

Hsph1 Cas9-CKO Strategy

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Design Date: 2019-11-26

Project Overview



Project Name Hsph1

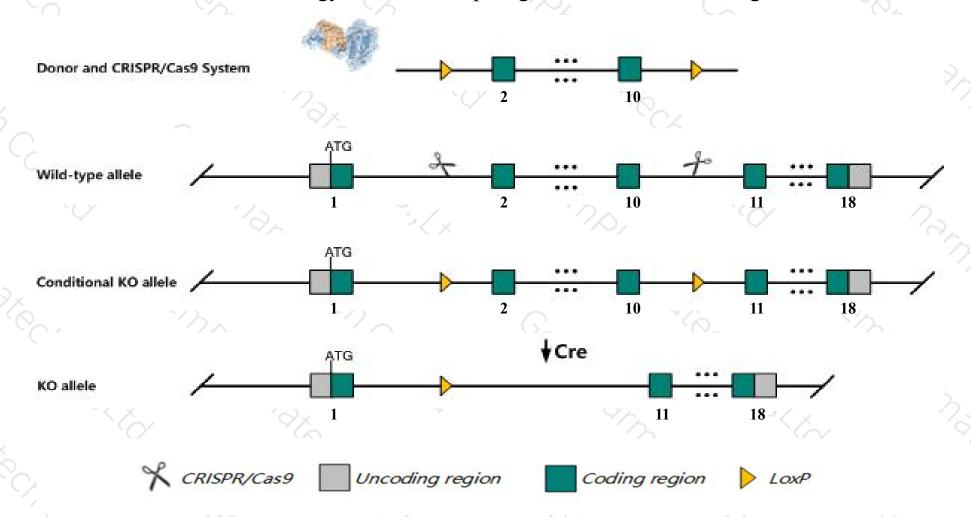
Project type Cas9-CKO

Strain background C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Hsph1* gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Hsph1* gene has 11 transcripts. According to the structure of *Hsph1* gene, exon2-exon10 of *Hsph1-211*(ENSMUST00000202361.3) transcript is recommended as the knockout region. The region contains 1271bp coding sequence.

 Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Hsph1* 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, Homozygous inactivation of this gene leads to decreased susceptibility to ischemic brain injury.
- The *Hsph1* gene is located on the Chr5. 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)



Hsph1 heat shock 105kDa/110kDa protein 1 [Mus musculus (house mouse)]

Gene ID: 15505, updated on 12-Aug-2019

Summary



Official Symbol Hsph1 provided by MGI

Official Full Name heat shock 105kDa/110kDa protein 1 provided by MGI

Primary source MGI:MGI:105053

See related Ensembl: ENSMUSG00000029657

Gene type protein coding
RefSeq status VALIDATED
Organism Mus musculus

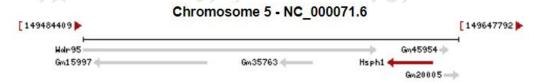
Lineage Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha;

Muroidea; Muridae; Murinae; Mus; Mus

Also known as 105kDa; Hsp105; Hsp110; hsp-E7I; Al790491; hsp110/105

Expression Broad expression in cortex adult (RPKM 30.2), CNS E11.5 (RPKM 29.9) and 25 other tissues See more

Orthologs human all



Transcript information (Ensembl)



The gene has 11 transcripts, all transcripts are shown below:

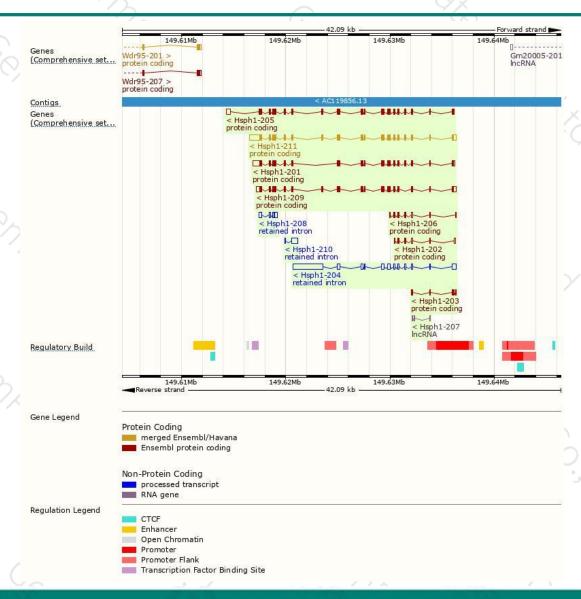
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Hsph1-211	ENSMUST00000202361.3	3802	858aa	Protein coding	CCDS19885	Q61699	TSL:1 GENCODE basic APPRIS P3
Hsph1-201	ENSMUST00000074846.13	3240	814aa	Protein coding	CCDS85010	Q61699	TSL:1 GENCODE basic APPRIS ALT
Hsph1-205	ENSMUST00000201452.3	3140	858aa	Protein coding	CCDS19885	Q61699	TSL:1 GENCODE basic APPRIS P3
Hsph1-209	ENSMUST00000202089.3	3054	817aa	Protein coding	12/	E9Q0U7	TSL:5 GENCODE basic
Hsph1-206	ENSMUST00000201559.3	661	<u>144aa</u>	Protein coding	187	D3Z3I9	CDS 3' incomplete TSL:5
Hsph1-202	ENSMUST00000200805.3	587	<u>94aa</u>	Protein coding	-	A0A0J9YTZ7	CDS 3' incomplete TSL:3
Hsph1-203	ENSMUST00000200825.1	416	<u>100aa</u>	Protein coding	828	D3Z027	CDS 3' incomplete TSL:2
Hsph1-204	ENSMUST00000201431.3	4764	No protein	Retained intron	1028	20	TSL:1
Hsph1-210	ENSMUST00000202137.1	752	No protein	Retained intron	187	7.6	TSL:2
Hsph1-208	ENSMUST00000201877.1	751	No protein	Retained intron	-	*1	TSL:2
Hsph1-207	ENSMUST00000201666.1	254	No protein	IncRNA	12	#8	TSL:5

