

# Klf14 Cas9-CKO Strategy

**Designer:** 

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**Design Date:** 

2019-7-26

# **Project Overview**



**Project Name** 

*Klf14* 

**Project type** 

Cas9-CKO

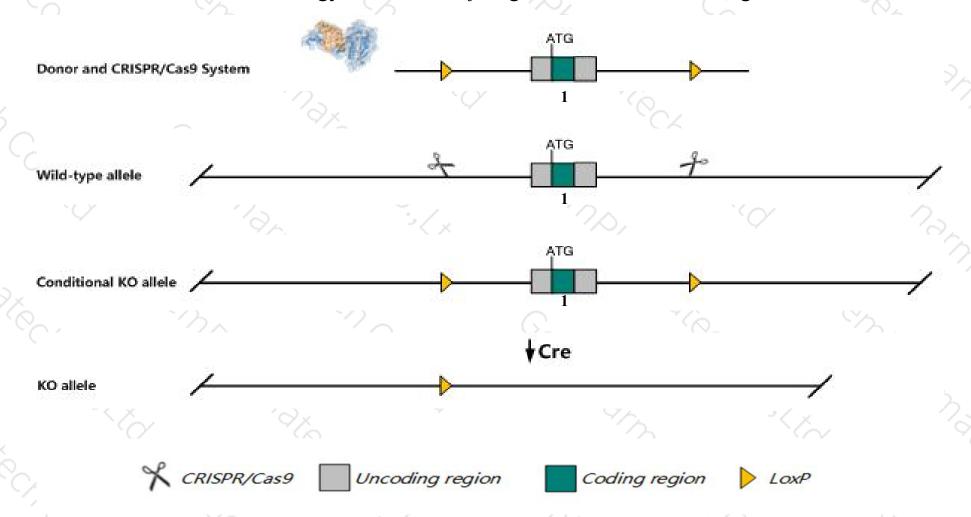
Strain background

C57BL/6JGpt

# Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Klf14* gene. The schematic diagram is as follows:



### **Technical routes**



- The *Klf14* gene has 1 transcript. According to the structure of *Klf14* gene, exon1 of *Klf14-201* (ENSMUST00000101589.4) transcript is recommended as the knockout region. The region contains all of the coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Klf14* 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, Mice homozygous for a null allele display genome instability and increased tumor incidence with age. Mice homozygous for a conditional allele knocked out in the liver exhibit decreased circulating HDL cholesterol levels.
- ➤ The *Klf14* gene is located on the Chr6. 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)



#### KIf14 Kruppel-like factor 14 [Mus musculus (house mouse)]

Gene ID: 619665, updated on 3-Feb-2019

#### Summary

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Official Symbol Klf14 provided by MGI

Official Full Name Kruppel-like factor 14 provided by MGI

Primary source MGI:MGI:3577024

See related Ensembl:ENSMUSG00000073209

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 5330411L03Rik, AL022736, BTEB5

Orthologs human all

# Transcript information (Ensembl)



The gene has 1 transcript, and the transcript is shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags	l
KIf14-201	ENSMUST00000101589.4	3056	325aa	Protein coding	CCDS51744	Q19A41	TSL:NA GENCODE basic APPRIS P1	

