

# *Lpar1* Cas9-KO Strategy

**Designer:**

**Lingyan Wu**

**Reviewer:**

**Jiayuan Yao**

**Design Date:**

**2019-7-22**

# Project Overview

**Project Name**

*Lpar1*

**Project type**

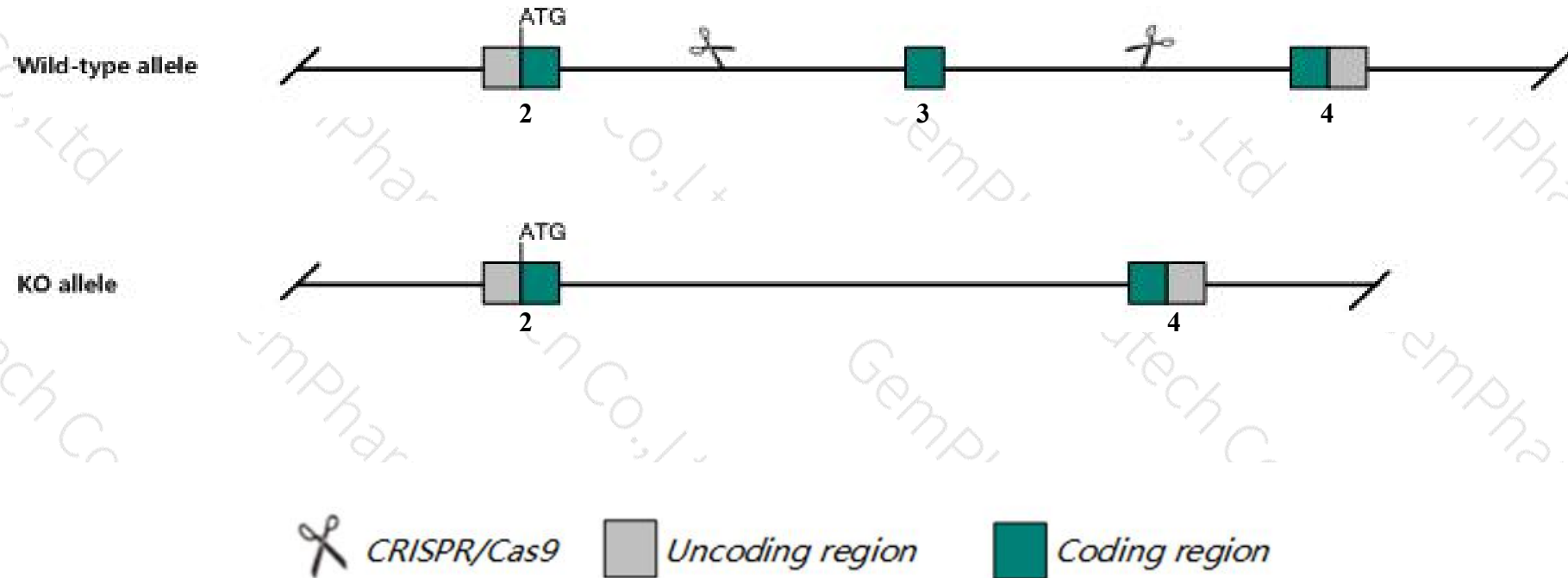
**Cas9-KO**

**Strain background**

**C57BL/6JGpt**

# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Lpar1* gene. The schematic diagram is as follows:



- The *Lpar1* gene has 8 transcripts. According to the structure of *Lpar1* gene, exon3 of *Lpar1*-203 (ENSMUST00000107571.7) transcript is recommended as the knockout region. The region contains 748bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Lpar1* gene. The brief process is as follows: CRISPR/Cas9 system

- According to the existing MGI data, Nullizygous mutations cause partial peri- and postnatal lethality, growth defects, craniofacial anomalies, and wide set eyes. Additional phenotypes include altered brain 5-HT and amino acids, reduced prepulse inhibition, impaired suckling, and increased apoptosis in sciatic nerve Schwann cells.
- The *Lpar1* gene is located on the Chr4. 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)

