

Wasf1 Cas9-KO Strategy

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Reviewer: Yanhua Shen

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Project Overview



Project Name Wasf1

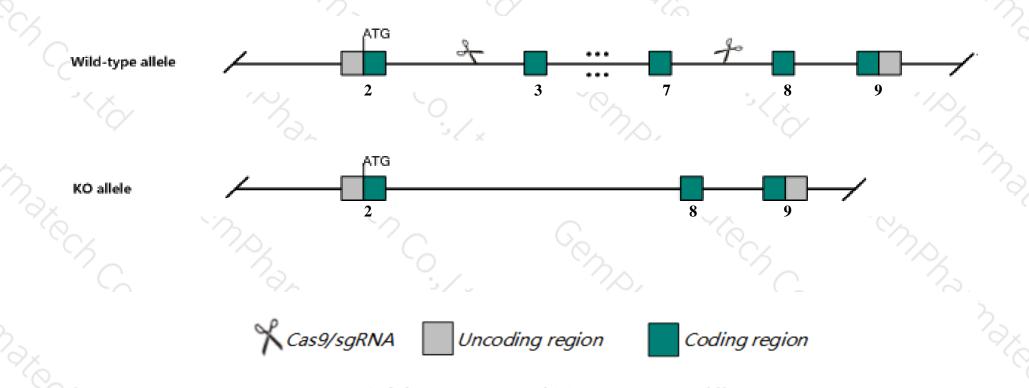
Project type Cas9-KO

Strain background C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Wasf1* gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Wasf1* gene has 2 transcripts. According to the structure of *Wasf1* gene, exon3-exon7 of *Wasf1-202*(ENSMUST00000105509.1) transcript is recommended as the knockout region. The region contains 760bp coding sequence.

 Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Wasf1* 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



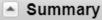
- ➤ According to the existing MGI data, Mutation of this gene has been associated with both morphological and functional defects of the central nervous system. Targeted mutagenesis has resulted in mice that display sensorimotor and cognitive defects similar to those exhibited by patients with 3p-syndrome mental retardation.
- ➤ The *Wasf1* gene is located on the Chr10. 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)



Wasf1 WAS protein family, member 1 [Mus musculus (house mouse)]

Gene ID: 83767, updated on 22-Oct-2019



? 💠

Official Symbol Wasf1 provided by MGI

Official Full Name WAS protein family, member 1 provided by MGI

Primary source MGI:MGI:1890563

See related Ensembl: ENSMUSG00000019831

Gene type protein coding
RefSeq status VALIDATED
Organism Mus musculus

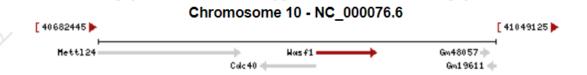
Lineage Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha;

Muroidea; Muridae; Murinae; Mus; Mus

Also known as Scar; WAVE; WAVE-1; Al195380; Al838537

Expression Biased expression in frontal lobe adult (RPKM 33.5), cortex adult (RPKM 31.9) and 9 other tissues See more

Orthologs <u>human</u> all



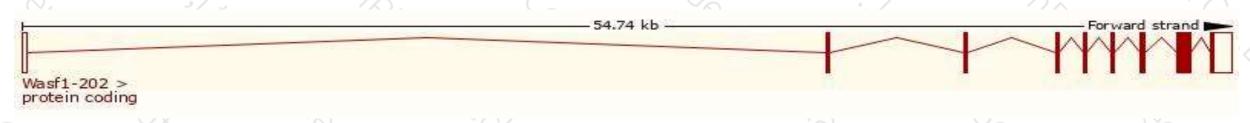
Transcript information (Ensembl)



The gene has 2 transcripts, all transcripts are shown below:

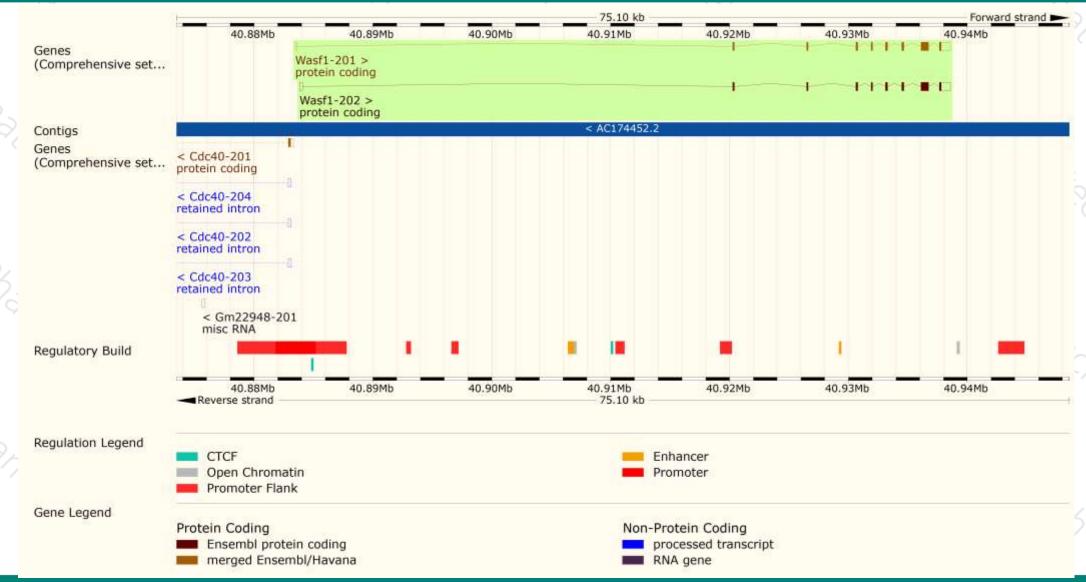
Name 🍦	Transcript ID	bp 🌲	Protein 🌲	Translation ID	Biotype 🍦	CCDS	UniProt 🍦	Flags		
Wasf1-202	ENSMUST00000105509.1	2719	<u>559aa</u>	ENSMUSP00000101148.1	Protein coding	CCDS23801 ₺	<u>Q8R5H6</u> ₽	TSL:5	GENCODE basic	APPRIS P1
Wasf1-201	ENSMUST00000019975.13	2609	<u>559aa</u>	ENSMUSP00000019975.7	Protein coding	CCDS23801 ₺	<u>Q8R5H6</u> ₽	TSL:1	GENCODE basic	APPRIS P1

The strategy is based on the design of Wasf1-202 transcript, The transcription is shown below



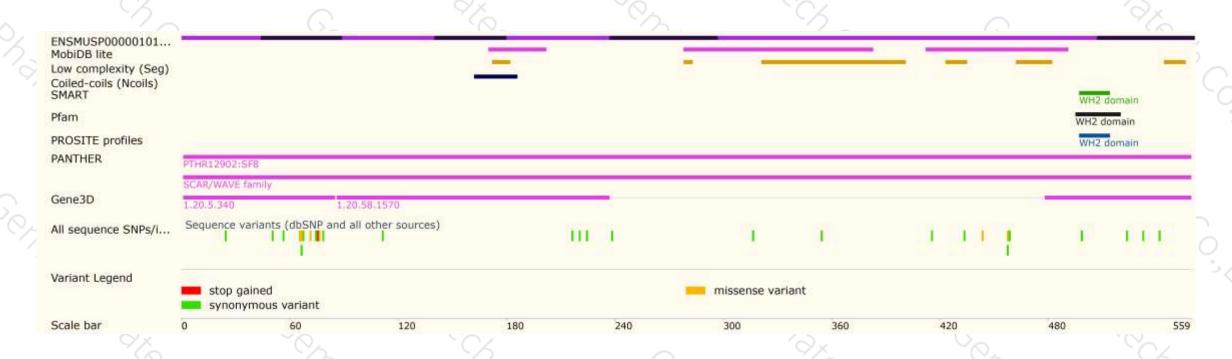
Genomic location distribution





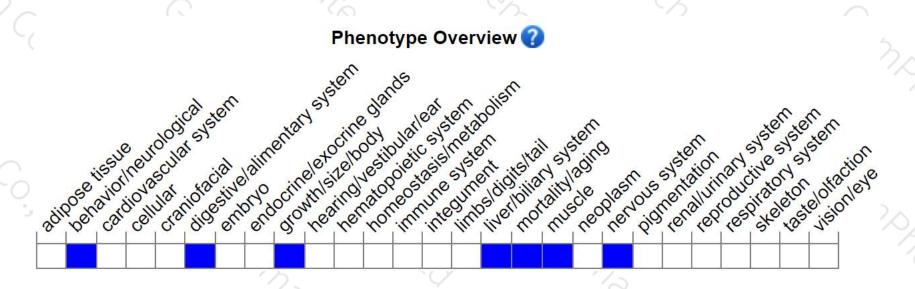
Protein domain





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, Mutation of this gene has been associated with both morphological and functional defects of the central nervous system. Targeted mutagenesis has resulted in mice that display sensorimotor and cognitive defects similar to those exhibited by patients with 3p-syndrome mental retardation.



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

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