

# Atp6v1b1 Cas9-CKO Strategy

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**Reviewer:** 

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## **Project Overview**



**Project Name** 

Atp6v1b1

**Project type** 

Cas9-CKO

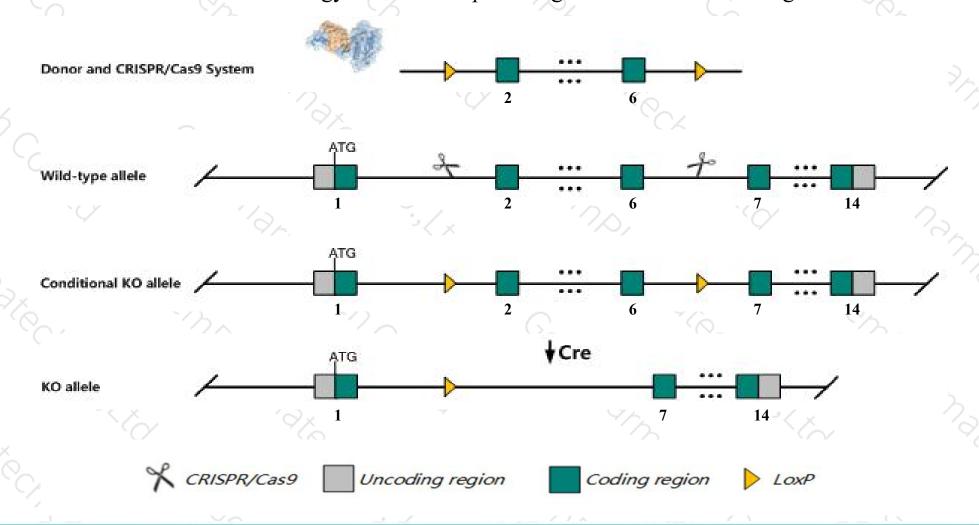
Strain background

C57BL/6JGpt

## Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Atp6v1b1* gene. The schematic diagram is as follows:



### Technical routes



- The *Atp6v1b1* gene has 6 transcripts. According to the structure of *Atp6v1b1* gene, exon2-exon6 of *Atp6v1b1-201* (ENSMUST0000006431.7) 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 *Atp6v1b1* 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 targeted mutation show impaired urinary acidification with a more severe metabolic acidosis and inappropriately alkaline urine after oral acid challenge. However, contrary to expectation, neither hearing nor inner ear morphology are impaired.
- The *Atp6v1b1* 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 loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at existing technological level.

### Gene information (NCBI)



#### Atp6v1b1 ATPase, H+ transporting, lysosomal V1 subunit B1 [ Mus musculus (house mouse) ]

Gene ID: 110935, updated on 19-Nov-2019







Official Symbol Atp6v1b1 provided by MGI

Official Full Name ATPase, H+ transporting, lysosomal V1 subunit B1 provided by MGI

Primary source MGI:MGI:103285

> See related Ensembl:ENSMUSG00000006269

Gene type protein coding RefSeq status VALIDATED Organism Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha;

Muroidea; Muridae; Murinae; Mus; Mus

Vpp3; Vpp-3; Atp6b1; AW208839; D630003L15; D630030L16Rik; D630039P21Rik Also known as

Biased expression in kidney adult (RPKM 52.9), mammary gland adult (RPKM 17.0) and 3 other tissues See more Expression

Orthologs human all

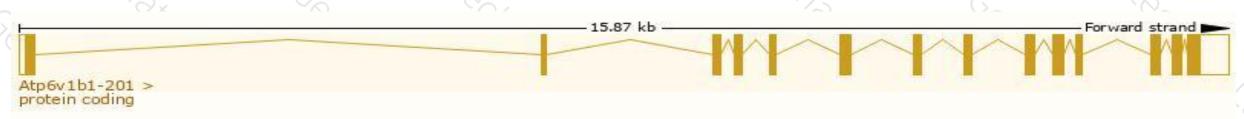
## Transcript information (Ensembl)



