

Abcb6 Cas9-CKO Strategy

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Project Overview



Project Name

Abcb6

Project type

Cas9-CKO

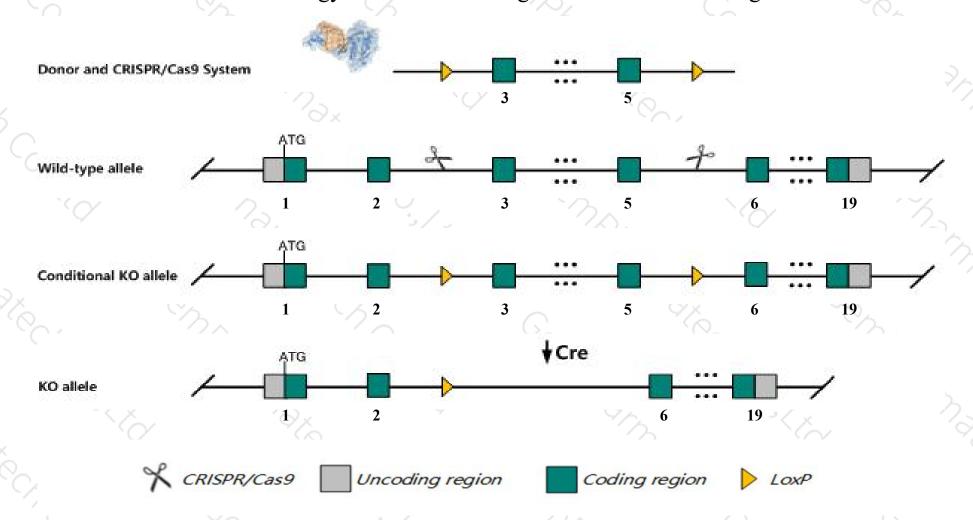
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Abcb6 gene. The schematic diagram is as follows:



Technical routes



- The *Abcb6* gene has 6 transcripts. According to the structure of *Abcb6* gene, exon3-exon5 of *Abcb6-201* (ENSMUST00000027396.14) transcript is recommended as the knockout region. The region contains 467bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Abcb6* 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 mutation display partial lethality, impaired stress erythropoiesis, and absence of ATP-dependent transport of Coproporphyrin III in mitochondria.
- > The N-terminal of Abcb6 gene will remain several amino acids ,it may remain the partial function of Abcb6 gene.
- ➤ The effect on transcript *Abcb6*-206 is unknown.
- > Transcript *Abcb6*-205 may be destroyed dierctly.
- > The *Abcb6* gene is located on the Chr1. 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)



Abcb6 ATP-binding cassette, sub-family B (MDR/TAP), member 6 [Mus musculus (house mouse)]

Gene ID: 74104, updated on 27-Feb-2020

Summary

△ ?

Official Symbol Abcb6 provided by MGI

Official Full Name ATP-binding cassette, sub-family B (MDR/TAP), member 6 provided by MGI

Primary source MGI:MGI:1921354

See related Ensembl:ENSMUSG00000026198

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 1200005B17Rik

Expression Ubiquitous expression in liver adult (RPKM 38.1), colon adult (RPKM 25.7) and 28 other tissues See more

Orthologs human all

▲ Genomic context



Location: 1; 1 C4

See Abcb6 in Genome Data Viewer

Exon count: 19

Annotation release	Status	Assembly	Chr	Location	in d
108	current	GRCm38.p6 (GCF_000001635.26)	1	NC_000067.6 (7517164075180392, complement)	
Build 37.2	previous assembly	MGSCv37 (GCF_000001635.18)	1	NC_000067.5 (7516821475176857, complement)	

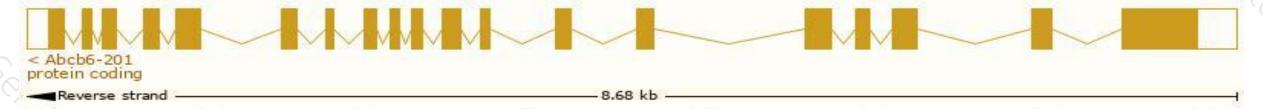
Transcript information (Ensembl)



The gene has 6 transcripts, all transcripts are shown below:

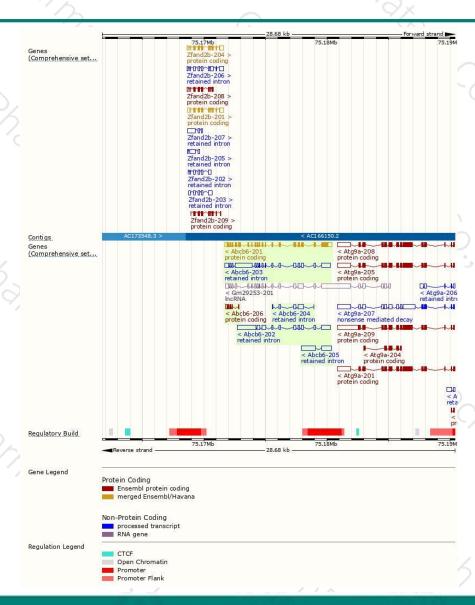
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Abcb6-201	ENSMUST00000027396.14	2966	842aa	Protein coding	CCDS15065	Q9DC29	TSL:1 GENCODE basic APPRIS P1
Abcb6-206	ENSMUST00000161215.1	416	<u>97aa</u>	Protein coding	D-	F6SGP4	CDS 5' incomplete TSL:5
Abcb6-203	ENSMUST00000160081.7	3772	No protein	Retained intron	1/4	ų.	TSL:2
Abcb6-202	ENSMUST00000159219.7	3625	No protein	Retained intron	82	-	TSL:2
Abcb6-205	ENSMUST00000161103.1	1966	No protein	Retained intron	1.5		TSL:1
Abcb6-204	ENSMUST00000160757.1	829	No protein	Retained intron	10-	-	TSL:5

The strategy is based on the design of *Abcb6-201* transcript, The transcription is shown below



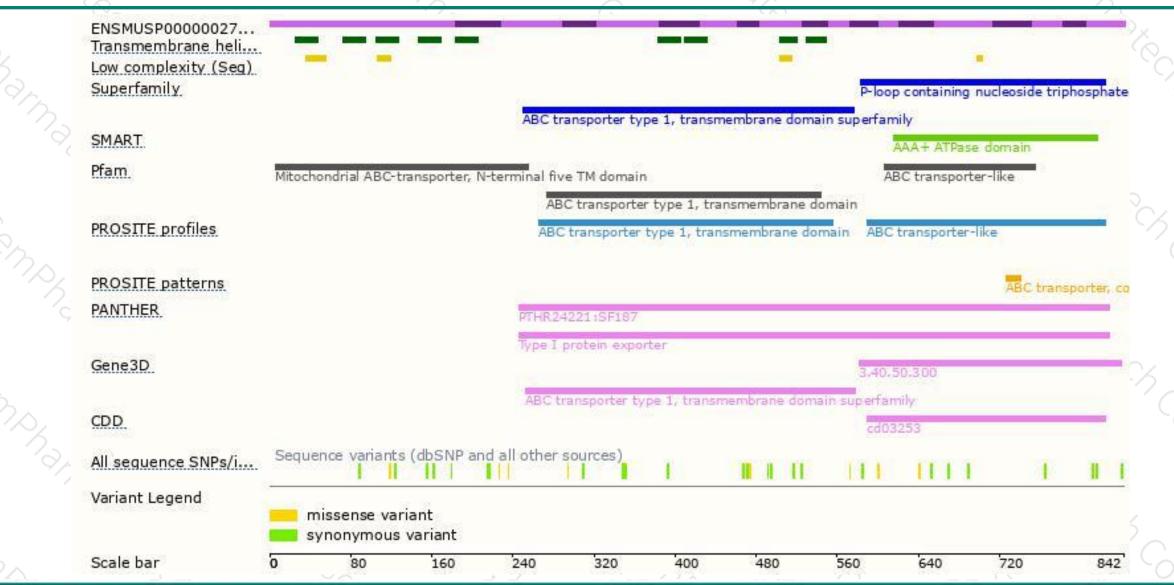
Genomic location distribution





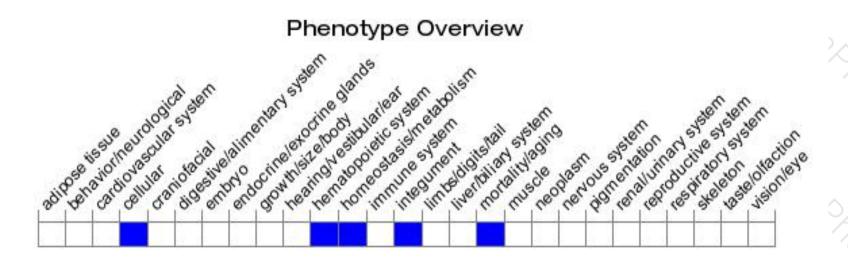
Protein domain





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 mutation display partial lethality, impaired stress erythropoiesis, and absence of ATP-dependent transport of Coproporphyrin III in mitochondria.



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





