

Usp2 Cas9-KO Strategy

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Project Overview



Project Name

Usp2

Project type

Cas9-KO

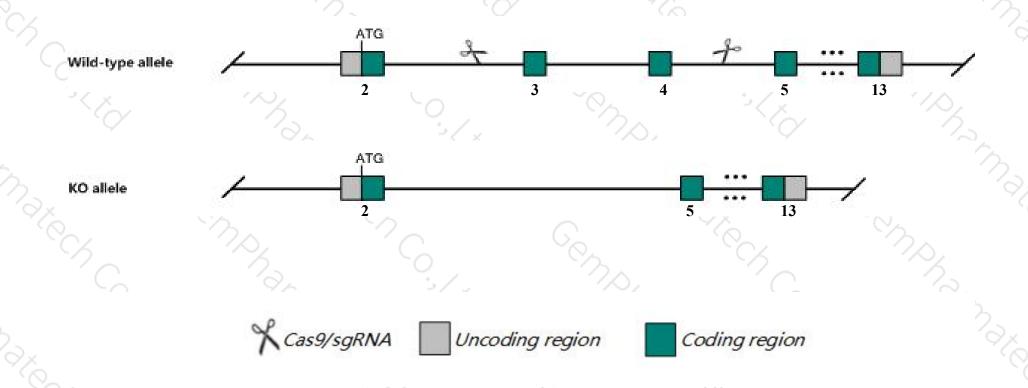
Strain background

C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Usp2* gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Usp2* gene has 9 transcripts. According to the structure of *Usp2* gene, exon3-exon4 of *Usp2-201*(ENSMUST00000034508.13) transcript is recommended as the knockout region. The region contains 175bp coding sequence Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Usp2* gene. The brief process is as follows: sgRNA was transcribed in vitro.Cas9 and sgRNA 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.

Notice



- ➤ According to the existing MGI data, Mice homozygous for a null mutation display severely reduced male fertility with defects in sperm motility.
- ➤ Transcript *Usp2*-205&209 may not be affected.
- \rightarrow The partial intron of Gm49380 gene will be deleted together in this strategy.
- > The *Usp2* gene is located on the Chr9. 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)



Usp2 ubiquitin specific peptidase 2 [Mus musculus (house mouse)]

Gene ID: 53376, updated on 31-Jan-2019

Summary

☆ ?

Official Symbol Usp2 provided by MGI

Official Full Name ubiquitin specific peptidase 2 provided by MGI

Primary source MGI:MGI:1858178

See related Ensembl:ENSMUSG00000032010

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 B930035K21Rik, Ubp41

Expression Biased expression in testis adult (RPKM 55.3), heart adult (RPKM 6.4) and 5 other tissuesSee more

Orthologs <u>human all</u>

Transcript information (Ensembl)



The gene has 9 transcripts, all transcripts are shown below:

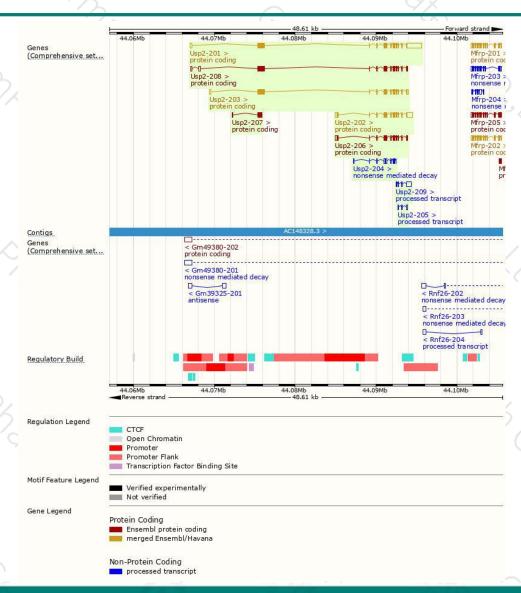
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Usp2-201	ENSMUST00000034508.13	3867	619aa	Protein coding	CCDS23094	088623	TSL:1 GENCODE basic
Usp2-208	ENSMUST00000177054.7	2370	<u>619aa</u>	Protein coding	CCDS23094	<u>088623</u>	TSL:1 GENCODE basic
Usp2-203	ENSMUST00000114830.8	2122	<u>619aa</u>	Protein coding	CCDS23094	<u>088623</u>	TSL:1 GENCODE basic
Usp2-202	ENSMUST00000065461.8	2057	396aa	Protein coding	CCDS23095	<u>088623</u>	TSL:1 GENCODE basic APPRIS P2
Usp2-206	ENSMUST00000176416.7	1486	<u>393aa</u>	Protein coding	5	<u>088623</u>	TSL:1 GENCODE basic APPRIS ALT1
Usp2-207	ENSMUST00000176671.1	659	<u>174aa</u>	Protein coding	*	H3BLN9	CDS 3' incomplete TSL:3
Usp2-204	ENSMUST00000175816.1	748	<u>43aa</u>	Nonsense mediated decay	-	H3BLH3	TSL:3
Usp2-209	ENSMUST00000177422.7	908	No protein	Processed transcript	24	82	TSL:2
Usp2-205	ENSMUST00000176022.1	417	No protein	Processed transcript	-	65	TSL:2
					N. 18 1800		Y

