# *Gorasp1* Cas9-KO Strategy

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**Reviewer: Daohua Xu**

**Design Date: 2021-6-3**

# Project Overview

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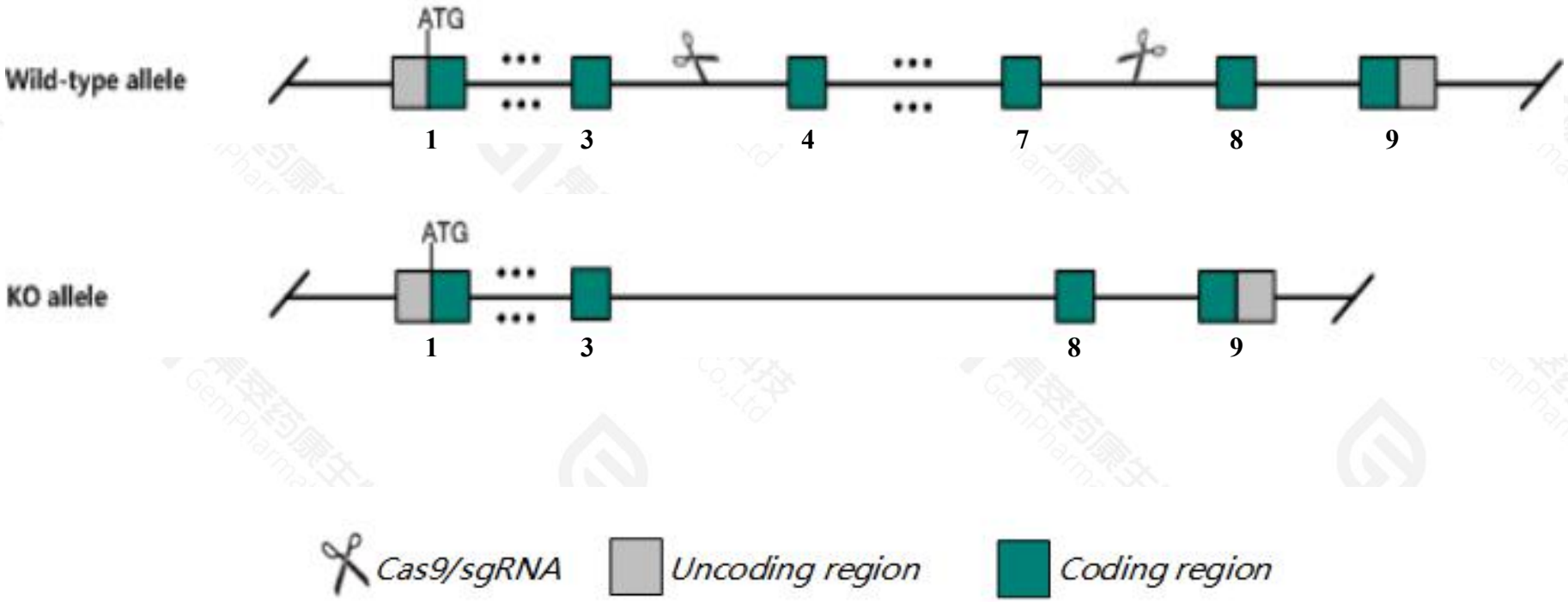
|---|---|
| **Project Name** | ***Gorasp1*** |
| **Project type** | **Cas9-KO** |
| **Strain background** | **C57BL/6JGpt** |

# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Gorasp1* gene. The schematic diagram is as follows:



# Technical routes

➢ The *Gorasp1* gene has 3 transcripts. According to the structure of *Gorasp1* gene, exon4-exon7 of *Gorasp1-201*(ENSMUST00000035099.9) transcript is recommended as the knockout region.The region contains 580bp coding sequence. Knock out the region will result in disruption of protein function.

➢ In this project we use CRISPR/Cas9 technology to modify *Gorasp1* gene. The brief process is as follows: sgRNA was transcribed in vitro.Cas9 and sgRNA 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.

# Notice

➢ According to the existing MGI data,mice homozygous for a targeted disruption of this gene are viable, fertile and healthy with no detectable tissue defects. However, immortalized mutant embryonic fibroblasts show loss of cis Golgi integrity and glycosylation defects.

