

Ift140 Cas9-CKO Strategy

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Project Overview



Project Name

Ift140

Project type

Cas9-CKO

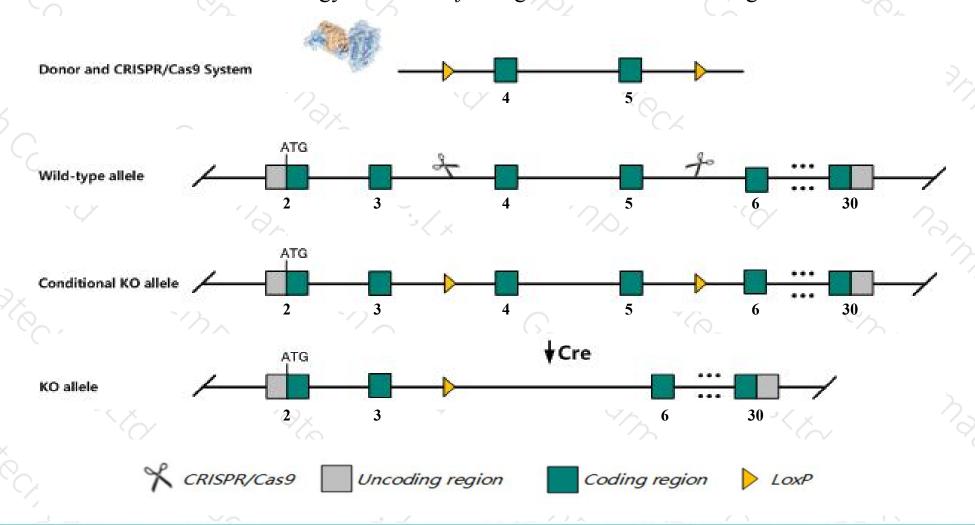
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Ift140* gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Ift140* gene has 9 transcripts. According to the structure of *Ift140* gene, exon4-exon5 of *Ift140-201*(ENSMUST00000024983.11) transcript is recommended as the knockout region. The region contains 265bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Ift140* gene. The brief process is as follows:CRISPR/Cas9 system 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 will be 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 reporter knock-out allele die at mid-gestation.

 Mice homozygous for an ENU-induced mutation exhibit cardiovascular defects and situs abnormalities.
- The *Ift140* gene is located on the Chr17. 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)



Ift140 intraflagellar transport 140 [Mus musculus (house mouse)]

Gene ID: 106633, updated on 9-Apr-2019

Summary

☆ ?

Official Symbol Ift140 provided by MGI

Official Full Name intraflagellar transport 140 provided by MGI

Primary source MGI:MGI:2146906

See related Ensembl: ENSMUSG00000024169

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 Al661311, Tce5, Wdtc2, mKIAA0590

Expression Broad expression in testis adult (RPKM 39.8), liver E14.5 (RPKM 18.4) and 23 other tissuesSee more

Orthologs human all

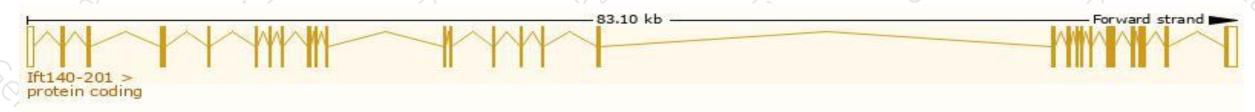
Transcript information (Ensembl)



