

Atp6v0a4 Cas9-KO Strategy

Designer:Qiong Zhou

Project Overview



Project Name

Atp6v0a4

Project type

Cas9-KO

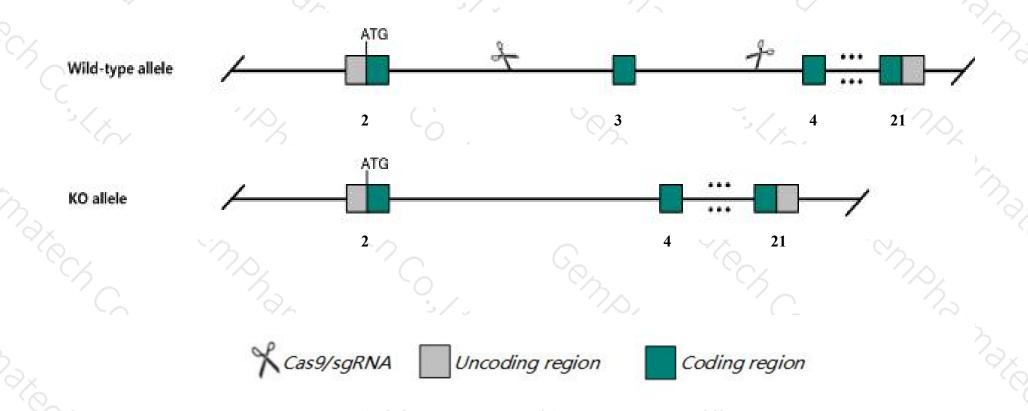
Strain background

C57BL/6J

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Atp6v0a4 gene. The schematic diagram is as follows:



Technical routes



- ➤ The Atp6v0a4 gene has 7 transcripts. According to the structure of Atp6v0a4 gene, exon3 of Atp6v0a4-201 (ENSMUST00000040259.7) transcript is recommended as the knockout region. The region contains 79bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Atp6v0a4* gene. The brief process is as follows: sgRNA was transcribed in vitro.Cas9 and sgRNA were microinjected into the fertilized eggs of C57BL/6J 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/6J mice.

Notice



- ➤ According to the existing MGI data, Mice homozygous for a null mutation display postnatal or premature lethality, hyperchloremic hypokalemic acidosis with hypocitraturia, inner ear defects, impaired hearing, and impaired olfaction.
- ➤ The Atp6v0a4 gene is located on the Chr6. 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)



Atp6v0a4 ATPase, H+ transporting, lysosomal V0 subunit A4 [Mus musculus (house mouse)]

Gene ID: 140494, updated on 31-Jan-2019

Summary

☆ ?

Official Symbol Atp6v0a4 provided by MGI

Official Full Name ATPase, H+ transporting, lysosomal V0 subunit A4 provided by MGI

Primary source MGI:MGI:2153480

See related Ensembl: ENSMUSG00000038600

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 Atp6n1b, a4

Expression Biased expression in kidney adult (RPKM 61.8) and placenta adult (RPKM 4.2)See more

Orthologs <u>human</u> all

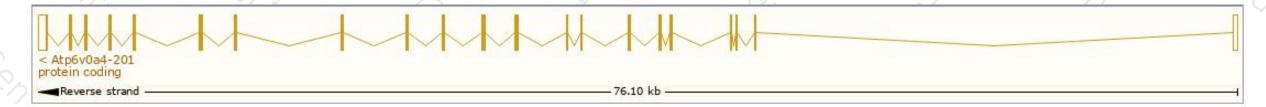
Transcript information (Ensembl)



