

P2ry2 Cas9-CKO Strategy

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Reviewer:

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Design Date:

2019-7-25

Project Overview



Project Name

P2ry2

Project type

Cas9-CKO

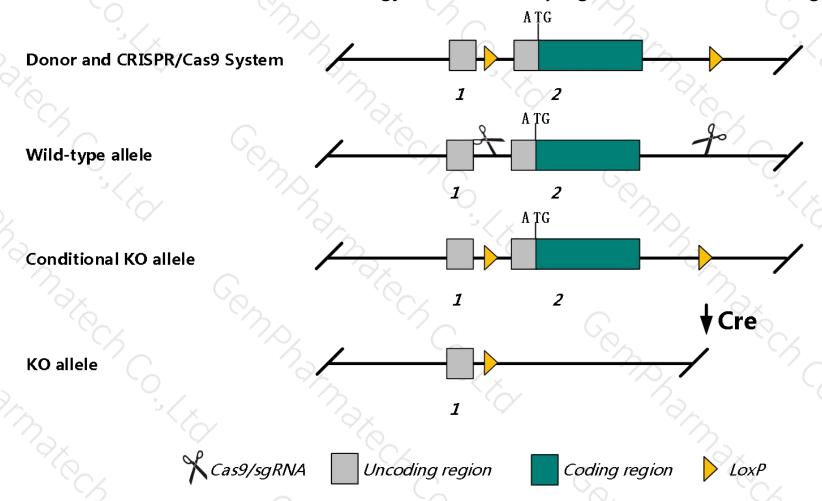
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *P2ry2* gene. The schematic diagram is as follows:



Technical routes



- The *P2ry2* gene has 6 transcripts. According to the structure of *P2ry2* gene, exon2 of *P2ry2-206*(ENSMUST00000208340.1) 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 *P2ry2* 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 disruption in this gene display reduced nucleotide-stimulated calcium secretion from lung fibroblasts and nasal and tracheal epithelial cells and chloride secretion from trachea and gallbladder. Induction of neuronal differentiation ATP gammaS is abolished.
- > The *P2ry2* gene is located on the Chr7. 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)



P2ry2 purinergic receptor P2Y, G-protein coupled 2 [Mus musculus (house mouse)]

Gene ID: 18442, updated on 31-Jan-2019

Summary

☆ ?

Official Symbol P2ry2 provided by MGI

Official Full Name purinergic receptor P2Y, G-protein coupled 2 provided by MGI

Primary source MGI:MGI:105107

See related Ensembl: ENSMUSG00000032860

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 P2U1, P2Y2

Expression Broad expression in heart adult (RPKM 12.3), small intestine adult (RPKM 10.9) and 18 other tissuesSee 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
P2ry2-201	ENSMUST00000037540.4	3011	<u>373aa</u>	Protein coding	CCDS21508	A0A0R4J289	TSL:1 GENCODE basic APPRIS P1
P2ry2-206	ENSMUST00000208340.1	2693	373aa	Protein coding	CCDS21508	A0A0R4J289	TSL:1 GENCODE basic APPRIS P1
P2ry2-204	ENSMUST00000207916.1	2372	<u>373aa</u>	Protein coding	CCDS21508	A0A0R4J289	TSL:1 GENCODE basic APPRIS P1
P2ry2-202	ENSMUST00000178340.2	2137	<u>373aa</u>	Protein coding	CCDS21508	A0A0R4J289	TSL:1 GENCODE basic APPRIS P1
P2ry2-203	ENSMUST00000207049.1	528	<u>21aa</u>	Protein coding	-	A0A140LIR2	CDS 3' incomplete TSL:3
P2ry2-205	ENSMUST00000208258.1	565	No protein	Retained intron		-8	TSL:NA

The strategy is based on the design of *P2ry2-206* transcript, The transcription is shown below

