

Hdac4 Cas9-KO Strategy

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Design Date: 2019-7-18

Project Overview



Project Name

Hdac4

Project type

Cas9-KO

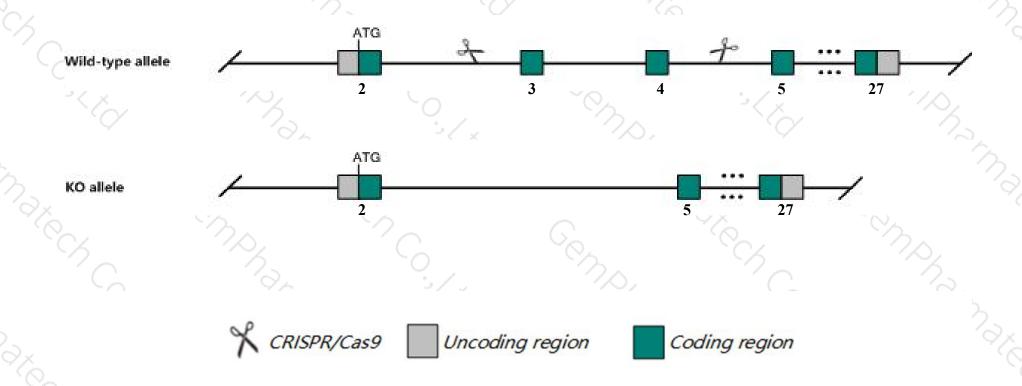
Strain background

C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Hdac4* gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Hdac4* gene has 12 transcripts. According to the structure of *Hdac4* gene, exon3-exon4 of *Hdac4-202*(ENSMUST00000097644.8) transcript is recommended as the knockout region. The region contains 314bp coding sequence.

 Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Hdac4* gene. The brief process is as follows: CRISPR/Cas9 system

Notice



- ➤ According to the existing MGI data, Mice homozygous for a gene trap allele exhibit increased thermal nociception threshold and seizures. Mice homozygous for a knock-out allele exhibit postnatal lethality, exencephaly, and abnormal skeleton morphology and physiology.
- > Transcript *Hdac4*-206 may not be affected.
- > The *Hdac4* 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)



Hdac4 histone deacetylase 4 [Mus musculus (house mouse)]

Gene ID: 208727, updated on 19-Mar-2019

Summary

☆ ?

Official Symbol Hdac4 provided by MGI

Official Full Name histone deacetylase 4 provided by MGI

Primary source MGI:MGI:3036234

See related Ensembl:ENSMUSG00000026313

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 4932408F19Rik, HD4

Expression Ubiquitous expression in adrenal adult (RPKM 12.3), ovary adult (RPKM 10.9) and 28 other tissuesSee more

Orthologs <u>human</u> all

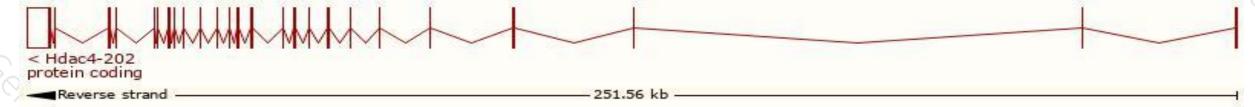
Transcript information (Ensembl)



The gene has 12 transcripts, all transcripts are shown below:

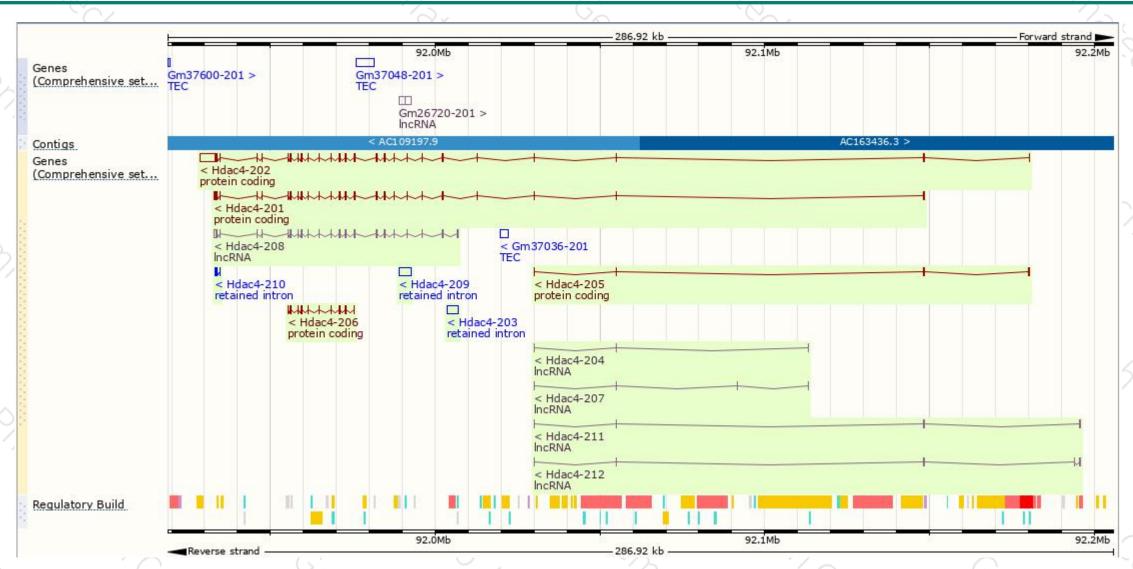
		1		``\ <u>+</u>			
Name 🛊	Transcript ID	bp 🌲	Protein	Biotype	CCDS .	UniProt	Flags
Hdac4-202	ENSMUST00000097644.8	8100	1076aa	Protein coding	CCDS48324 ₽	Q6NZM9®	TSL:1 GENCODE basic APPRIS P1
Hdac4-201	ENSMUST00000008995.14	3937	<u>1076aa</u>	Protein coding	CCDS48324 ₺	Q6NZM9 ₽	TSL:1 GENCODE basic APPRIS P1
Hdac4-206	ENSMUST00000187308.1	938	312aa	Protein coding		A0A087WQ92₺	CDS 5' and 3' incomplete TSL:5
Hdac4-205	ENSMUST00000186002.2	849	<u>112aa</u>	Protein coding	-	<u>A0A087WSF0</u> ₽	CDS 3' incomplete TSL:3
Hdac4-209	ENSMUST00000189730.1	3576	No protein	Retained intron	729	돧	TSL:NA
Hdac4-203	ENSMUST00000185267.1	3191	No protein	Retained intron	1926	8	TSL:NA
Hdac4-210	ENSMUST00000191327.1	589	No protein	Retained intron	1972	2	TSL:2
Hdac4-208	ENSMUST00000189303.6	3287	No protein	IncRNA	1.01		TSL:1
Hdac4-212	ENSMUST00000212867.1	797	No protein	IncRNA	3.5	=	TSL:5
Hdac4-211	ENSMUST00000212408.1	650	No protein	IncRNA		-	TSL:5
Hdac4-207	ENSMUST00000187622.1	410	No protein	IncRNA	*	-	TSL:3
Hdac4-204	ENSMUST00000185341.6	407	No protein	IncRNA	-	-	TSL:2

The strategy is based on the design of *Hdac4-202* transcript, The transcription is shown below



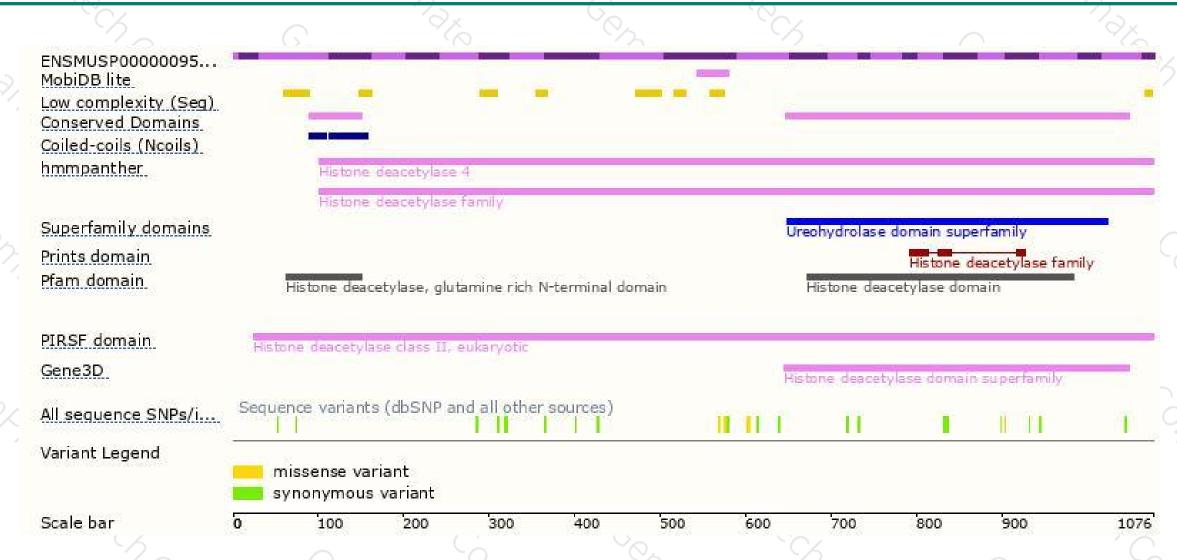
Genomic location distribution





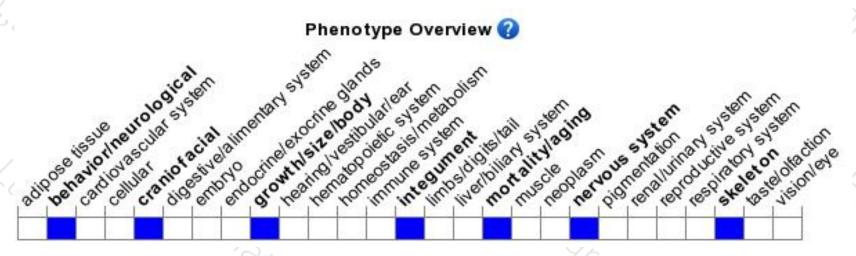
Protein domain





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, Mice homozygous for a gene trap allele exhibit increased thermal nociception threshold and seizures. Mice homozygous for a knock-out allele exhibit postnatal lethality, exencephaly, and abnormal skeletomorphology and physiology.



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





