

Cyp2e1 Cas9-CKO Strategy To hope of the company of the compan

Designer: Lixin Lv

Project Overview



Project Name

Cyp2e1

Project type

Cas9-CKO

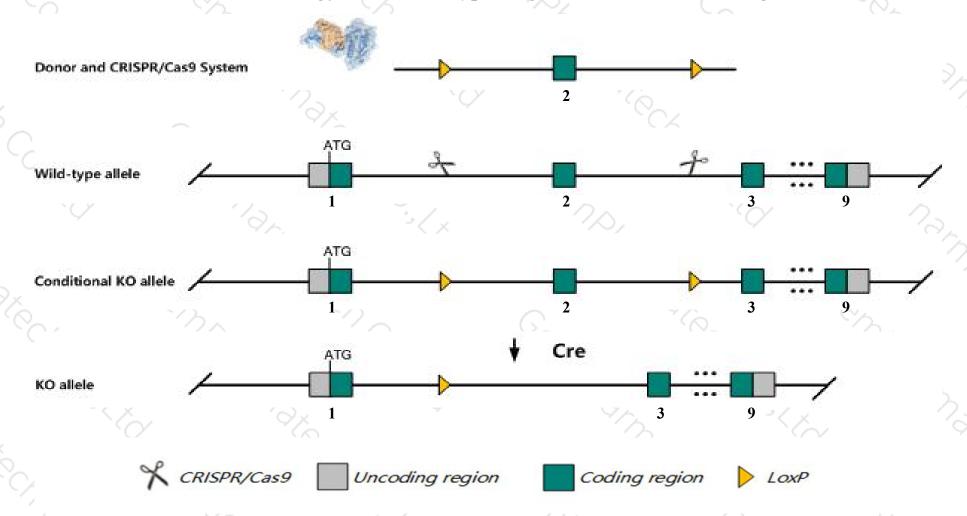
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Cyp2e1 gene. The schematic diagram is as follows:



Technical routes



- The *Cyp2e1* gene has 4 transcripts. According to the structure of *Cyp2e1* gene, exon2 of *Cyp2e1-201* (ENSMUST00000026552.8) transcript is recommended as the knockout region. The region contains 160bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Cyp2e1* 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 exhibit altered responses to xenobiotics including decreased urethane-induced tumors and allylnitrile- or acetamenophen-associated mortality but increased allylnitrile-induced vestibular function loss.
- > The *Cyp2e1* 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)



Cyp2e1 cytochrome P450, family 2, subfamily e, polypeptide 1 [Mus musculus (house mouse)]

Gene ID: 13106, updated on 19-Mar-2019

Summary

☆ ?

Official Symbol Cyp2e1 provided by MGI

Official Full Name cytochrome P450, family 2, subfamily e, polypeptide 1 provided by MGI

Primary source MGI:MGI:88607

See related Ensembl: ENSMUSG00000025479

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 CYPIIE1, Cyp2e

Expression Biased expression in liver adult (RPKM 981.7), subcutaneous fat pad adult (RPKM 122.6) and 2 other tissuesSee more

Orthologs <u>human</u> all

Transcript information (Ensembl)