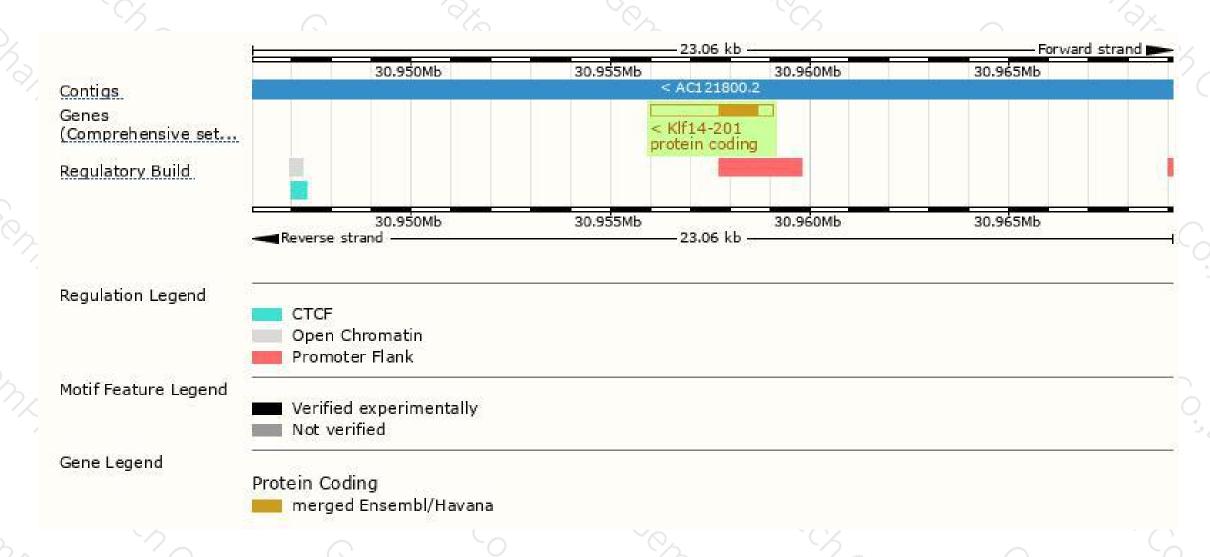
The strategy is based on the design of *Klf14-201* transcript, The transcription is shown below

< Klf14-201
protein coding

Reverse strand — 3.06 kb —

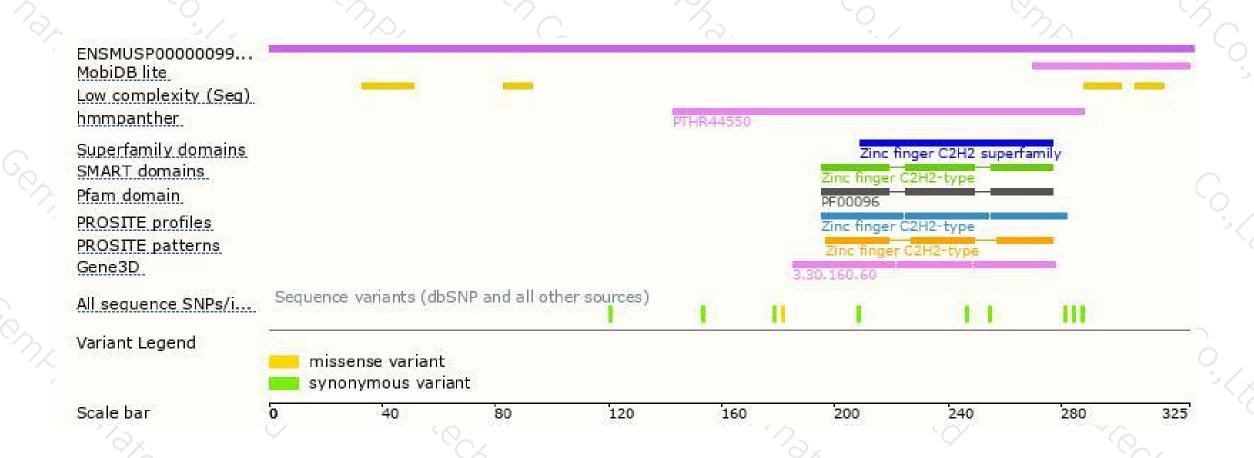
### Genomic location distribution





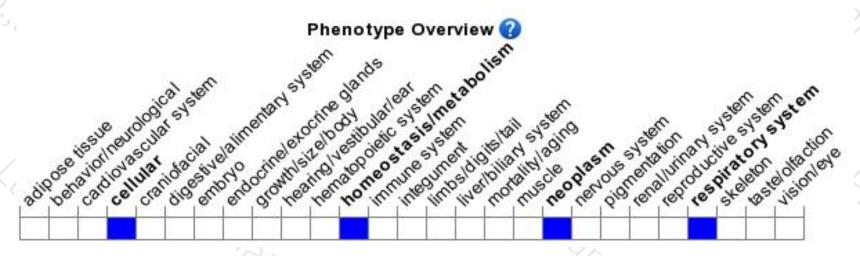
### Protein domain





# 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 display genome instability and increased tumor incidence with age. Mice homozygous for a conditional allele knocked out in the liver exhibit decreased circulating HDL cholesterol levels.



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