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

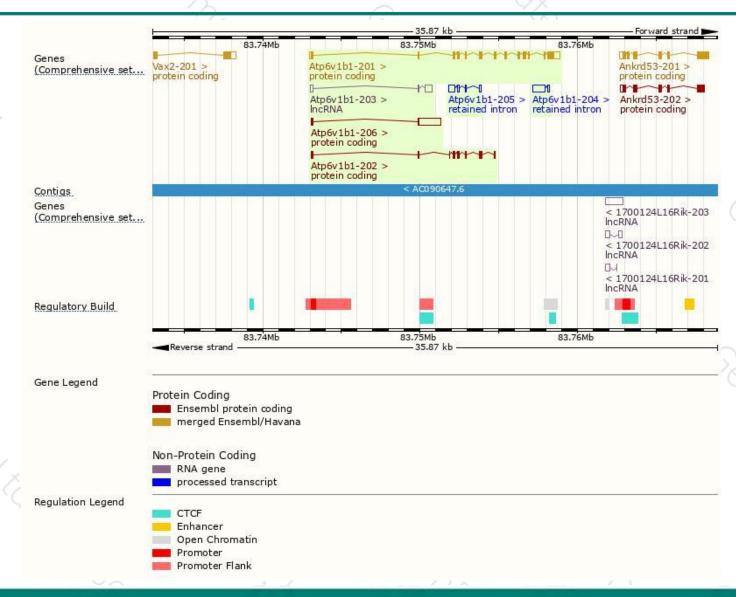
Name 🍦	Transcript ID 🗼	bp 🌲	Protein 🍦	Translation ID 🗼	Biotype 🍦	CCDS 🍦	UniProt 🍦	Flags
Atp6v1b1-201	ENSMUST00000006431.7	1999	<u>513aa</u>	ENSMUSP00000006431.6	Protein coding	CCDS20283₽	Q91YH6@	TSL:1 GENCODE basic APPRIS P1
Atp6v1b1-206	ENSMUST00000206911.1	1591	<u>68aa</u>	ENSMUSP00000146154.1	Protein coding	100	A0A0U1RPW7₽	TSL:1 GENCODE basic
Atp6v1b1-202	ENSMUST00000205763.1	647	215aa	ENSMUSP00000145710.1	Protein coding	32 <del>0</del> 33	A0A0U1RNU9₽	CDS 5' and 3' incomplete TSL:5
Atp6v1b1-204	ENSMUST00000206052.1	935	No protein	-	Retained intron	32 <del>0</del> 3	88	TSL:5
Atp6v1b1-205	ENSMUST00000206652.1	645	No protein	-	Retained intron	1200	( <del>-</del> 6)	TSL:5
Atp6v1b1-203	ENSMUST00000205867.1	653	No protein	-	IncRNA	120	: <del>-</del> ::	TSL:3

The strategy is based on the design of Atp6v1b1-201 transcript, The transcription is shown below



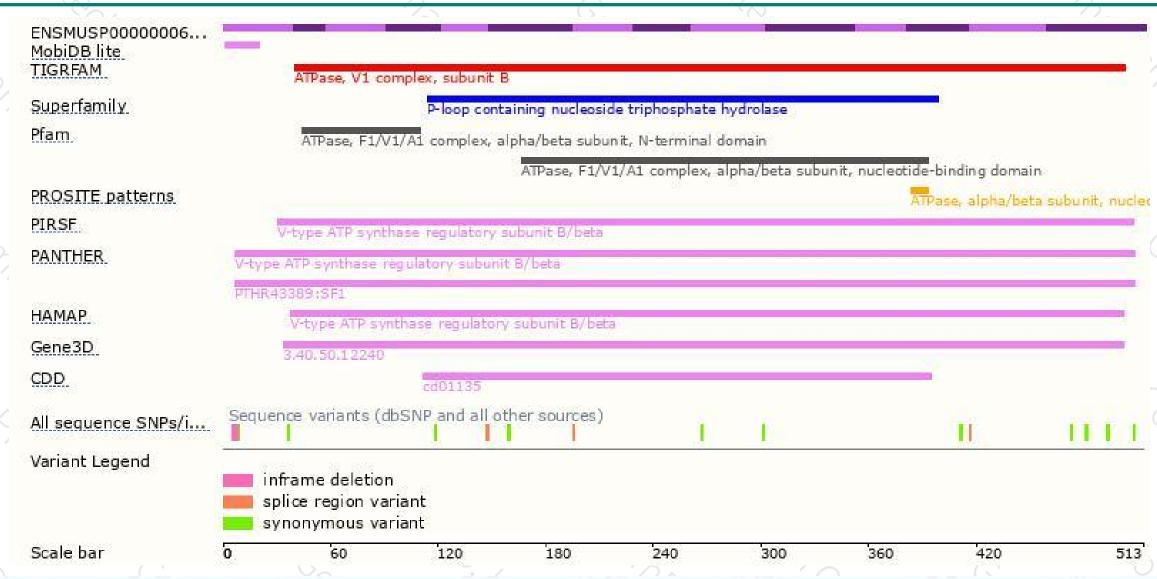
### Genomic location distribution





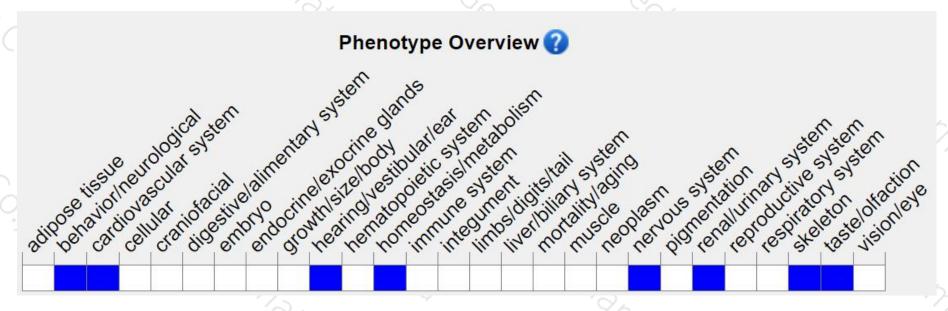
### Protein domain





## 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 targeted mutation show impaired urinary acidification with a more severe metabolic acidosis and inappropriately alkaline urine after oral acid challenge. However, contrary to expectation, neither hearing nor inner ear morphology are impaired.



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





