

# Mlxip Cas9-CKO Strategy

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# **Project Overview**



Project Name Mlxip

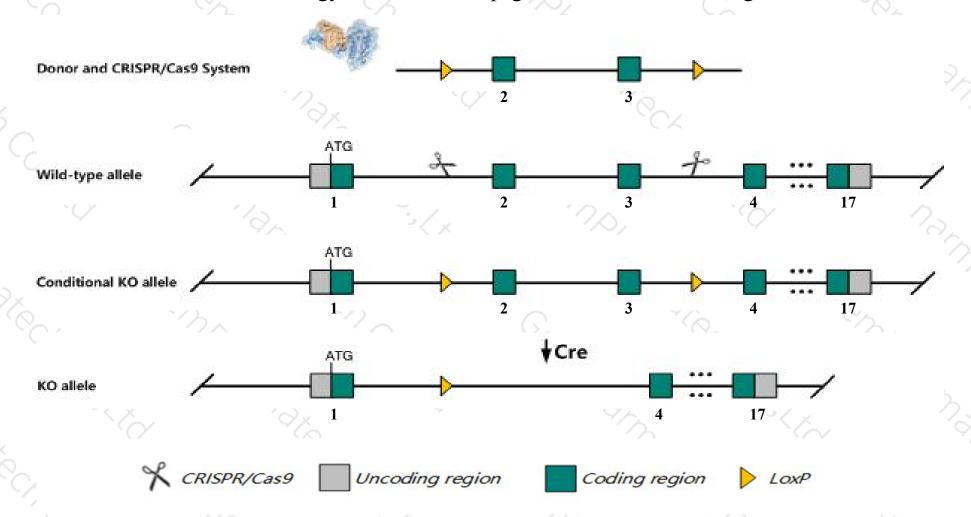
Project type Cas9-CKO

Strain background C57BL/6JGpt

## Conditional Knockout strategy



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



### Technical routes



- The *Mlxip* gene has 4 transcripts. According to the structure of *Mlxip* gene, exon2-exon3 of *Mlxip-201* (ENSMUST00000068237.11) transcript is recommended as the knockout region. The region contains 193bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Mlxip* 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 metabolic and performance abnormalities in response to mild exercise and improved sprint exercise performance.
- The *Mlxip* gene is located on the Chr5. 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)



#### Mixip MLX interacting protein [Mus musculus (house mouse)]

Gene ID: 208104, updated on 31-Jan-2019

#### Summary

☆ ?

Official Symbol Mixip provided by MGI

Official Full Name MLX interacting protein provided by MGI

Primary source MGI:MGI:2141183

See related Ensembl:ENSMUSG00000038342

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 AW228700, Mir, bHLHe36

Expression Ubiquitous expression in spleen adult (RPKM 18.7), colon adult (RPKM 18.0) and 28 other tissuesSee more

Orthologs human all

# Transcript information (Ensembl)



The gene has 4 transcripts, all transcripts are shown below:

| Name      | Transcript ID         | bp   | Protein      | Biotype                 | CCDS      | UniProt | Flags                         |
|-----------|-----------------------|------|--------------|-------------------------|-----------|---------|-------------------------------|
| Mixip-201 | ENSMUST00000068237.11 | 7303 | <u>917aa</u> | Protein coding          | CCDS19662 | G5E8D8  | TSL:1 GENCODE basic APPRIS P1 |
| MIxip-202 | ENSMUST00000111596.7  | 2653 | <u>614aa</u> | Protein coding          | CCDS39268 | B2RQ56  | TSL:1 GENCODE basic           |
| Mixip-203 | ENSMUST00000135961.1  | 5510 | 241aa        | Nonsense mediated decay |           | F6UPM1  | CDS 5' incomplete TSL:1       |
| Mixip-204 | ENSMUST00000199458.1  | 405  | No protein   | IncRNA                  | · ·       | 20      | TSL:3                         |

