Kdm6b CKO Strategy

Designer: Composition of Co. Line

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Project Overview



Project Name

Kdm6b

Project type

CKO

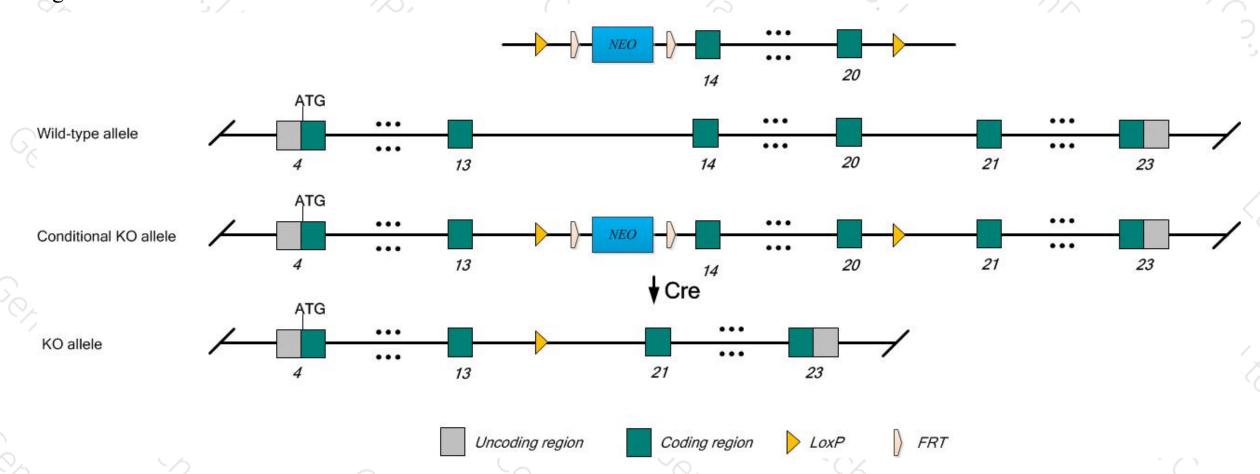
Strain background

C57BL/6N

Conditional Knockout strategy



This model will use ES cell mediated homologous recombination technology to edit the *Kdm6b* gene. The schematic diagram is as follows:



Technical routes



- The *Kdm6b* gene has 2 transcripts. According to the structure of *Kdm6b* gene, exon14-20 of *Kdm6b-201* (ENSMUST00000094077.4) transcript is recommended as the knockout region. The region contains 937bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use ES cell targeting and homologous recombination technology to modify *Kdm6b* gene. The brief process is as follows: the targeting vector is constructed in vitro and the ES cells are electroporated. The ES clones screened by PCR and southern are injected into the donor blastocysts and transplanted into the pseudo-pregnant mother to further develop chimeric mice. A stable F1 generation mouse model was obtained by mating chimeric mice with C57BL/6N mice.
- > The flox mice was knocked out after mating with mice expressing Cre recombinase, resulting in the loss of function of the target gene in specific tissues or cell types.

Notice



- According to the existing MGI data, Mice homozygous for a null allele show perinatal death, thick alveolar septum, and absence of air space in the lungs. Mice homozygous for a different null allele die neonatally displaying abnormal lung development, dwarfism, kyphosis, short limbs, and a severe delay in endochondral ossification.
- The KO region contains functional region of the *Kdm6bos-201* gene. Knockout the region may affect the function of *Kdm6bos-201* gene.
- The position of *Kdm6b* gene and *Tmem88-201* gene is adjacent. Knockout the region may affect the function of the *Tmem88-201* gene.
- The *Kdm6b* gene is located on the Chr11. 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 loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.

Gene information (NCBI)



Kdm6b KDM1 lysine (K)-specific demethylase 6B [Mus musculus (house mouse)]

Gene ID: 216850, updated on 3-Mar-2019

Summary

Official Symbol Kdm6b provided by MGI

Official Full Name KDM1 lysine (K)-specific demethylase 6B provided by MGI

Primary source MGI:MGI:2448492

See related Ensembl: ENSMUSG00000018476

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 Jmjd3; BC038313; 1700064E03Rik

Expression Ubiquitous expression in thymus adult (RPKM 16.9), duodenum adult (RPKM 10.2) and 25 other tissues See more

Orthologs human all

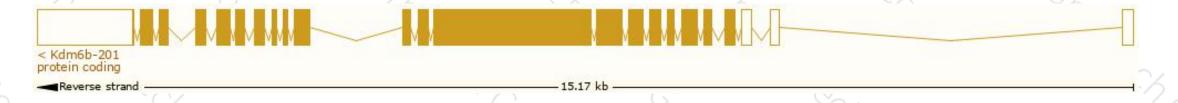
Transcript information (Ensembl)



The gene has 2 transcripts, and all transcripts are shown below:

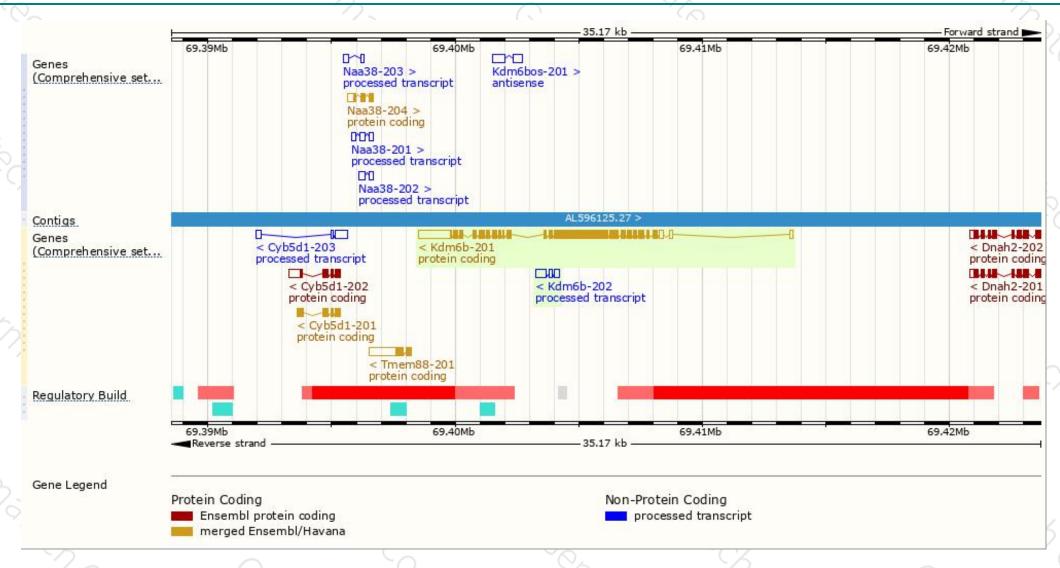
Name 🍦	Transcript ID 👙	bp 🌲	Protein 🍦	Biotype	CCDS 🍦	UniProt 🍦	Flags
Kdm6b-201	ENSMUST00000094077.4	6654	<u>1641aa</u>	Protein coding	<u>CCDS24895</u> ₽	Q5NCY0₽	TSL:5 GENCODE basic APPRIS P1
Kdm6b-202	ENSMUST00000156562.1	743	No protein	Processed transcript	÷	8 5	TSL:2

The strategy is based on the design of *Kdm6b*-201 transcript, The transcription is shown below



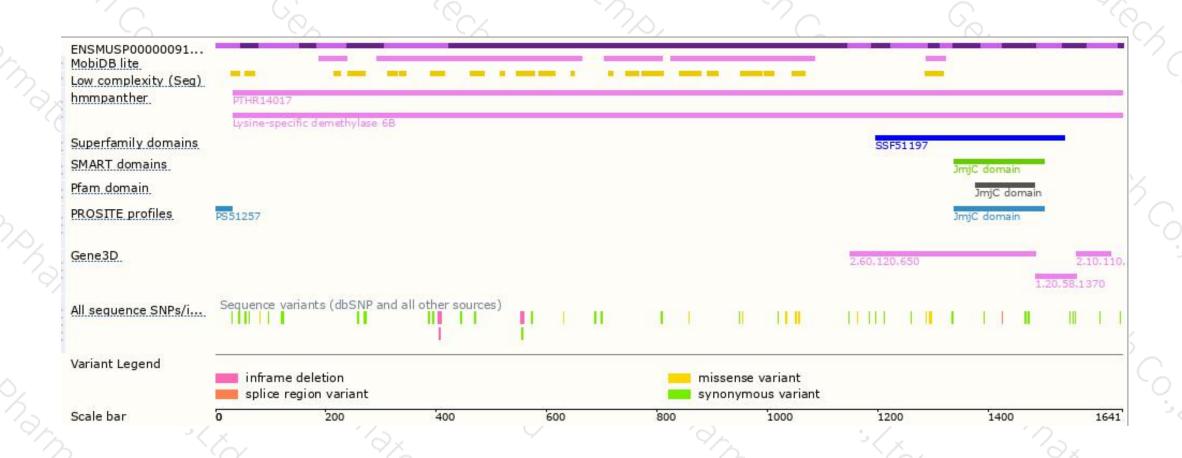
Genomic location distribution





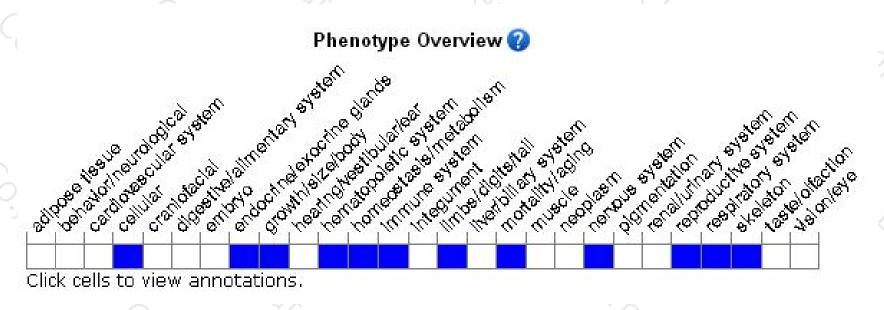
Protein domain





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 null allele show perinatal death, thick alveolar septum, and absence of air space in the lungs. Mice homozygous for a different null allele die neonatally displaying abnormal lung development, dwarfism, kyphosis, short limbs, and a severe delay in endochondral ossification.

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





