

Gpr174 Cas9-CKO Strategy

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Project Overview



Project Name

Gpr174

Project type

Cas9-CKO

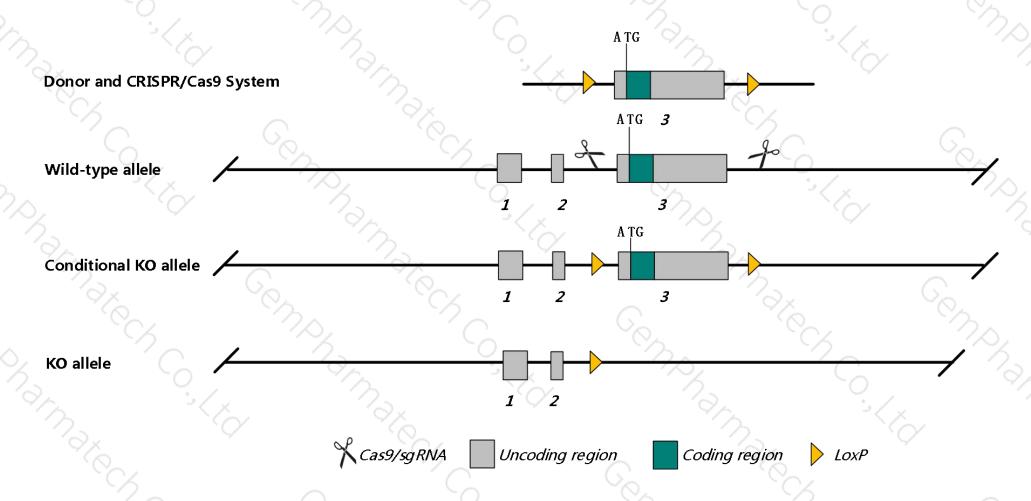
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Gpr174* gene. The schematic diagram is as follows:



Technical routes



- The *Gpr174* gene has 7 transcripts. According to the structure of *Gpr174* gene, exon3 of *Gpr174-207* (ENSMUST00000178838.1) 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 *Gpr174* 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, Male mice hemizygous for a knock-out allele exhibit abnormal T cell proliferation, abnormal regulatory T cell physiology and decreased susceptibility to EAE.
- The *Gpr174* gene is located on the ChrX. 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)



Gpr174 G protein-coupled receptor 174 [Mus musculus (house mouse)]

Gene ID: 213439, updated on 19-Mar-2019

Summary

☆ ?

Official Symbol Gpr174 provided by MGI

Official Full Name G protein-coupled receptor 174 provided by MGI

Primary source MGI:MGI:2685222

See related Ensembl: ENSMUSG00000073008

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 Gm376, Lypsr3

Expression Low expression observed in reference datasetSee more

Orthologs <u>human</u> all

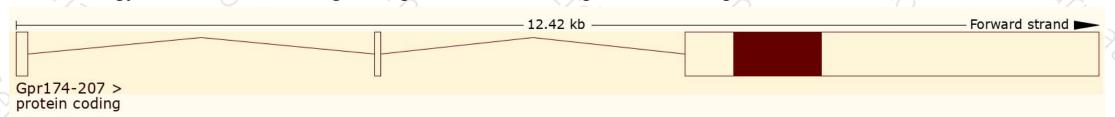
Transcript information (Ensembl)



The gene has 7 transcripts, all transcripts are shown below:

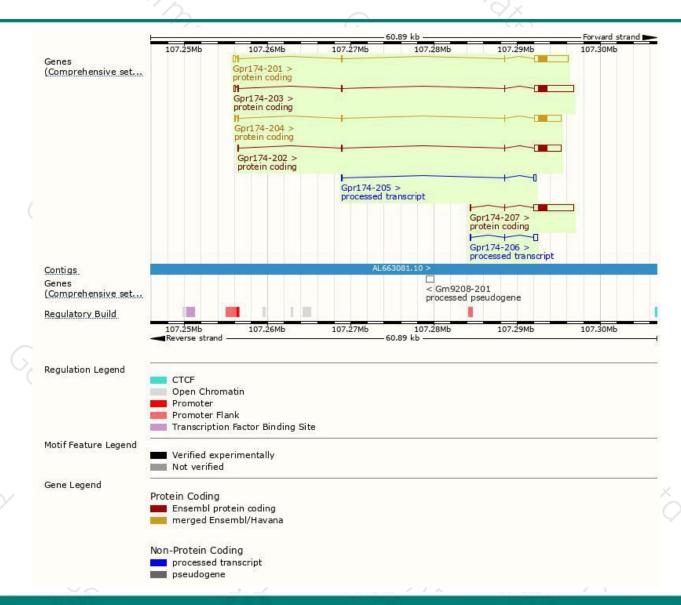
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Gpr174-203	ENSMUST00000118820.7	5112	<u>335aa</u>	Protein coding	CCDS30348	Q3U507	TSL:1 GENCODE basic APPRIS P1
Gpr174-207	ENSMUST00000178838.1	4955	<u>335aa</u>	Protein coding	CCDS30348	Q3U507	TSL:2 GENCODE basic APPRIS P1
Gpr174-201	ENSMUST00000101294.8	4614	<u>335aa</u>	Protein coding	CCDS30348	Q3U507	TSL:1 GENCODE basic APPRIS P1
Gpr174-204	ENSMUST00000120971.7	3588	<u>335aa</u>	Protein coding	CCDS30348	Q3U507	TSL:1 GENCODE basic APPRIS P1
Gpr174-202	ENSMUST00000117310.7	3536	<u>335aa</u>	Protein coding	CCDS30348	Q3U507	TSL:1 GENCODE basic APPRIS P1
Gpr174-206	ENSMUST00000156709.7	639	No protein	Processed transcript	6-80	-	TSL:2
Gpr174-205	ENSMUST00000138031.7	455	No protein	Processed transcript	(2)	0.27	TSL:3

The strategy is based on the design of *Gpr174-207* transcript, The transcription is shown below



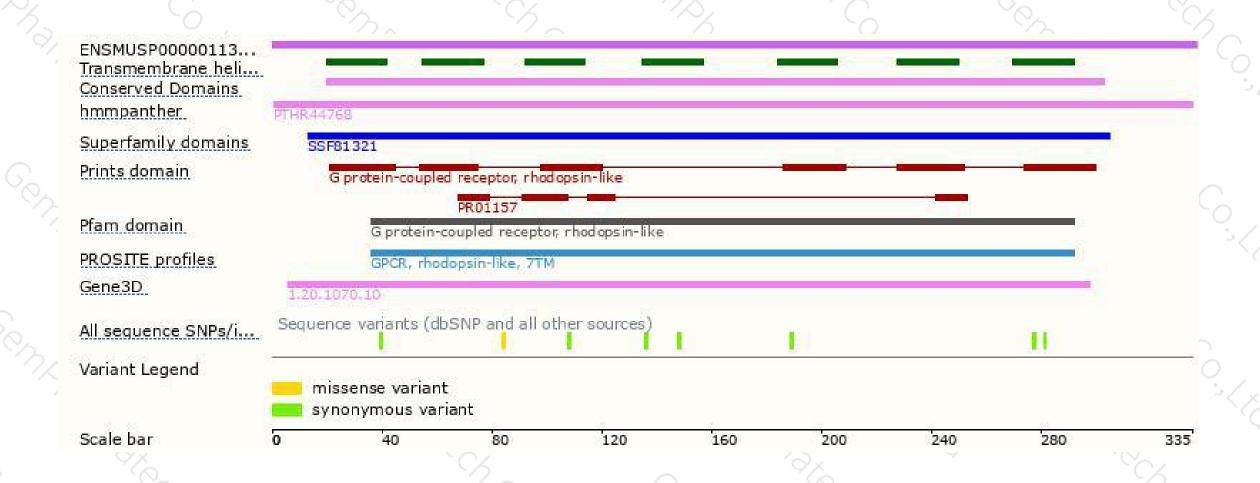
Genomic location distribution





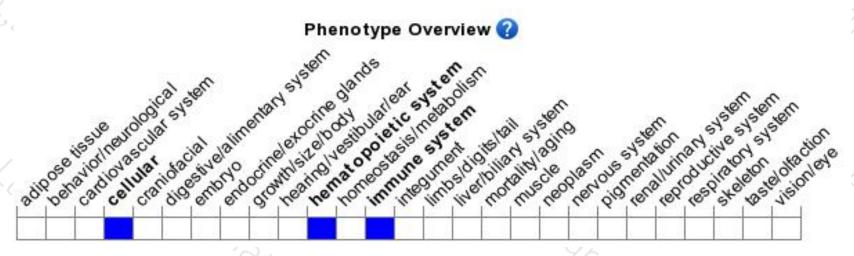
Protein domain





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, Male mice hemizygous for a knock-out allele exhibit abnormal T cell proliferation, abnormal regulatory T cell physiology and decreased susceptibility to EAE.



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