The strategy is based on the design of *Hsph1-211* transcript, The transcription is shown below



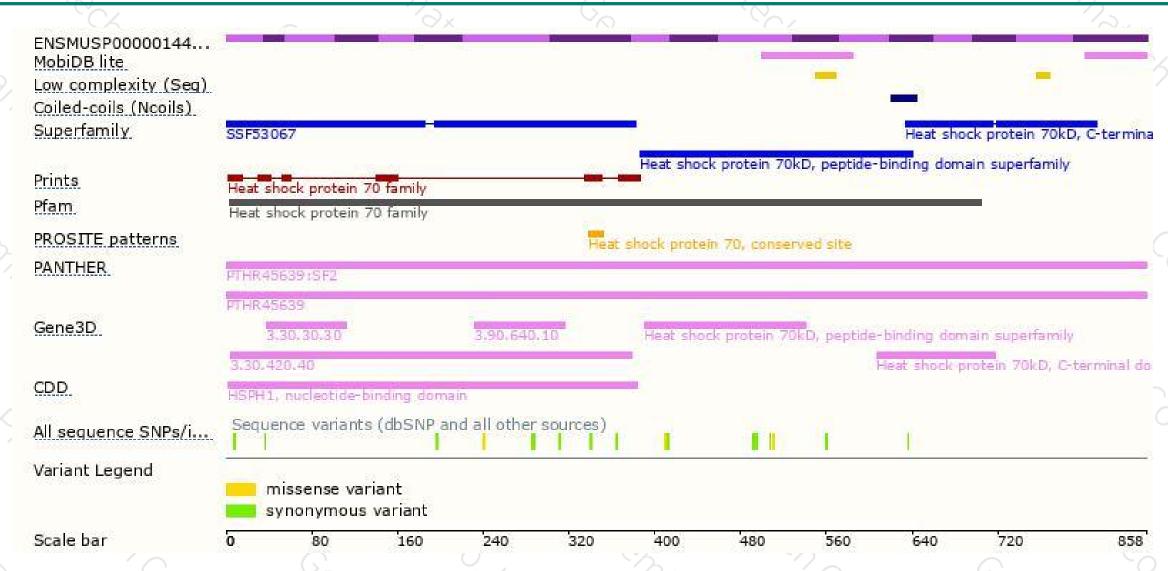
Genomic location distribution





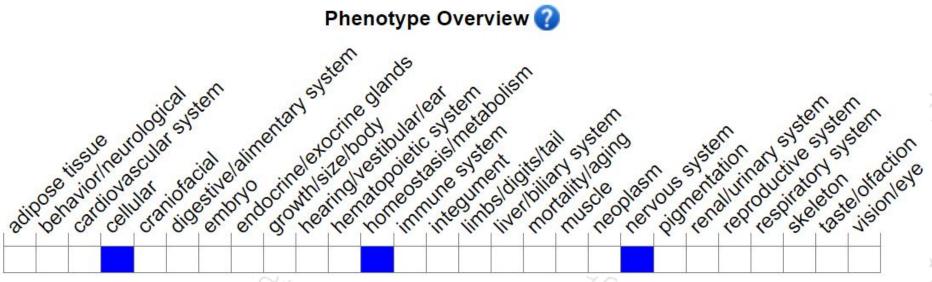
Protein domain





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, Homozygous inactivation of this gene leads to decreased susceptibility to ischemic brain injury.



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





