

# Liph Cas9-KO Strategy

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Reviewer: Huimin Su

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# **Project Overview**



**Project Name** 

Liph

**Project type** 

Cas9-KO

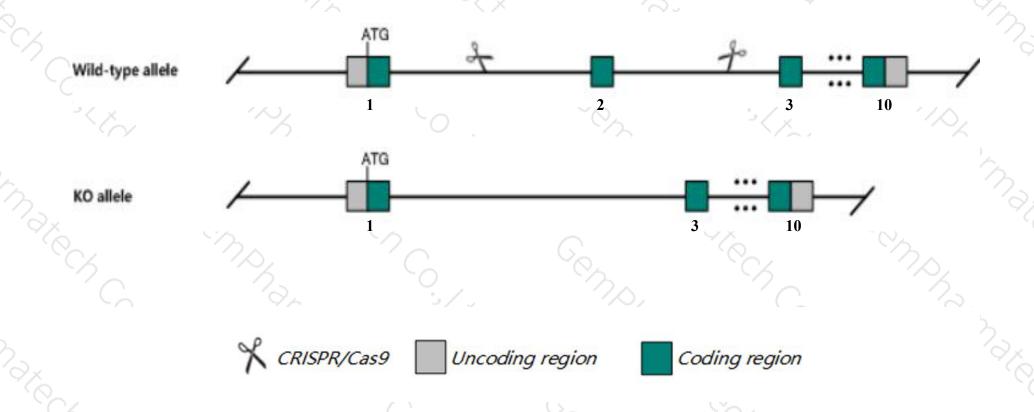
Strain background

C57BL/6JGpt

# **Knockout strategy**



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



### **Technical routes**



- ➤ The *Liph* gene has 6 transcripts. According to the structure of *Liph* gene, exon2 of *Liph-201*(ENSMUST00000060673.7) transcript is recommended as the knockout region. The region contains 368bp coding sequence.

  Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Liph* gene. The brief process is as follows: CRISPR/Cas9 system v

### **Notice**



- > According to the existing MGI data, Mice homozygous for a knock-out allele exhibit wavy vibrissae and wavy and matted coats associated with impaired inner rooth sheath formation.
- The *Liph* gene is located on the Chr16. 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)



#### Liph lipase, member H [ Mus musculus (house mouse) ]

Gene ID: 239759, updated on 13-Mar-2020

#### Summary

☆ ?

Official Symbol Liph provided by MGI

Official Full Name lipase, member H provided by MGI

Primary source MGI:MGI:2388029

See related Ensembl: ENSMUSG00000044626

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 P3; H06; lpd1; lpd2; Lpdlr; PLA1B; mPA-PLA1; D16Wsu119e; C130037N08Rik

Expression Biased expression in placenta adult (RPKM 8.8), large intestine adult (RPKM 8.0) and 7 other tissues See more

Orthologs human all

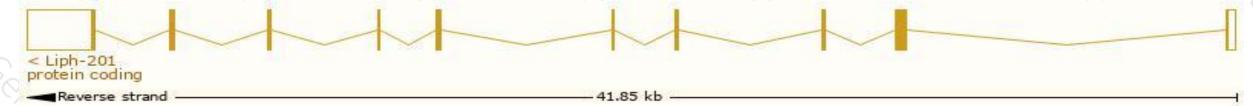
# Transcript information (Ensembl)



The gene has 6 transcripts, all transcripts are shown below:

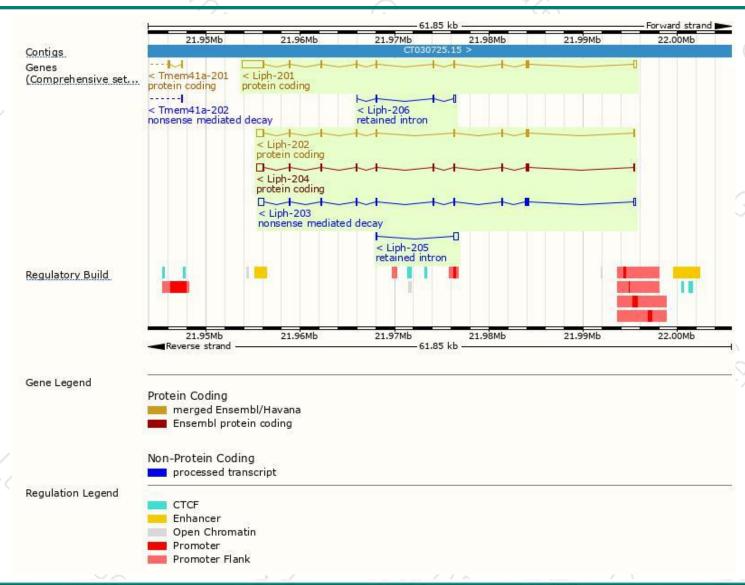
| Name 🍦   | Transcript ID 👙       | bp 🍦 | Protein 🍦  | Biotype                 | CCDS       | UniProt #           | Flags                         |
|----------|-----------------------|------|------------|-------------------------|------------|---------------------|-------------------------------|
| Liph-201 | ENSMUST00000060673.7  | 3923 | 451aa      | Protein coding          | CCDS37296₽ | Q8CIV3函             | TSL:1 GENCODE basic APPRIS P2 |
| Liph-202 | ENSMUST00000074230.11 | 2051 | 421aa      | Protein coding          | CCDS37295₽ | Q8CIV3個             | TSL:1 GENCODE basic           |
| Liph-204 | ENSMUST00000231766.1  | 2118 | 449aa      | Protein coding          |            | Q8CIV3₽             | GENCODE basic APPRIS ALT1     |
| Liph-203 | ENSMUST00000231682.1  | 2065 | 146aa      | Nonsense mediated decay | -          | <u>A0A338P6M6</u> ₽ | -                             |
| Liph-205 | ENSMUST00000232120.1  | 690  | No protein | Retained intron         | -          | 856                 | •                             |
| Liph-206 | ENSMUST00000232673.1  | 535  | No protein | Retained intron         | -          | 856                 | -                             |

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



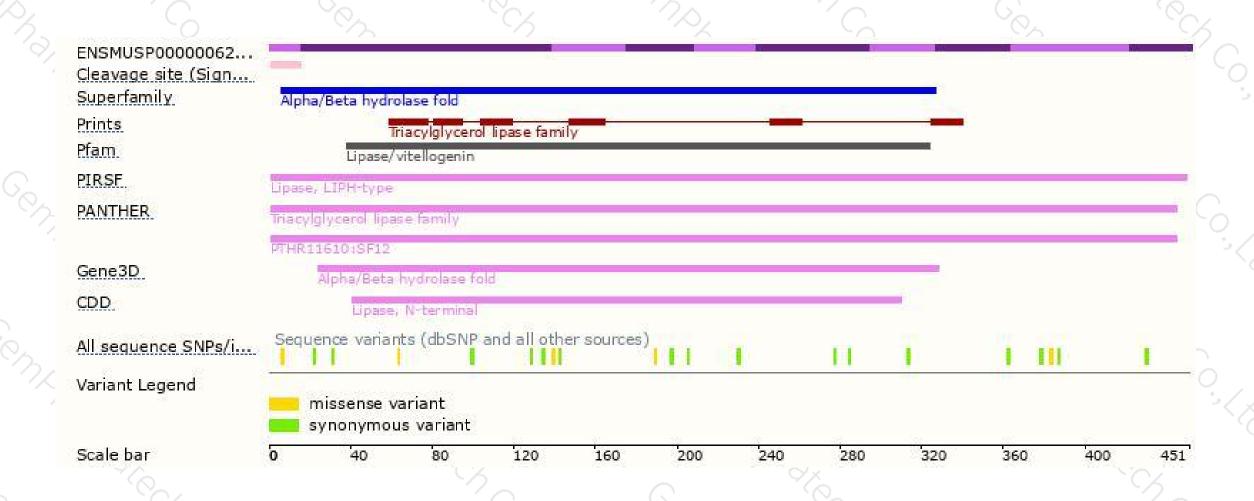
### Genomic location distribution





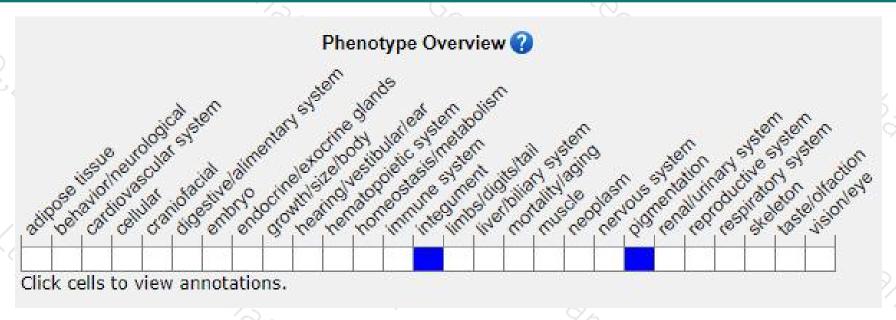
### Protein domain





## 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 wavy vibrissae and wavy and matted coats associated with impaired inner rooth sheath formation.



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





