

Ywhaq Cas9-CKO Strategy

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Reviewer:

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Design Date:

2019-10-16

Project Overview



Project Name

Ywhaq

Project type

Cas9-CKO

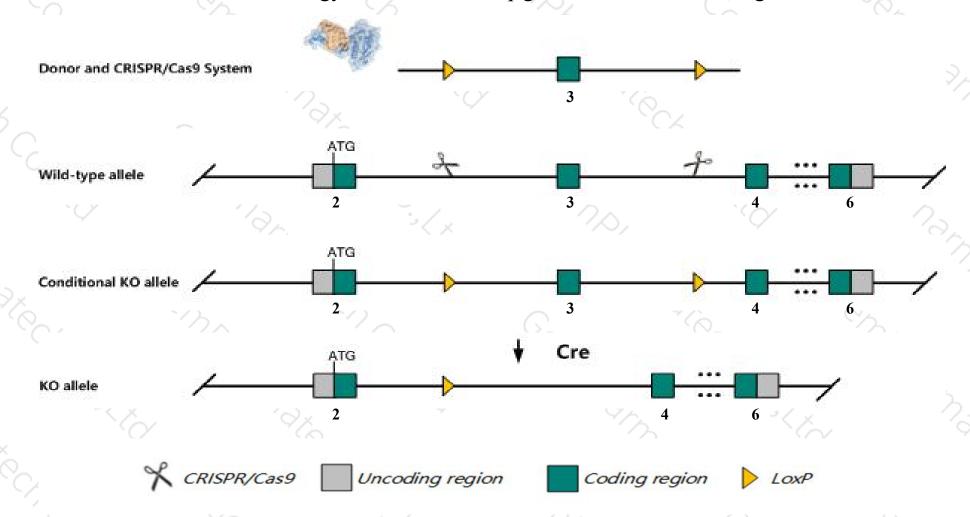
Strain background

C57BL/6JGpt

Conditional Knockout strategy



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



Technical routes



- The *Ywhaq* gene has 5 transcripts. According to the structure of *Ywhaq* gene, exon3 of *Ywhaq-202*(ENSMUST00000135088.8) transcript is recommended as the knockout region. The region contains 124bp coding sequence.

 Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Ywhaq* 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, Embryos homozygous for a gene trap allele are developmentally delayed and die by E14 with no specific cardiac defects; however, heterozygotes develop larger myocardial infarctions with increased post-infarction cardiac remodeling while cultured cardiomyocytes are sensitized to proapoptotic stimuli.
- The Ywhaq gene is located on the Chr12. 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)



Ywhaq tyrosine 3-monooxygenase/tryptophan 5-monooxygenase activation protein theta [Mus musculus (house mouse)]

Gene ID: 22630, updated on 10-Oct-2019

Summary

☆ ?

Official Symbol Ywhaq provided by MGI

Official Full Name tyrosine 3-monooxygenase/tryptophan 5-monooxygenase activation protein theta provided by MGI

Primary source MGI:MGI:891963

See related Ensembl: ENSMUSG00000076432

Gene type protein coding
RefSeq status PROVISIONAL
Organism Mus musculus

Lineage Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae;

Murinae; Mus; Mus

Also known as R74690; AA409740; AU021156; 2700028P07Rik

Expression Ubiquitous expression in CNS E18 (RPKM 170.8), CNS E14 (RPKM 138.7) and 28 other tissues See more

Orthologs human all

Transcript information (Ensembl)



The gene has 5 transcripts, all transcripts are shown below:

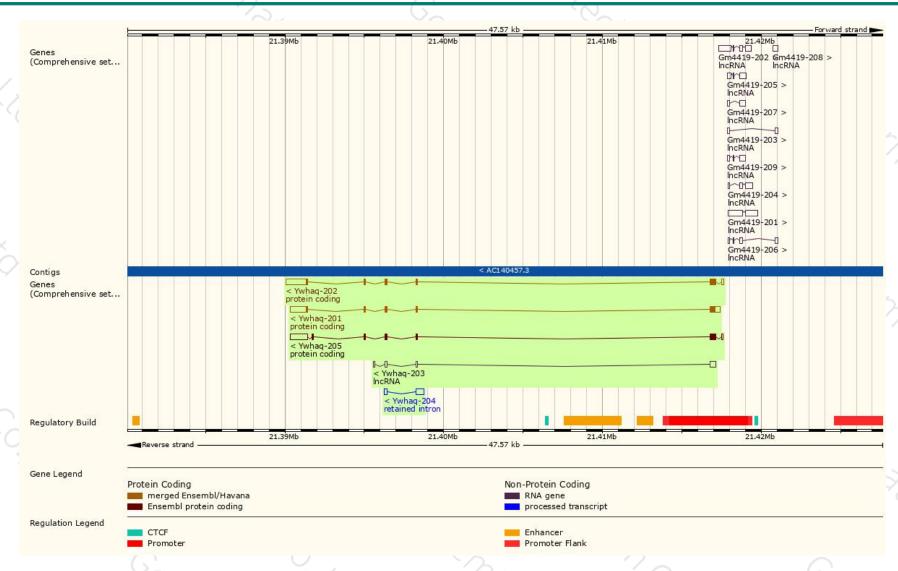
Name A	Transcript ID	bp 🌲	Protein	Biotype 🍦	CCDS 🍦	UniProt	Flags
Ywhaq-201	ENSMUST00000103002.7	2110	245aa	Protein coding	CCDS25837 ₺	A3KML3 ₽ P68254 ₽	TSL:1 GENCODE basic APPRIS P1
Ywhaq-202	ENSMUST00000135088.8	2197	245aa	Protein coding	CCDS25837 ₺	<u>A3KML3</u> & <u>F6VW30</u> & <u>P68254</u> &	TSL:1 GENCODE basic APPRIS P1
Ywhaq-203	ENSMUST00000140710.1	703	No protein	IncRNA	979		TSL:3
Ywhaq-204	ENSMUST00000146206.1	682	No protein	Retained intron	-	-	TSL:2
Ywhaq-205	ENSMUST00000155480.8	1970	243aa	Protein coding	1570	F6YY69 & P68254 &	TSL:1 GENCODE basic

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



Genomic location distribution





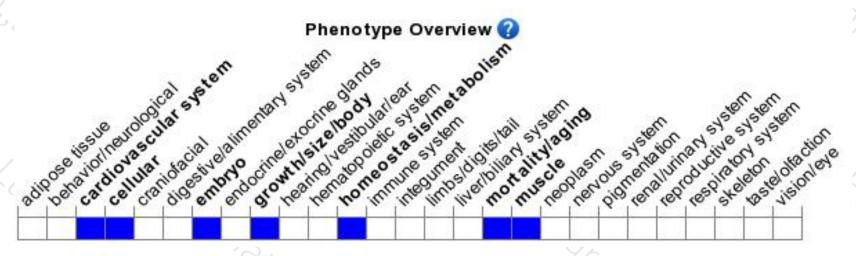
Protein domain





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, embryos homozygous for a gene trap allele are developmentally delayed and die by E14 with no specific cardiac defects; however, heterozygotes develop larger myocardial infarctions with increased post-infarction cardiac remodeling while cultured cardiomyocytes are sensitized to proapoptotic stimuli.



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