➢ The partial intron of *Cx3cr1* gene will be deleted together in this strategy.

➢ The N-terminal of *Gorasp1* gene will remain several amino acids,it may remain the partial function of *Gorasp1* gene.

➢ The *Gorasp1* gene is located on the Chr9. 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）

集萃药康 GemPharmatech

## Gorasp1 golgi reassembly stacking protein 1 [Mus musculus (house mouse)]

Gene ID: 74498, updated on 19-Jan-2021

### Summary

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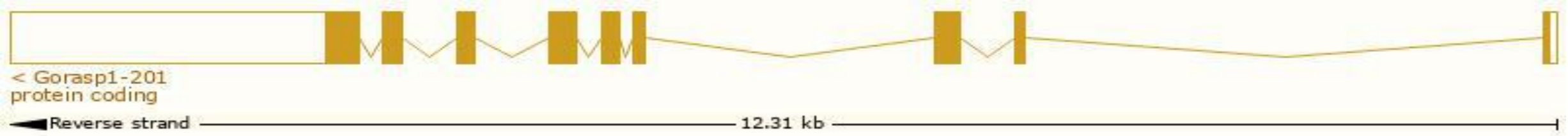
|---|---|
| **Official Symbol** | Gorasp1 provided by MGI |
| **Official Full Name** | golgi reassembly stacking protein 1 provided by MGI |
| **Primary source** | MGI:MGI:1921748 |
| **See related** | Ensembl:ENSMUSG00000032513 |
| **Gene type** | protein coding |
| **RefSeq status** | PROVISIONAL |
| **Organism** | Mus musculus |
| **Lineage** | Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus |
| **Also known as** | 5430411C10Rik, GOL, GOLPH5, GRA, GRASP65, P6, P65 |
| **Expression** | Ubiquitous expression in adrenal adult (RPKM 25.3), genital fat pad adult (RPKM 21.1) and 28 other tissues See more |
| **Orthologs** | human all |

# Transcript information（Ensembl）

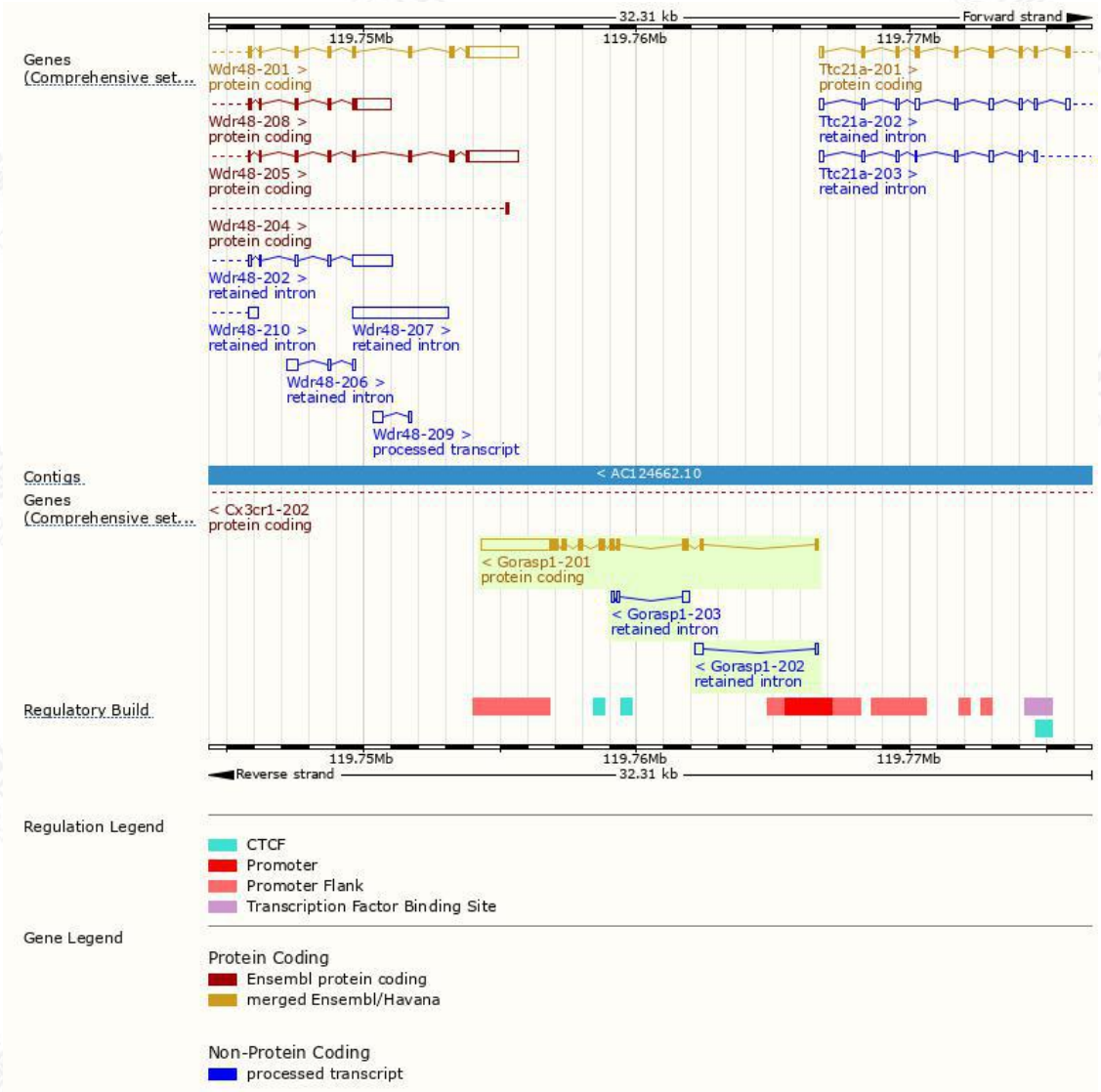
The gene has 3 transcripts,all transcripts are shown below:

| Name | Transcript ID | bp | Protein | Biotype | CCDS | UniProt | Flags |
|---|---|---|---|---|---|---|---|
| **Gorasp1-201** | ENSMUST00000035099.9 | 3906 | 446aa | Protein coding | CCDS23619 | | TSL:1 , GENCODE basic , APPRIS P1 , |
| **Gorasp1-203** | ENSMUST00000214118.2 | 434 | No protein | Retained intron | - | | TSL:2 , |
| **Gorasp1-202** | ENSMUST00000213409.2 | 386 | No protein | Retained intron | - | | TSL:2 , |

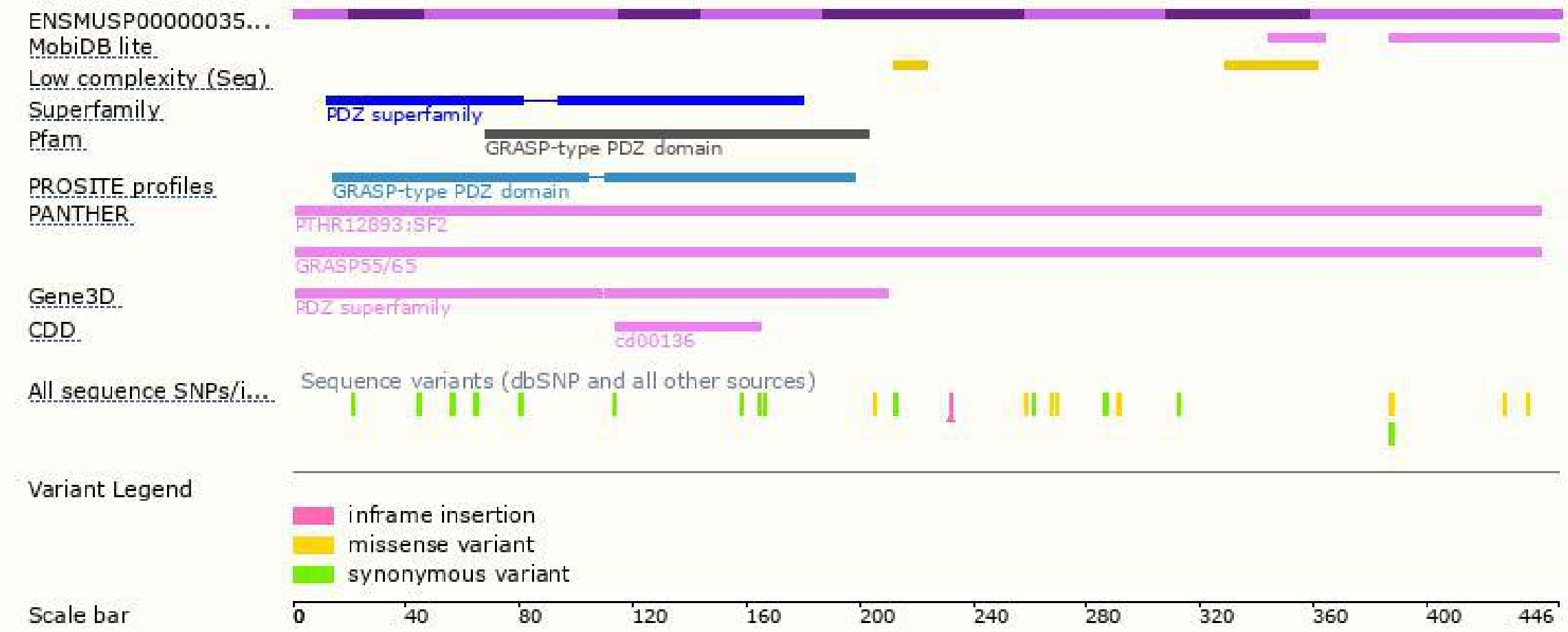
The strategy is based on the design of *Gorasp1-201* transcript,the transcription is shown below:



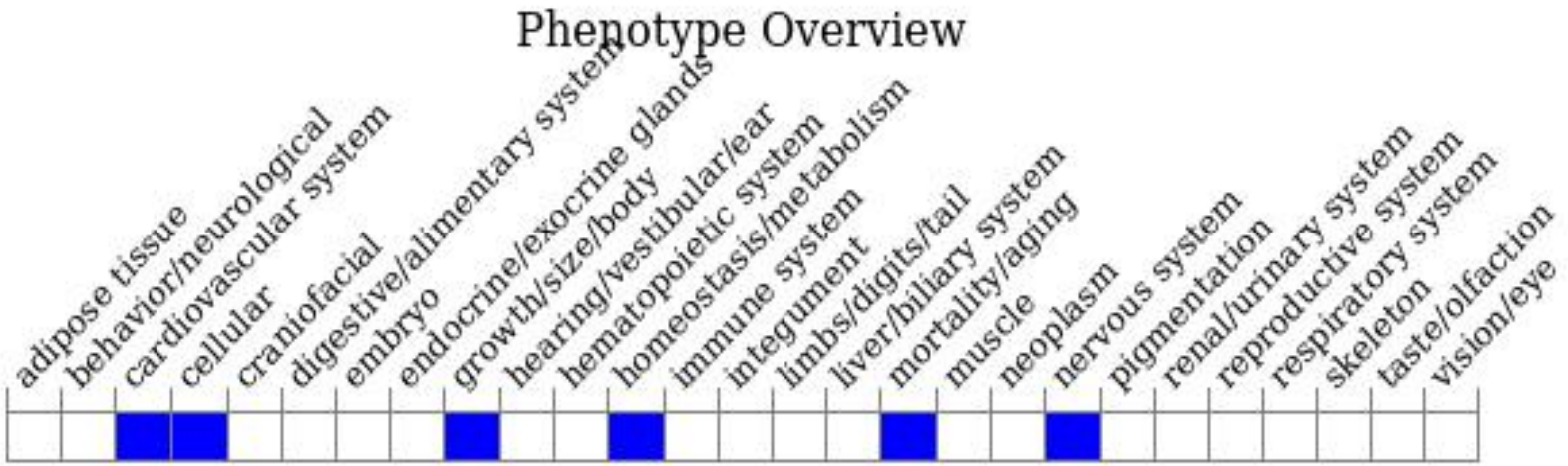
# Genomic location distribution



# 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 targeted disruption of this gene are viable, fertile and healthy with no detectable tissue defects. However, immortalized mutant embryonic fibroblasts show loss of cis Golgi integrity and glycosylation defects.

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

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