

Kidins220 Cas9-KO Strategy

Designer: Zihe Cui

Reviewer: Xueting Zhang

Design Date: 2020-7-22

Project Overview



Project Name

Kidins220

Project type

Cas9-KO

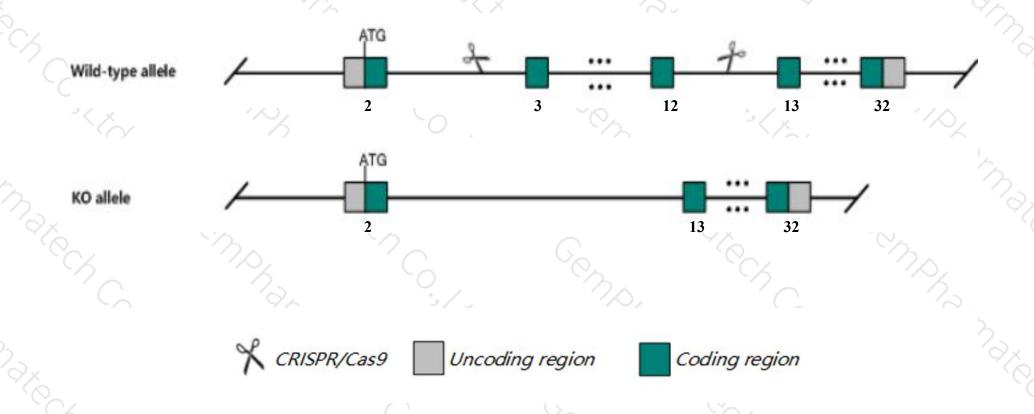
Strain background

C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Kidins220* gene. The schematic diagram is as follows:



Technical routes



- The *Kidins220* gene has 10 transcripts. According to the structure of *Kidins220* gene, exon3-exon12 of *Kidins220*-201(ENSMUST00000066652.6) transcript is recommended as the knockout region. The region contains 1168bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Kidins220* gene. The brief process is as follows: CRISPR/Cas9 system 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.

Notice



- > According to the existing MGI data,mice homozygous for a knock-out allele exhibit embryonic lethality. Mice heterozygous for a knock-out allele exhibit decreased dendritic complexity in the barrel somatosensory cortex and dentate gyrus neurons.
- ➤ Transcript *Kidins220*-209 may not be affected.
- > The *Kidins220* 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 the gene knockout on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.

Gene information (NCBI)



Kidins220 kinase D-interacting substrate 220 [Mus musculus (house mouse)]

Gene ID: 77480, updated on 26-Jun-2020

Summary



Official Symbol Kidins220 provided by MGI

Official Full Name kinase D-interacting substrate 220 provided by MGI

Primary source MGI:MGI:1924730

See related Ensembl: ENSMUSG00000036333

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 Al194387; Al316525; mKIAA1250; 3110039L19Rik; C330002l19Rik

Expression Broad expression in CNS E18 (RPKM 54.8), whole brain E14.5 (RPKM 42.9) and 25 other tissues See more

Orthologs <u>human</u> all

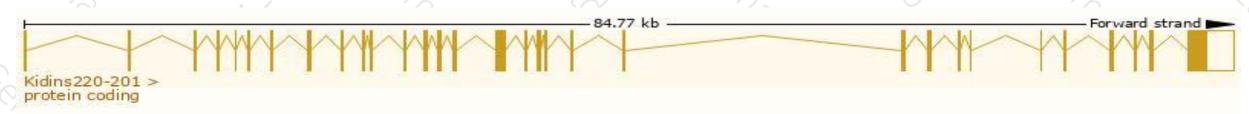
Transcript information (Ensembl)



The gene has 10 transcripts, all transcripts are shown below:

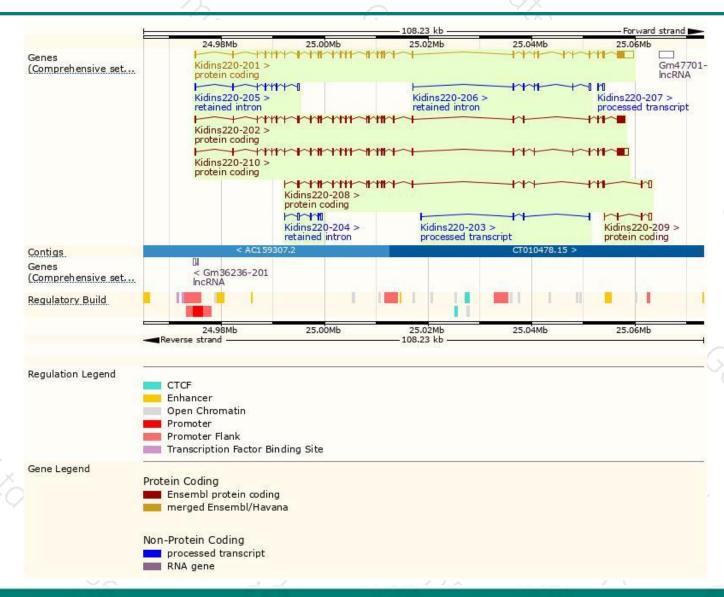
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Kidins220-201	ENSMUST00000066652.6	7409	1793aa	Protein coding	CCDS36424	E9Q9B7	TSL:5 GENCODE basic
Kidins220-210	ENSMUST00000222941.1	6277	<u>1763aa</u>	Protein coding	==	A0A1Y7VME9	TSL:5 GENCODE basic APPRIS P1
Kidins220-202	ENSMUST00000220459.1	5381	<u>1672aa</u>	Protein coding	-	A0A1Y7VMH7	TSL:1 GENCODE basic
Kidins220-208	ENSMUST00000222013.1	3998	<u>1182aa</u>	Protein coding	-	A0A1Y7VNF8	CDS 5' incomplete TSL:1
Kidins220-209	ENSMUST00000222481.1	627	<u>61aa</u>	Protein coding	-	A0A1Y7VK25	CDS 5' incomplete TSL:3
Kidins220-203	ENSMUST00000220622.1	736	No protein	Processed transcript	-	-	TSL:2
Kidins220-207	ENSMUST00000221622.1	440	No protein	Processed transcript		-	TSL:2
Kidins220-205	ENSMUST00000221378.1	1115	No protein	Retained intron	-	-	TSL:1
Kidins220-206	ENSMUST00000221423.1	947	No protein	Retained intron	-	-	TSL:1
Kidins220-204	ENSMUST00000221050.1	636	No protein	Retained intron	-8	-	TSL:2

The strategy is based on the design of *Kidins220-201* transcript, the transcription is shown below:



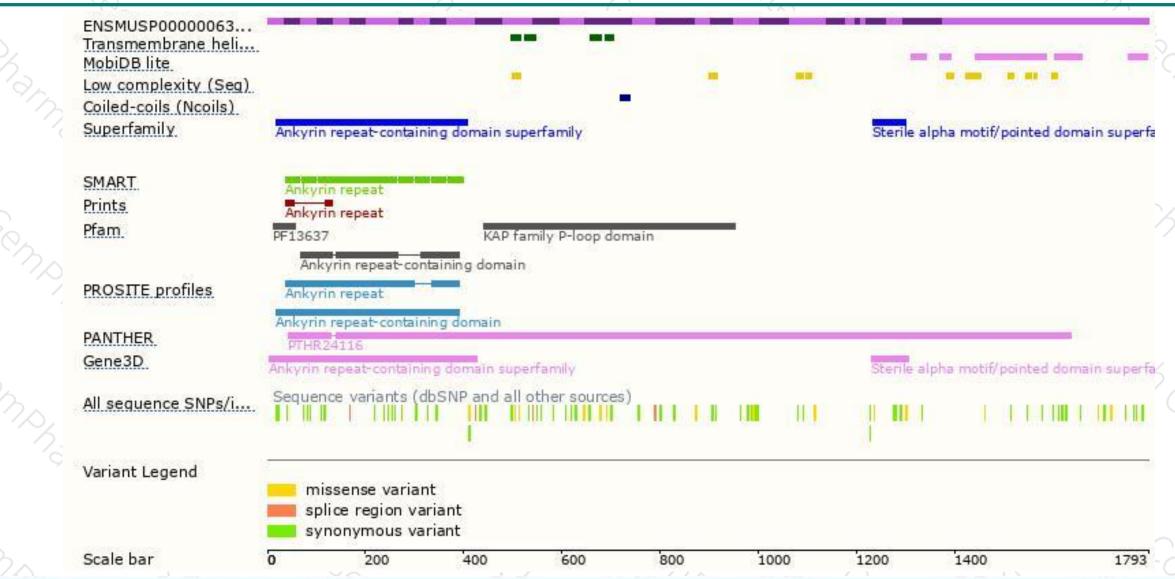
Genomic location distribution





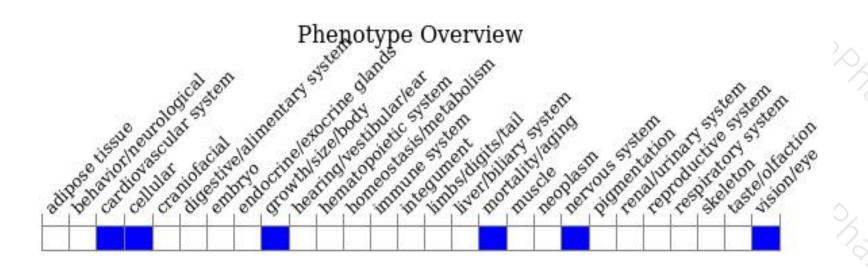
Protein domain





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 embryonic lethality. Mice heterozygous for a knock-out allele exhibit decreased dendritic complexity in the barrel somatosensory cortex and dentate gyrus neurons.



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





