

Ush2a Cas9-KO Strategy

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Project Overview



Project Name

Ush2a

Project type

Cas9-KO

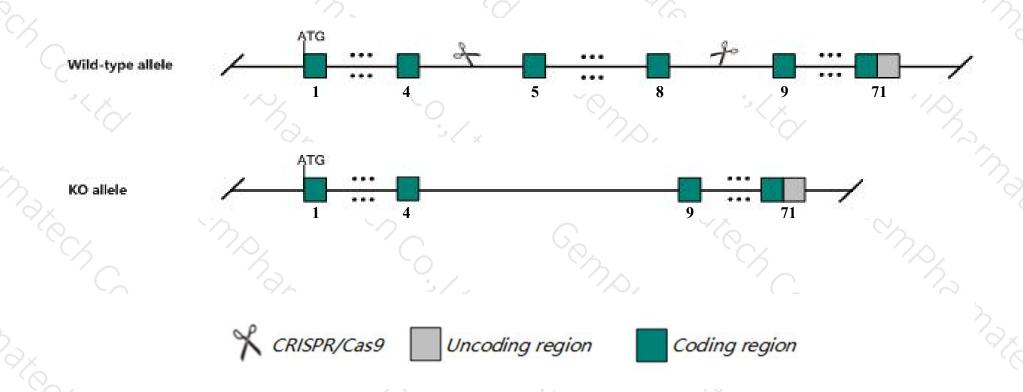
Strain background

C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Ush2a* gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Ush2a* gene has 5 transcripts. According to the structure of *Ush2a* gene, exon5-exon8 of *Ush2a-201* (ENSMUST00000060479.13) transcript is recommended as the knockout region. The region contains 796bp coding sequence Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Ush2a* gene. The brief process is as follows: CRISPR/Cas9 system

Notice



- ➤ According to the existing MGI data, Mice homozygous for a knock-out allele display progressive retinal photoreceptor degeneration along with significantly reduced a- and b-wave amplitudes, and a moderate but nonprogressive high-frequency hearing loss associated with widespread loss of outer hair cells in the basal turn of the cochlea.
- The *Ush2a* 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)



Ush2a usherin [Mus musculus (house mouse)]

Gene ID: 22283, updated on 10-Oct-2019

Summary

Official Symbol Ush2a provided by MGI

Official Full Name usherin provided by MGI

Primary source MGI:MGI:1341292

See related Ensembl: ENSMUSG00000026609

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 Gm676; Gm983; Ushrn; Mush2a; A930011D15Rik; A930037M10Rik

Expression Biased expression in testis adult (RPKM 1.3) and ovary adult (RPKM 0.0) See more

Orthologs human all

Transcript information (Ensembl)



The gene has 5 transcripts, all transcripts are shown below:

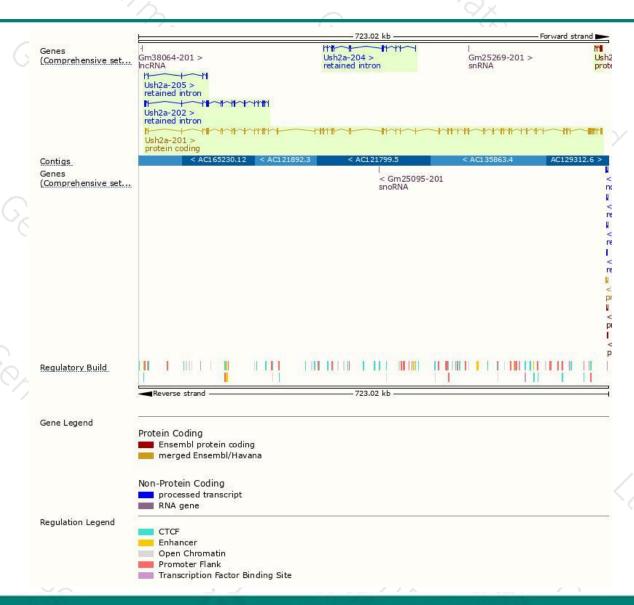
Name	Transcript ID	bp	Protein	Biotype	ccds	UniProt	Flags
Ush2a-201	ENSMUST00000060479.13	15695	5193aa	Protein coding	CCDS15607	Q2Q147	TSL:1 GENCODE basic APPRIS P1
Ush2a-203	ENSMUST00000127077.1	750	<u>237aa</u>	Protein coding	-	F6TQ19	CDS 5' incomplete TSL:5
Ush2a-202	ENSMUST00000124358.1	5126	No protein	Retained intron		1920	TSL:1
Ush2a-204	ENSMUST00000142159.1	2282	No protein	Retained intron	-	323	TSL:1
Ush2a-205	ENSMUST00000142189.7	1236	No protein	Retained intron	5	(5)	TSL:1

The strategy is based on the design of *Ush2a-201* transcript, The transcription is shown below



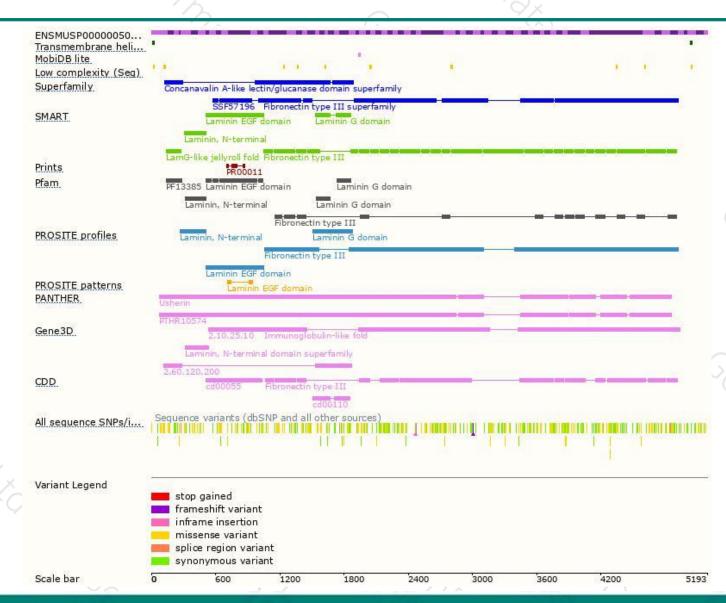
Genomic location distribution





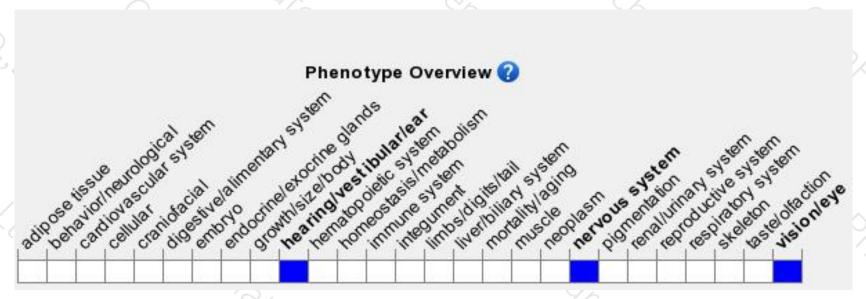
Protein domain





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 display progressive retinal photoreceptor degeneration along with significantly reduced a- and b-wave amplitudes, and a moderate but nonprogressive high-frequency hearing loss associated with widespread loss of outer hair cells in the basal turn of the cochlea.



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





