

Atp6v1h Cas9-KO Strategy

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Project Overview



Project Name Atp6v1h

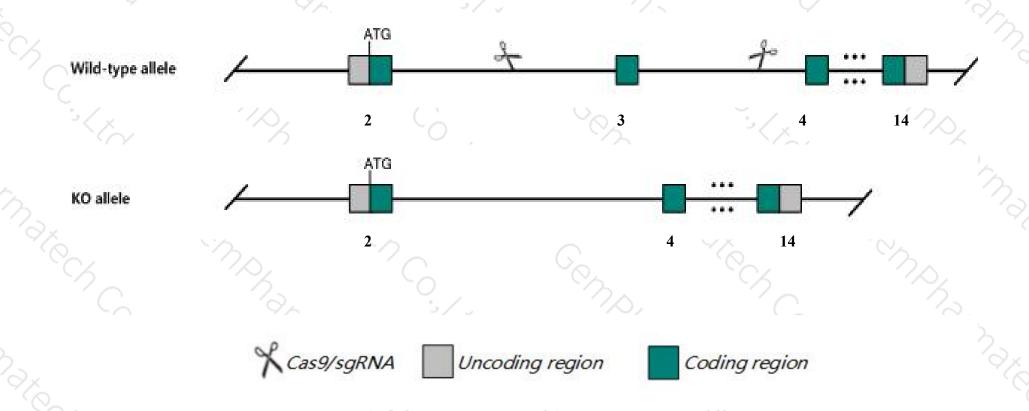
Project type Cas9-KO

Strain background C57BL/6J

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Atp6v1h* gene. The schematic diagram is as follows:



Technical routes



- ➤ The Atp6v1h gene has 8 transcripts. According to the structure of Atp6v1h gene, exon3 of Atp6v1h-201

 (ENSMUST00000044369.12) transcript is recommended as the knockout region. The region contains 103bp coding sequence Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Atp6v1h* 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 knock-out allele exhibit embryonic lethality. Mice heterozygous for the allele exhibit bone loss with altered bone absorption and decreased bone formation.
- The *Atp6v1h* 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 the gene knockout on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.

Gene information (NCBI)



Atp6v1h ATPase, H+ transporting, lysosomal V1 subunit H [Mus musculus (house mouse)]

Gene ID: 108664, updated on 7-Apr-2019

Summary

☆ ?

Official Symbol Atp6v1h provided by MGI

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

Primary source MGI:MGI:1914864

See related Ensembl: ENSMUSG00000033793

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 0710001F19Rik, AU022349, CGI-11, SFD, SFDalpha, SFDbeta, VMA13

Expression Ubiquitous expression in placenta adult (RPKM 22.7), CNS E18 (RPKM 15.8) and 28 other tissuesSee more

Orthologs <u>human</u> all

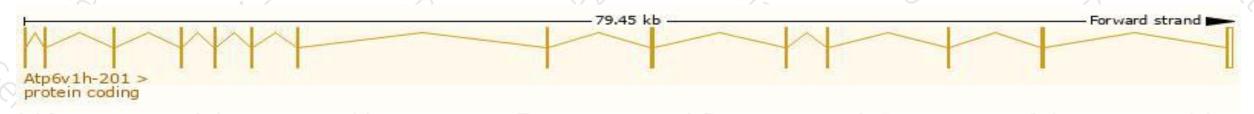
Transcript information (Ensembl)



The gene has 8 transcripts, all transcripts are shown below:

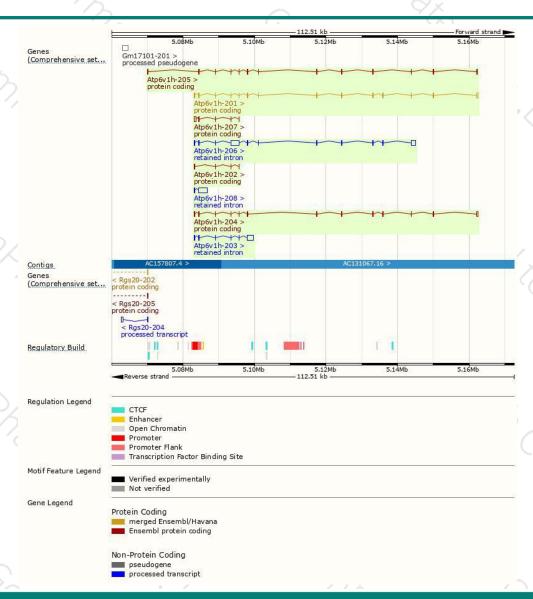
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Atp6v1h-201	ENSMUST00000044369.12	2049	483aa	Protein coding	CCDS14808	Q8BVE3	TSL:1 GENCODE basic APPRIS P3
Atp6v1h-204	ENSMUST00000192698.2	1811	<u>465aa</u>	Protein coding	CCDS78541	A0A0A6YX18	TSL:5 GENCODE basic APPRIS ALT1
Atp6v1h-205	ENSMUST00000192847.5	1662	441aa	Protein coding	2	A0A0A6YWP6	TSL:5 GENCODE basic
Atp6v1h-207	ENSMUST00000194676.5	970	140aa	Protein coding	(c)	A0A0A6YVU0	CDS 3' incomplete TSL:3
Atp6v1h-202	ENSMUST00000192029.5	385	<u>77aa</u>	Protein coding	-	A0A0A6YW86	CDS 3' incomplete TSL:3
Atp6v1h-206	ENSMUST00000194301.5	4829	No protein	Retained intron	- 8		TSL:5
Atp6v1h-208	ENSMUST00000194978.5	2510	No protein	Retained intron	2	-	TSL:1
Atp6v1h-203	ENSMUST00000192142.1	2318	No protein	Retained intron	8	=	TSL:1
			2 2000			7.5 % 3	7 200

The strategy is based on the design of Atp6v1h-201 transcript, The transcription is shown below



Genomic location distribution





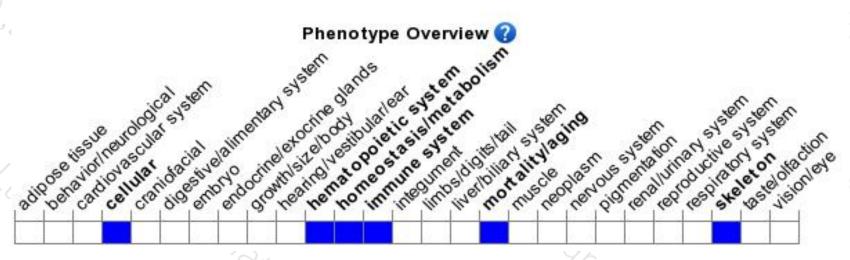
Protein domain





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 knock-out allele exhibit embryonic lethality. Mice heterozygous for the allele exhibit bone loss with altered bone absorption and decreased bone formation.



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

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