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

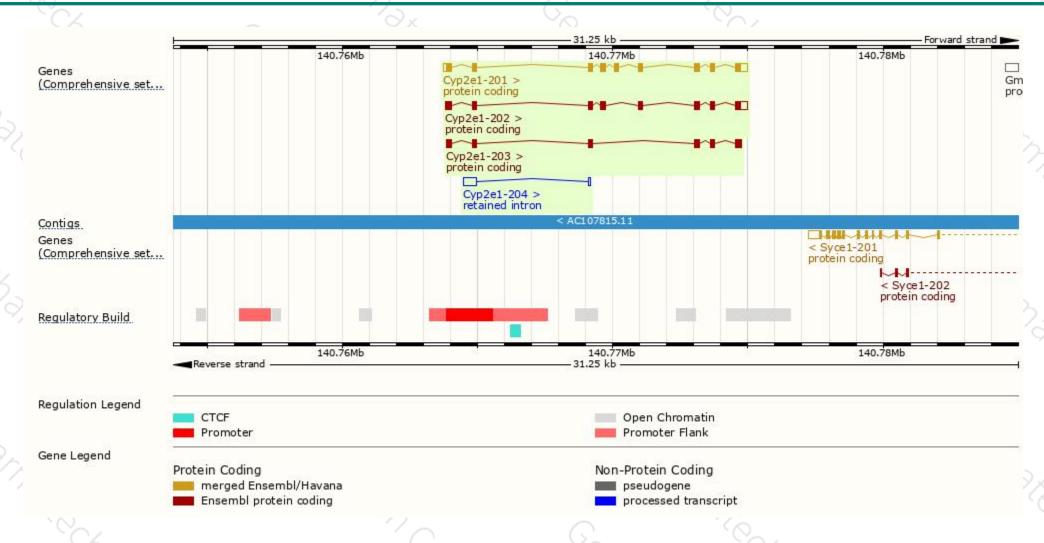
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Cyp2e1-201	ENSMUST00000026552.8	1833	493aa	Protein coding	CCDS21985	Q05421	TSL:1 GENCODE basic APPRIS P1
Cyp2e1-202	ENSMUST00000209253.1	1582	<u>434aa</u>	Protein coding	- 8	A0A1B0GSV7	TSL:5 GENCODE basic
Cyp2e1-203	ENSMUST00000210235.1	1012	<u>333aa</u>	Protein coding	20	Q0PGA1	TSL:1 GENCODE basic
Cyp2e1-204	ENSMUST00000210403.1	569	No protein	Retained intron	29	12	TSL:2

The strategy is based on the design of Cyp2e1-201 transcript, The transcription is shown below



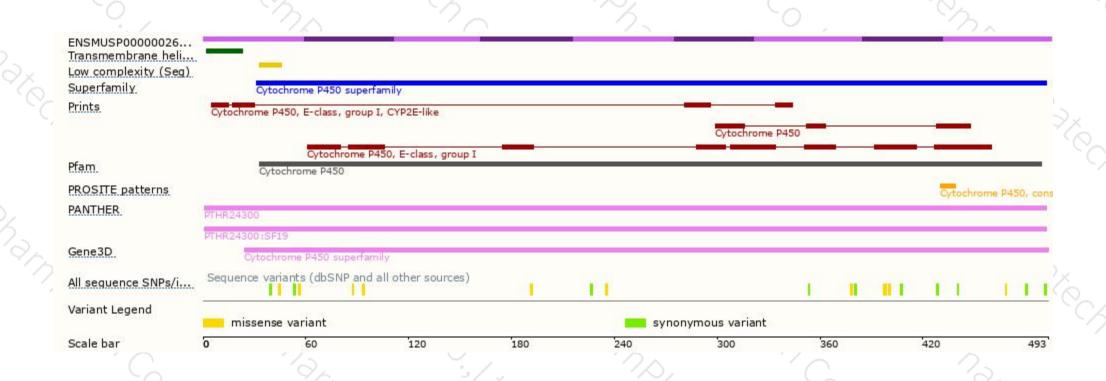
Genomic location distribution





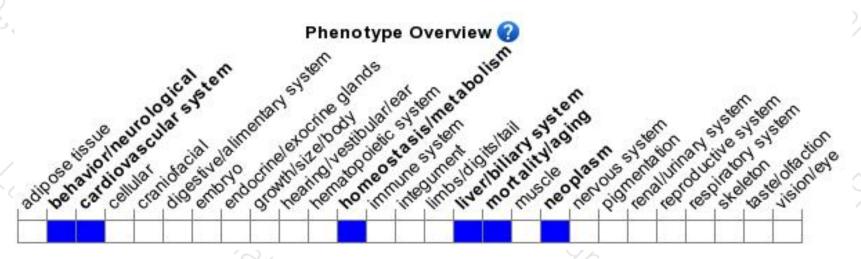
Protein domain





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 null allele exhibit altered responses to xenobiotics including decreased urethane-induced tumors and allylnitrile- or acetamenophen-associated mortality but increased allylnitrile-induced vestibular function loss.



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