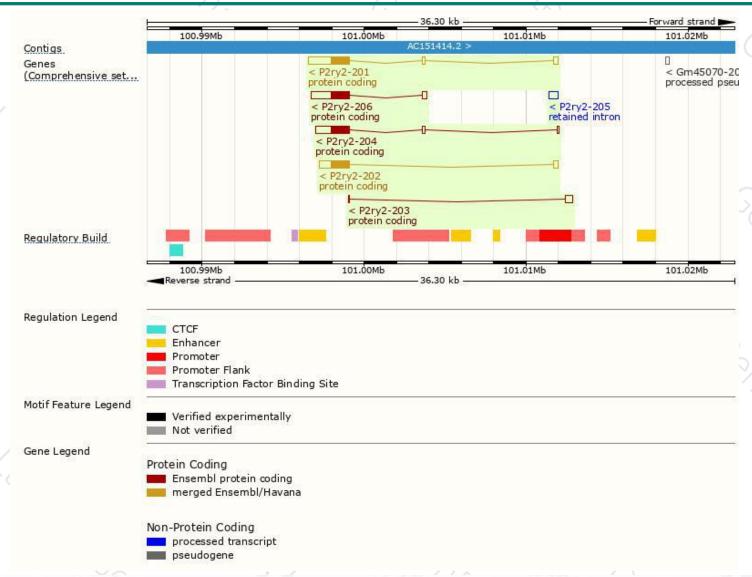
< P2ry2-206
protein coding

Reverse strand

7.16 kb

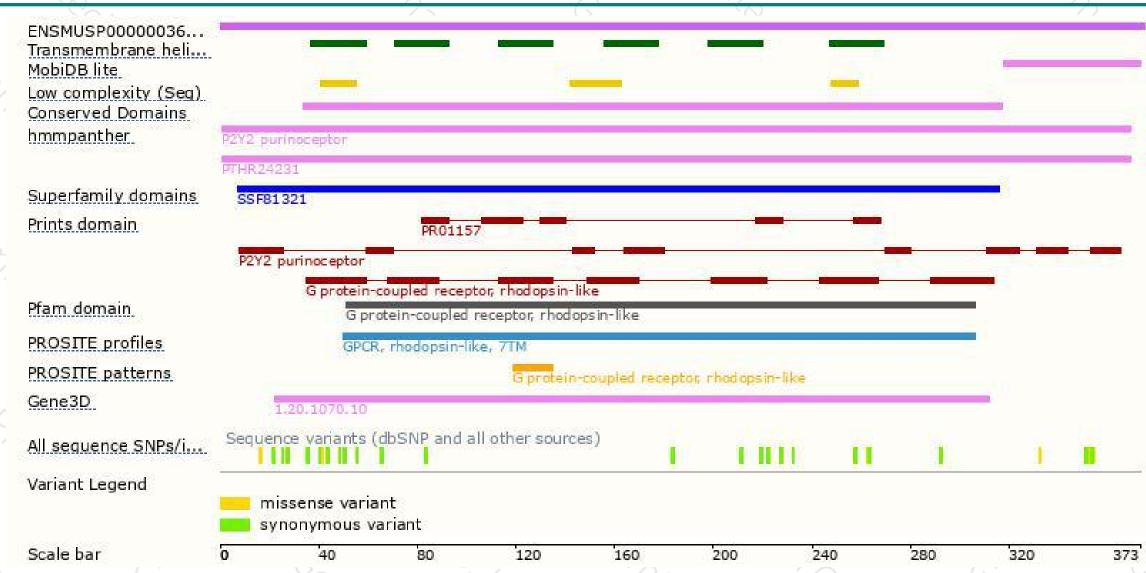
Genomic location distribution





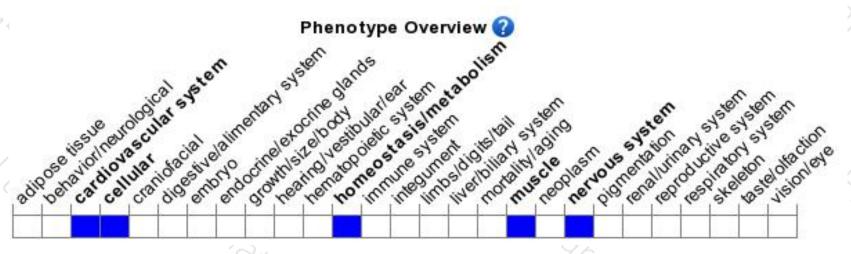
Protein domain





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 disruption in this gene display reduced nucleotide-stimulated calcium secretion from lung fibroblasts and nasal and tracheal epithelial cells and chloride secretion from gallbladder. Induction of neuronal differentiation ATP gammaS is abolished.



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