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

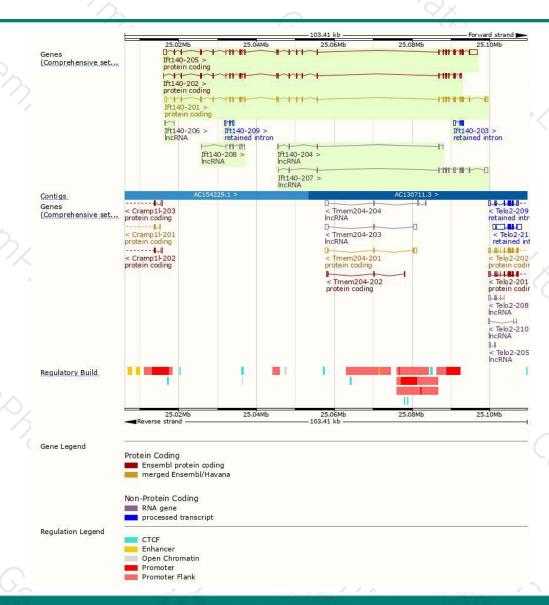
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Ift140-201	ENSMUST00000024983.11	5489	<u>1464aa</u>	Protein coding	CCDS28505	E9PY46	TSL:1 GENCODE basic APPRIS P2
Ift140-205	ENSMUST00000142000.8	6942	<u>1421aa</u>	Protein coding	8 7	A0A3B2W3F6	TSL:1 GENCODE basic APPRIS ALT2
Ift140-202	ENSMUST00000137386.7	3774	1232aa	Protein coding	β <u>L</u>	E9Q682	CDS 3' incomplete TSL:5
Ift140-203	ENSMUST00000139300.1	908	No protein	Retained intron	i i	726	TSL:3
lft140-209	ENSMUST00000232788.1	846	No protein	Retained intron	85	250	
lft140-207	ENSMUST00000153895.7	2938	No protein	IncRNA		793	TSL:1
Ift140-208	ENSMUST00000156945.1	740	No protein	IncRNA	94	(2)	TSL:3
lft140-204	ENSMUST00000140692.7	734	No protein	IncRNA	i a	323	TSL:3
Ift140-206	ENSMUST00000151776.1	343	No protein	IncRNA	85	257	TSL:2

The strategy is based on the design of *Ift140-201* transcript, The transcription is shown below



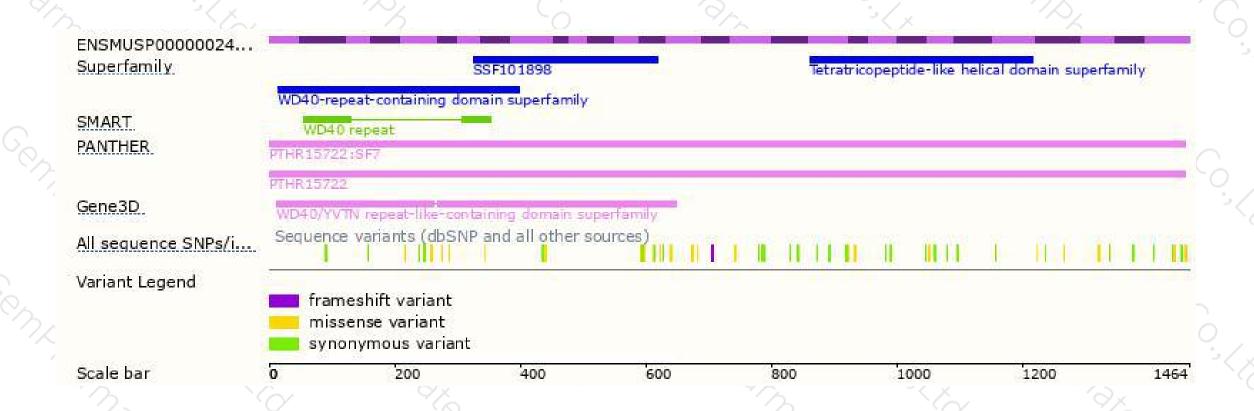
Genomic location distribution





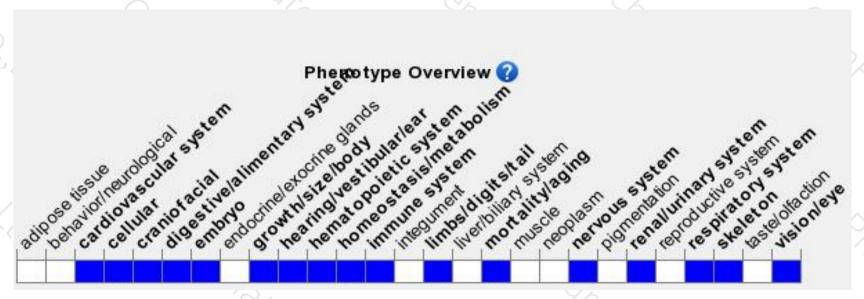
Protein domain





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 reporter knock-out allele die at mid-gestation. Mice homozygous for an ENU-induced mutation exhibit cardiovascular defects and situs abnormalities.



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