The strategy is based on the design of *Usp2-201* transcript, The transcription is shown below



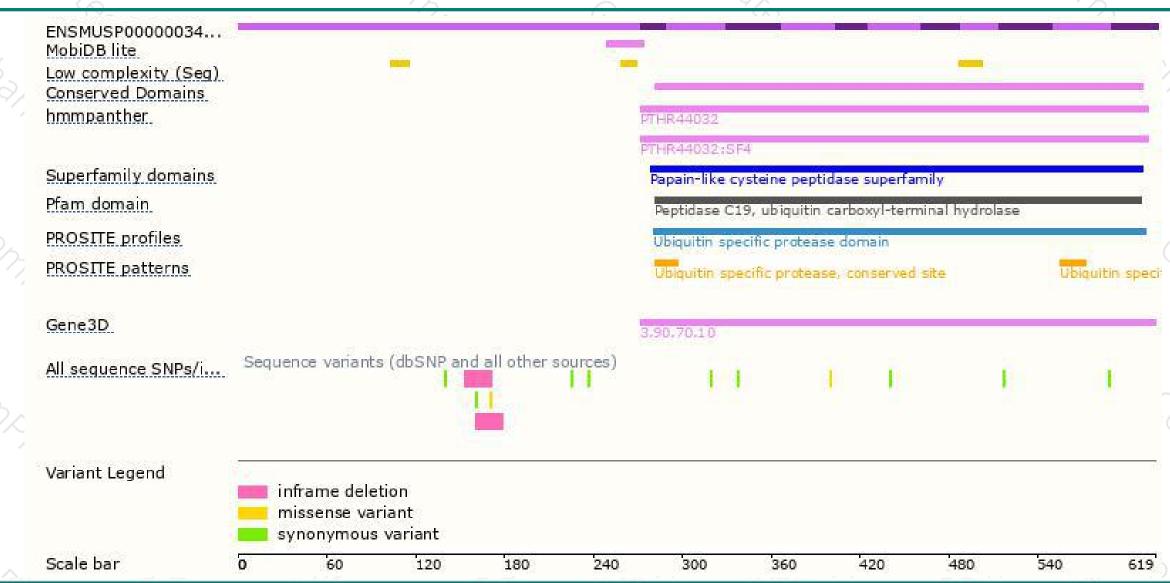
Genomic location distribution





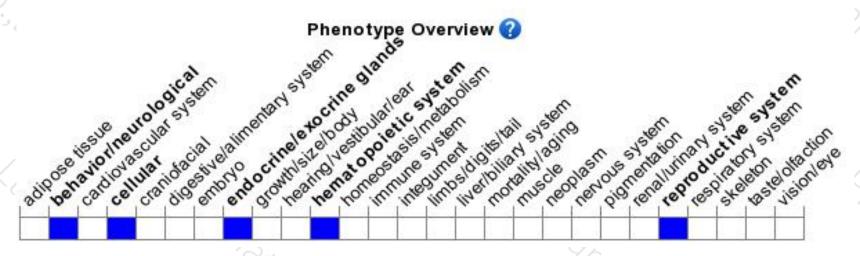
Protein domain





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 severely reduced male fertility with defects in sperm motility.



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

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