

Ift80 Cas9-KO Strategy

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Reviewer:

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Design Date:

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Project Overview



Project Name

Ift80

Project type

Cas9-KO

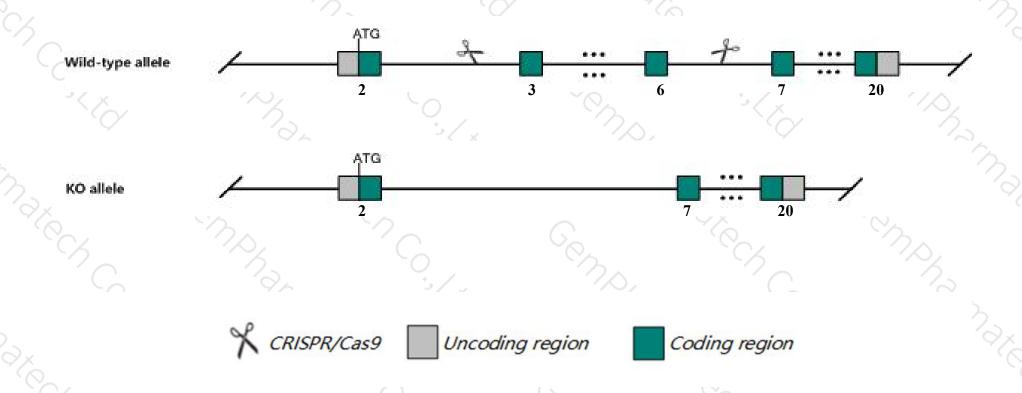
Strain background

C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Ift80 gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Ift80* gene has 12 transcripts. According to the structure of *Ift80* gene, exon3-exon6 of *Ift80-202* (ENSMUST00000107812.7) transcript is recommended as the knockout region. The region contains 512bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Ift80* gene. The brief process is as follows: CRISPR/Cas9 system v

Notice



- > According to the existing MGI data, Mice homozygous for a hypomorphic gene trap allele exhibit partial perinatal lethality, decreased body size, postnatal growth retardation, shortened long bones, constricted thoracic cage, periaxial polydactyly, and small cranium.
- > The *Ift80* gene is located on the Chr3. 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)



Ift80 intraflagellar transport 80 [Mus musculus (house mouse)]

Gene ID: 68259, updated on 31-Jan-2019

Summary

☆ ?

Official Symbol Iff80 provided by MGI

Official Full Name intraflagellar transport 80 provided by MGI

Primary source MGI:MGI:1915509

See related Ensembl:ENSMUSG00000027778

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 4921524P20Rik, Wdr56, mKIAA1374

Expression Broad expression in testis adult (RPKM 8.7), CNS E11.5 (RPKM 6.3) and 21 other tissuesSee more

Orthologs <u>human</u> all

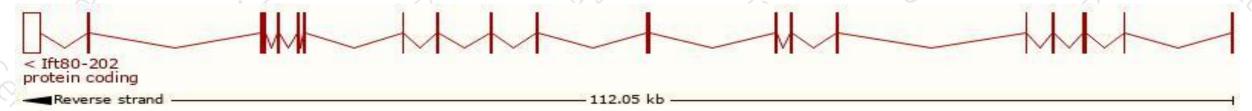
Transcript information (Ensembl)



The gene has 12 transcripts, all transcripts are shown below:

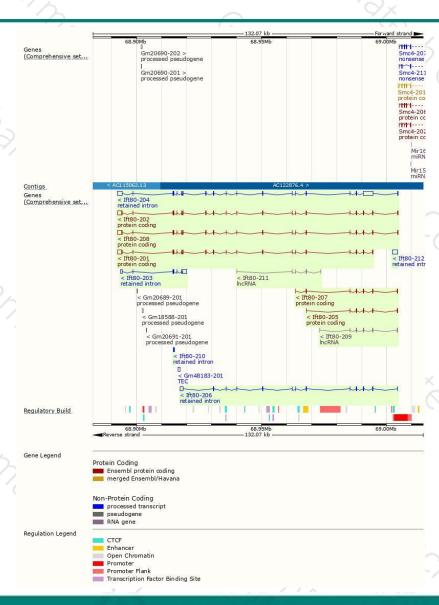
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Ift80-202	ENSMUST00000107812.7	4063	<u>777aa</u>	Protein coding	CCDS38451	Q8K057	TSL:5 GENCODE basic APPRIS P1
Ift80-201	ENSMUST00000029347.13	3880	<u>777aa</u>	Protein coding	CCDS38451	Q8K057	TSL:5 GENCODE basic APPRIS P1
Ift80-208	ENSMUST00000169064.7	4064	<u>777aa</u>	Protein coding	2	Q8K057	TSL:1 GENCODE basic APPRIS P1
Ift80-207	ENSMUST00000154741.7	755	209aa	Protein coding	-	D3YWI8	CDS 3' incomplete TSL:5
Ift80-205	ENSMUST00000148031.1	741	<u>158aa</u>	Protein coding	ē	D3Z157	CDS 3' incomplete TSL:5
Ift80-204	ENSMUST00000136448.7	7616	No protein	Retained intron	5	-	TSL:2
Ift80-203	ENSMUST00000136176.1	2860	No protein	Retained intron	2	(4)	TSL:1
Ift80-206	ENSMUST00000152502.7	2608	No protein	Retained intron	-	7527	TSL:5
Ift80-212	ENSMUST00000192467.1	2231	No protein	Retained intron		-	TSL:NA
Ift80-210	ENSMUST00000176189.1	241	No protein	Retained intron	5	-	TSL:3
Ift80-209	ENSMUST00000176135.1	653	No protein	IncRNA	2	(4)	TSL:3
Ift80-211	ENSMUST00000176754.1	525	No protein	IncRNA	2	752.5	TSL:2