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

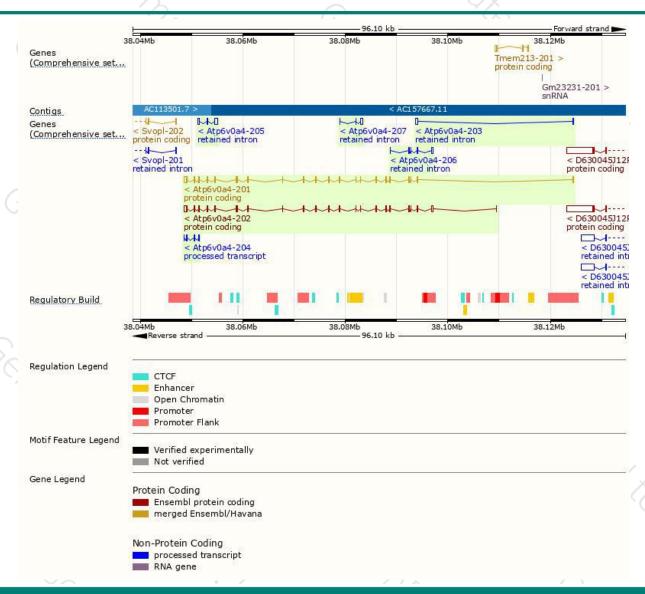
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Atp6v0a4-202	ENSMUST00000114908.4	3307	<u>833aa</u>	Protein coding	CCDS20010	Q920R6	TSL:1 GENCODE basic APPRIS P1
Atp6v0a4-201	ENSMUST00000040259.7	3285	<u>833aa</u>	Protein coding	CCDS20010	Q920R6	TSL:1 GENCODE basic APPRIS P1
Atp6v0a4-204	ENSMUST00000132736.1	508	No protein	Processed transcript	ų.	2	TSL:3
Atp6v0a4-205	ENSMUST00000135594.1	988	No protein	Retained intron	2	2	TSL:3
Atp6v0a4-206	ENSMUST00000138385.1	666	No protein	Retained intron	ā		TSL:5
Atp6v0a4-207	ENSMUST00000144752.1	591	No protein	Retained intron	-		TSL:5
Atp6v0a4-203	ENSMUST00000130816.1	554	No protein	Retained intron	9	2	TSL:2

The strategy is based on the design of Atp6v0a4-201 transcript, The transcription is shown below



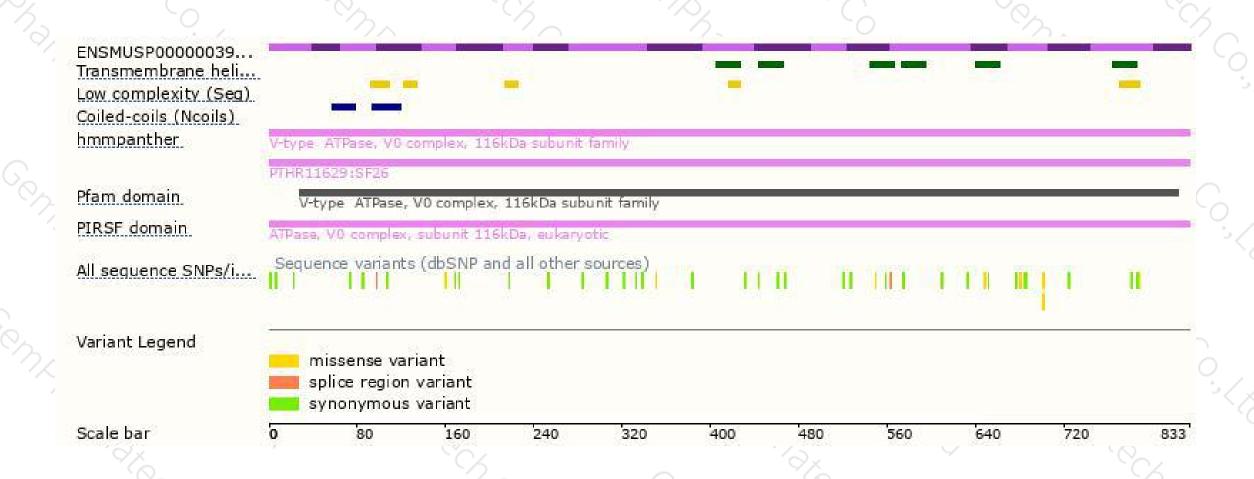
Genomic location distribution





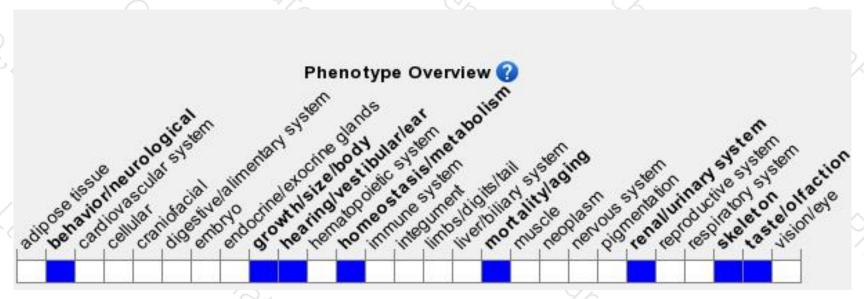
Protein domain





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 postnatal or premature lethality, hyperchloremic hypokalemic acidosis with hypocitraturia, inner ear defects, impaired hearing, and impaired olfaction.



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

Tel: 025-5864 1534





