Chchd10 Cas9-KO Strategy

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Design Date: 2019-8-5

Project Overview



Project Name

Chchd10

Project type

Cas9-KO

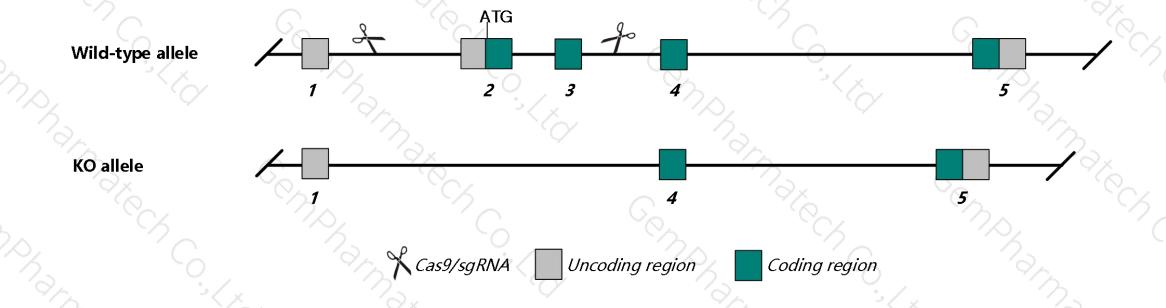
Strain background

C57BL/6J

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Chchd10* gene. The schematic diagram is as follows:



Technical routes



- The *Chchd10* gene has 3 transcripts. According to the structure of *Chchd10* gene, exon2-exon3 of *Chchd10*-203 (ENSMUST00000219839.1) transcript is recommended as the knockout region. The region contains start codon ATG. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Chchd10* gene. The brief process is as follows: sgRNA was transcribed in vitro.Cas9, sgRNA were microinjected into the fertilized eggs of C57BL/6J 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/6J mice.

Notice



- According to the existing MGI data, Homozygous deletion results in mild mitochondrial respiration anomalies in skeletal muscle.
- ➤ The KO region contains functional region of the *Mmp11* gene.Knockout the region may affect the function of *Mmp11* gene.
- ➤ The *Chchd10* gene is located on the Chr10. 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)



Chchd10 coiled-coil-helix-coiled-coil-helix domain containing 10 [Mus musculus (house mouse)]

Gene ID: 103172, updated on 23-Oct-2018

Summary

Official Symbol Chchd10 provided by MGI

Official Full Name coiled-coil-helix-coiled-coil-helix domain containing 10 provided by MGI

Primary source MGI:MGI:2143558

See related Ensembl: ENSMUSG00000049422

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 Ndg2; Al267078; 1620401E04Rik

Expression Biased expression in adrenal adult (RPKM 1343.9), duodenum adult (RPKM 646.7) and 10 other tissues See more

Orthologs human all

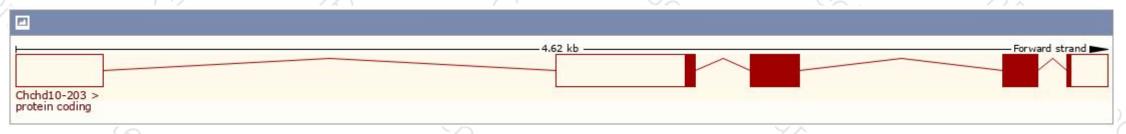
Transcript information (Ensembl)



The gene has 3 transcripts, and all transcripts are shown below:

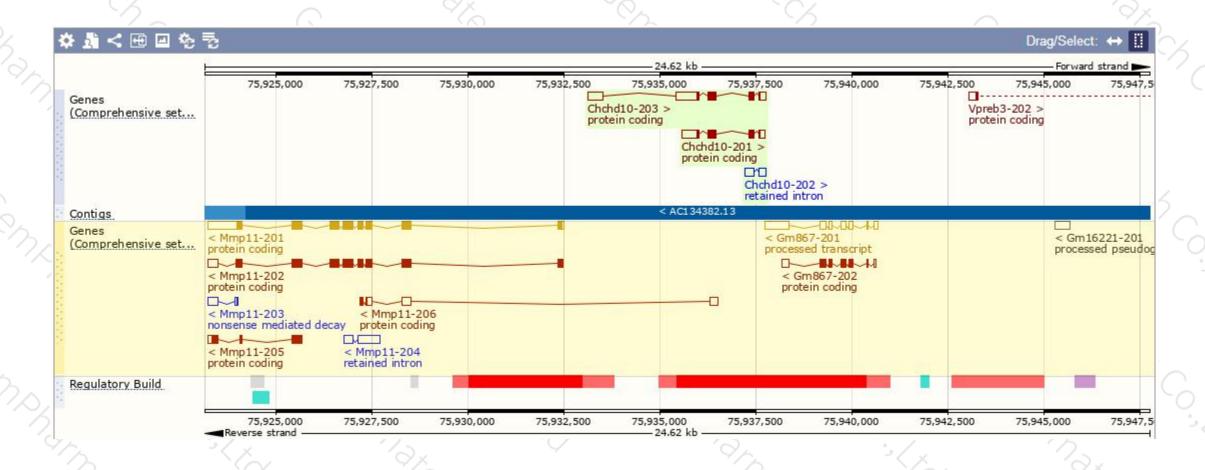
Show/hide columns (1 hidden)										
Name	Transcript ID	bp 🛊	Protein	Biotype	CCDS	UniProt 4	RefSeq		Flags	4
Chchd10-203	ENSMUST00000219839.1	1488	<u>138aa</u>	Protein coding	CCDS23938 ₽	Q7TNL9₽	120	TSL:1	GENCODE basic	APPRIS P1
Chchd10-201	ENSMUST00000058906.6	939	<u>138aa</u>	Protein coding	CCDS23938 €	Q7TNL9@	NM_175329@ NP_780538@	TSL:1	GENCODE basic	APPRIS P1
Chchd10-202	ENSMUST00000219435.1	405	No protein	Retained intron	-	-	-		TSL:2	

The strategy is based on the design of Chchd10-203 transcript, The transcription is shown below



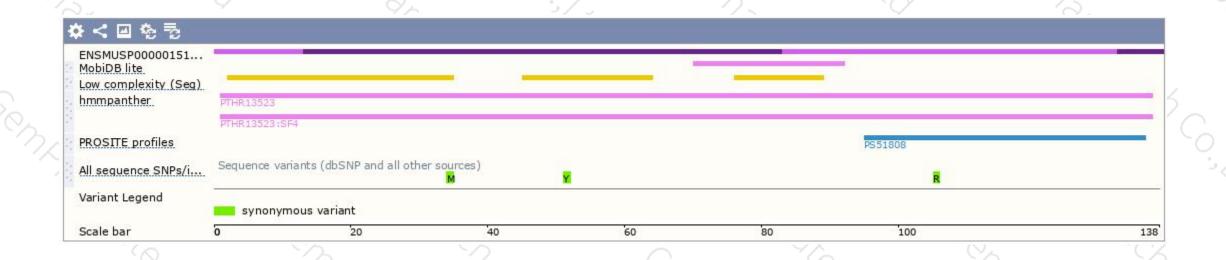
Genomic location distribution





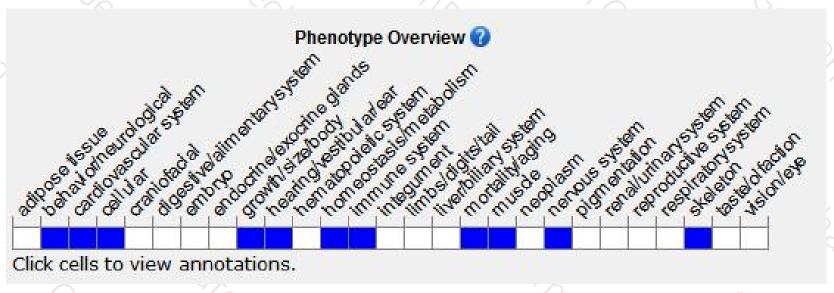
Protein domain





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, Homozygous deletion results in mild mitochondrial respiration anomalies in skeletal muscle.

If you have any questions, you are welcome to inquire. Tel: 025-5864 1534