The strategy is based on the design of *Ift80-202* transcript, The transcription is shown below



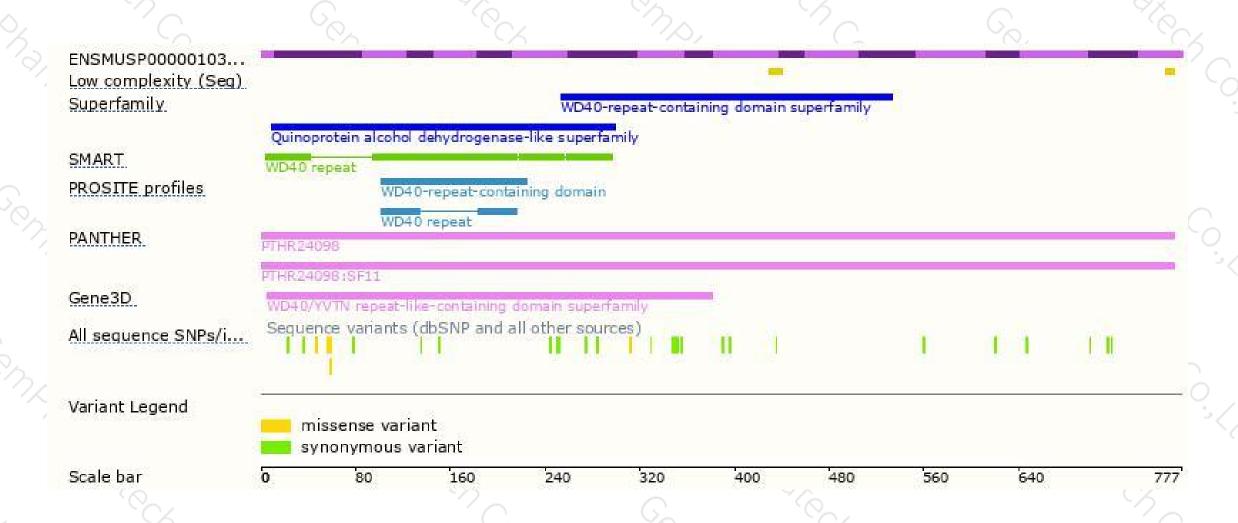
Genomic location distribution





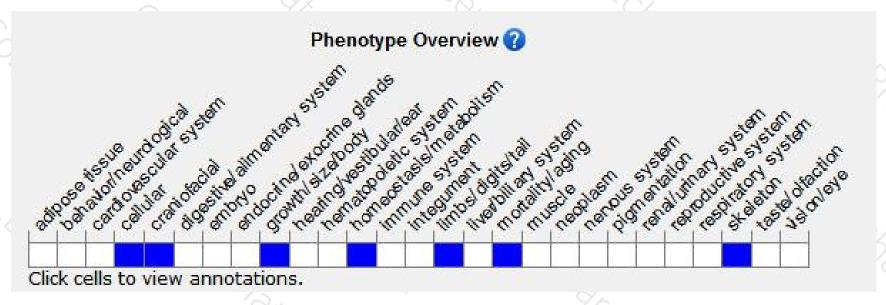
Protein domain





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, Mice homozygous for a hypomorphic gene trap allele exhibit partial perinatal lethality, decreased body size, postnatal growth retardation, shortened long bones, constricted thoracic cage, periaxial polydactyly, and small cranium.



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