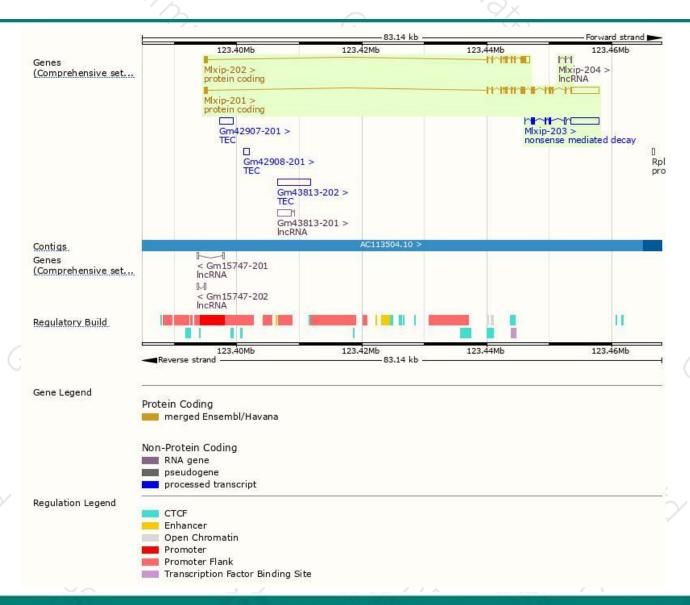
The strategy is based on the design of Mlxip-201 transcript, The transcription is shown below

63.13 kb

Forward strand 
Mixip-201 > protein coding

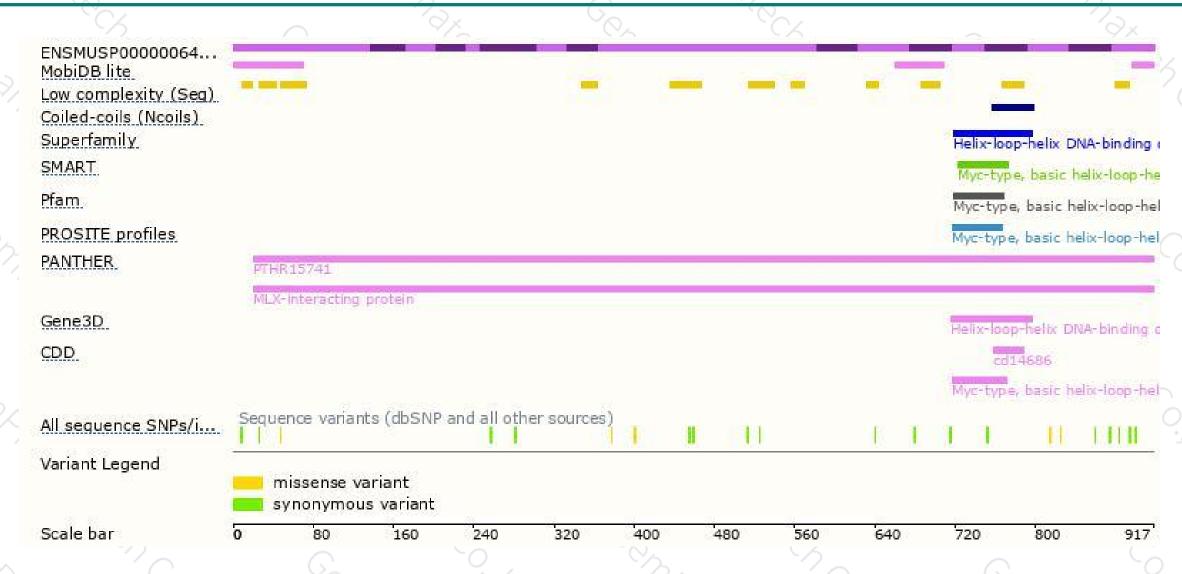
### Genomic location distribution





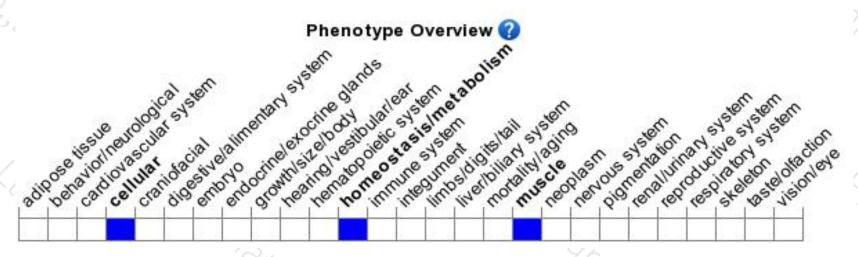
### Protein domain





# 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 metabolic and performance abnormalities in response to mild exercise and improved sprint exercise performance.



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





