

Cdc14b Cas9-KO Strategy

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Design Date: 2019-7-22

Project Overview



Project Name

Cdc14b

Project type

Cas9-KO

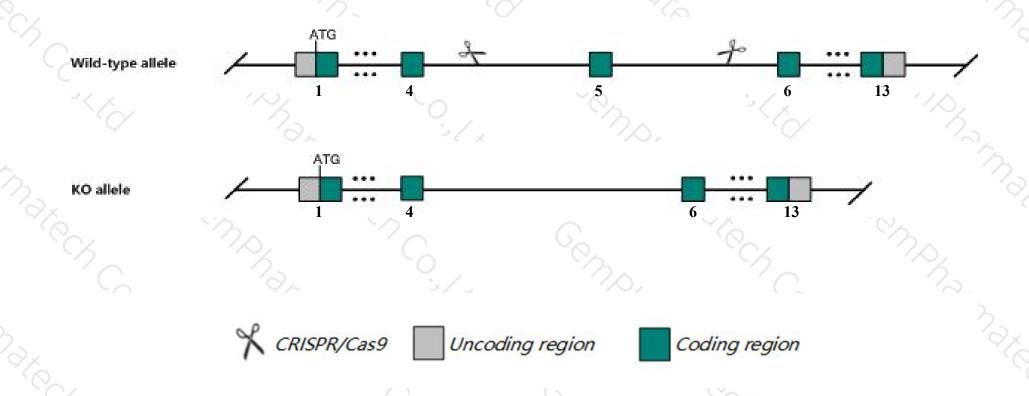
Strain background

C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Cdc14b* gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Cdc14b* gene has 11 transcripts. According to the structure of *Cdc14b* gene, exon5 of *Cdc14b-201* (ENSMUST00000039318.15) transcript is recommended as the knockout region. The region contains 77bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Cdc14b* gene. The brief process is as follows: CRISPR/Cas9 system

Notice



- ➤ According to the existing MGI data, Mice homozygous for a knock-out allele exhibit premature aging, including premature cataracts and kyphosis; reduced fertility, particularly in female mice; and impaired contextual conditioning.
- The *Cdc14b* gene is located on the Chr13. 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)



Cdc14b CDC14 cell division cycle 14B [Mus musculus (house mouse)]

Gene ID: 218294, updated on 31-Jan-2019

Summary

☆ ?

Official Symbol Cdc14b provided by MGI

Official Full Name CDC14 cell division cycle 14B provided by MGI

Primary source MGI:MGI:2441808

See related Ensembl:ENSMUSG00000033102

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 2810432N10Rik, A530086E13Rik, AA472821, CDC14B3, Cdc14B1

Expression Ubiquitous expression in testis adult (RPKM 6.1), placenta adult (RPKM 2.9) and 28 other tissuesSee more

Orthologs <u>human</u> all

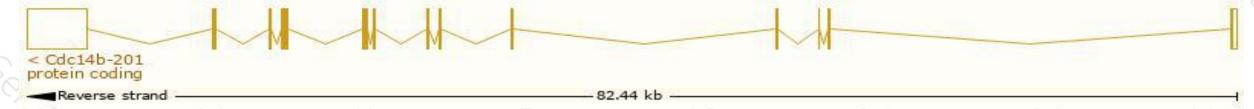
Transcript information (Ensembl)



The gene has 11 transcripts, all transcripts are shown below:

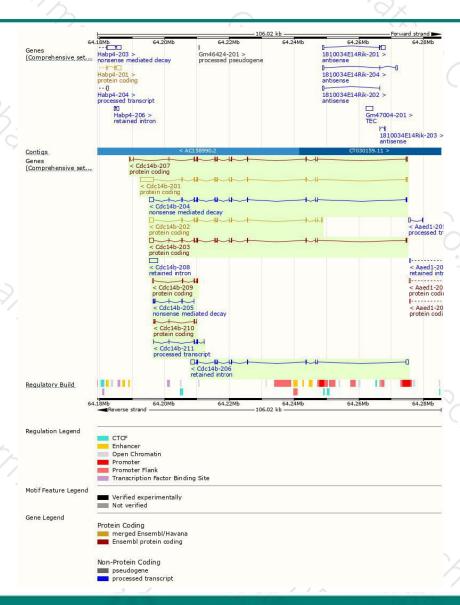
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Cdc14b-201	ENSMUST00000039318.15	5859	<u>485aa</u>	Protein coding	CCDS26598	Q6PFY9	TSL:1 GENCODE basic APPRIS P3
Cdc14b-203	ENSMUST00000109770.1	3052	485aa	Protein coding	CCDS26598	Q6PFY9	TSL:1 GENCODE basic APPRIS P3
Cdc14b-202	ENSMUST00000109769.9	2886	448aa	Protein coding	CCDS49292	Q6PFY9	TSL:1 GENCODE basic APPRIS ALT2
Cdc14b-207	ENSMUST00000221634.1	1953	492aa	Protein coding	29	A0A1Y7VJB9	TSL:1 GENCODE basic
Cdc14b-209	ENSMUST00000222713.1	790	<u>153aa</u>	Protein coding	5	A0A1Y7VJ26	CDS 5' incomplete TSL:3
Cdc14b-210	ENSMUST00000222766.1	474	<u>124aa</u>	Protein coding	*	A0A1Y7VKG2	CDS 5' incomplete TSL:3
Cdc14b-204	ENSMUST00000221139.1	3075	<u>485aa</u>	Nonsense mediated decay	-	Q6PFY9	TSL:1
Cdc14b-205	ENSMUST00000221217.1	628	<u>63aa</u>	Nonsense mediated decay	20	A0A1Y7VK12	CDS 5' incomplete TSL:3
Cdc14b-211	ENSMUST00000223116.1	752	No protein	Processed transcript	5	15	TSL:5
Cdc14b-206	ENSMUST00000221437.1	2841	No protein	Retained intron		89	TSL:1
Cdc14b-208	ENSMUST00000221788.1	2565	No protein	Retained intron	24	ķ <u>u</u>	TSL:NA
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The strategy is based on the design of Cdc14b-201 transcript, The transcription is shown below



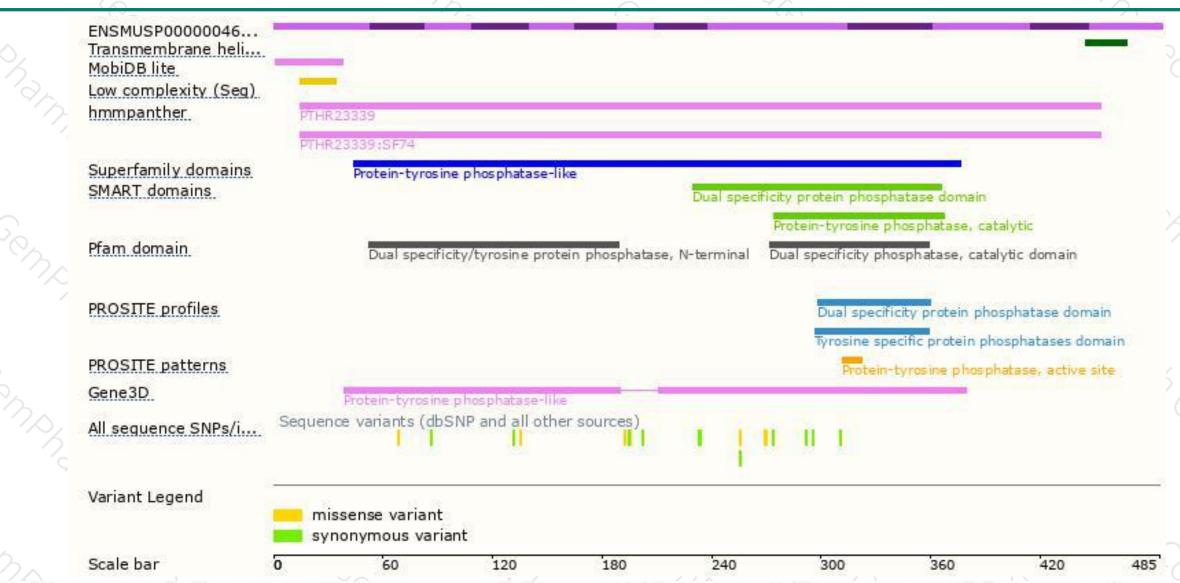
Genomic location distribution





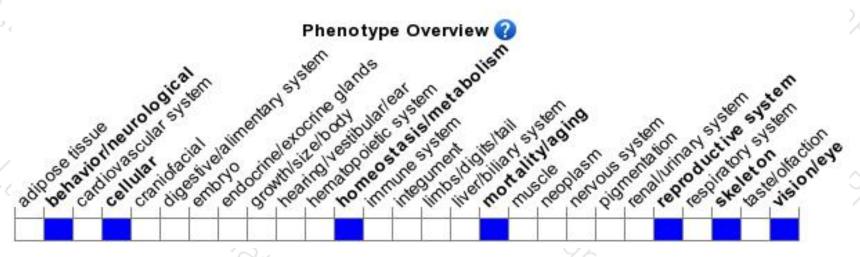
Protein domain





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 knock-out allele exhibit premature aging, including premature cataracts and kyphosis; reduced fertility, particularly in female mice; and impaired contextual conditioning.



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





