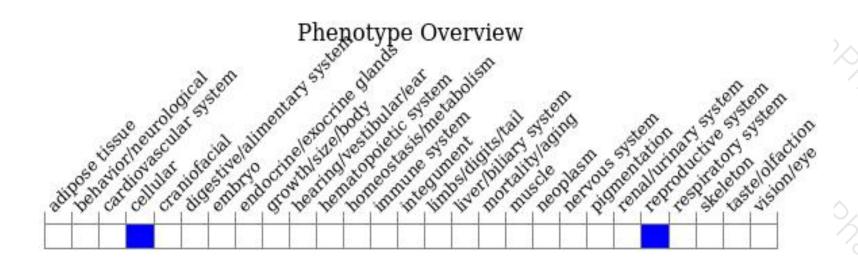
## Protein domain





## 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.

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