## Lpar1 lysophosphatidic acid receptor 1 [Mus musculus (house mouse)]

Gene ID: 14745, updated on 31-Jan-2019

### Summary



|                           |                                                                                                                                                                           |
|---------------------------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| <b>Official Symbol</b>    | Lpar1 provided by <a href="#">MGI</a>                                                                                                                                     |
| <b>Official Full Name</b> | lysophosphatidic acid receptor 1 provided by <a href="#">MGI</a>                                                                                                          |
| <b>Primary source</b>     | <a href="#">MGI:MGI:108429</a>                                                                                                                                            |
| <b>See related</b>        | <a href="#">Ensembl:ENSMUSG00000038668</a>                                                                                                                                |
| <b>Gene type</b>          | protein coding                                                                                                                                                            |
| <b>RefSeq status</b>      | VALIDATED                                                                                                                                                                 |
| <b>Organism</b>           | <a href="#">Mus musculus</a>                                                                                                                                              |
| <b>Lineage</b>            | Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus |
| <b>Also known as</b>      | AI326300, Edg2, Gpcr26, Kdt2, IpA1, vzg-1                                                                                                                                 |
| <b>Expression</b>         | Ubiquitous expression in limb E14.5 (RPKM 20.9), colon adult (RPKM 17.1) and 24 other tissues <a href="#">See more</a>                                                    |
| <b>Orthologs</b>          | <a href="#">human</a> <a href="#">all</a>                                                                                                                                 |

# Transcript information (Ensembl)

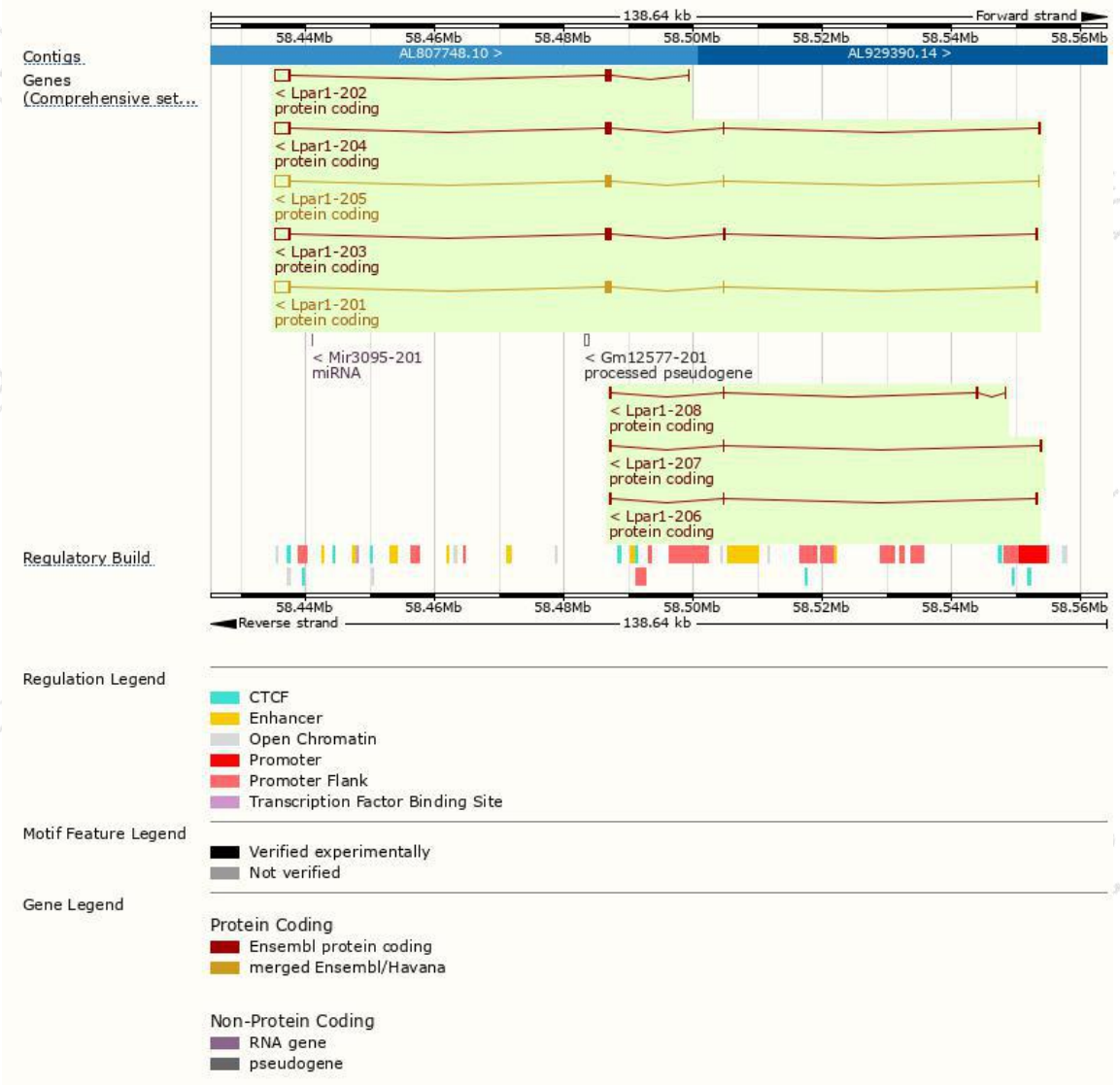
The gene has 8 transcripts,all transcripts are shown below:

| Name      | Transcript ID                         | bp   | Protein               | Biotype        | CCDS                      | UniProt                       | Flags                         |
|-----------|---------------------------------------|------|-----------------------|----------------|---------------------------|-------------------------------|-------------------------------|
| Lpar1-203 | <a href="#">ENSMUST00000107571.7</a>  | 3543 | <a href="#">364aa</a> | Protein coding | <a href="#">CCDS18212</a> | <a href="#">P61793 Q544V2</a> | TSL:5 GENCODE basic APPRIS P1 |
| Lpar1-201 | <a href="#">ENSMUST00000055018.10</a> | 3512 | <a href="#">364aa</a> | Protein coding | <a href="#">CCDS18212</a> | <a href="#">P61793 Q544V2</a> | TSL:1 GENCODE basic APPRIS P1 |
| Lpar1-204 | <a href="#">ENSMUST00000107574.7</a>  | 3437 | <a href="#">364aa</a> | Protein coding | <a href="#">CCDS18212</a> | <a href="#">P61793 Q544V2</a> | TSL:5 GENCODE basic APPRIS P1 |
| Lpar1-205 | <a href="#">ENSMUST00000107575.8</a>  | 3421 | <a href="#">364aa</a> | Protein coding | <a href="#">CCDS18212</a> | <a href="#">P61793 Q544V2</a> | TSL:1 GENCODE basic APPRIS P1 |
| Lpar1-202 | <a href="#">ENSMUST00000107570.1</a>  | 3244 | <a href="#">346aa</a> | Protein coding | <a href="#">CCDS71391</a> | <a href="#">P61793</a>        | TSL:1 GENCODE basic           |
| Lpar1-208 | <a href="#">ENSMUST00000155170.7</a>  | 514  | <a href="#">75aa</a>  | Protein coding | -                         | <a href="#">A2AMI9</a>        | CDS 3' incomplete TSL:5       |
| Lpar1-207 | <a href="#">ENSMUST00000147354.7</a>  | 454  | <a href="#">56aa</a>  | Protein coding | -                         | <a href="#">A2AMJ0</a>        | CDS 3' incomplete TSL:2       |
| Lpar1-206 | <a href="#">ENSMUST00000145361.1</a>  | 335  | <a href="#">45aa</a>  | Protein coding | -                         | <a href="#">A2AMJ1</a>        | CDS 3' incomplete TSL:2       |

The strategy is based on the design of *Lpar1-203* 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, Nullizygous mutations cause partial peri- and postnatal lethality, growth defects, craniofacial anomalies, and wide set eyes. Additional phenotypes include altered brain 5-HT and amino acids, reduced prepulse inhibition, impaired suckling, and increased apoptosis in sciatic nerve Schwann cells.

If you have any questions, you are welcome to inquire.

Tel: 400-9660890

