

# H1f8 Cas9-KO Strategy

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Reviewer: Huimin Su

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## **Project Overview**



**Project Name** 

H1f8

**Project type** 

Cas9-KO

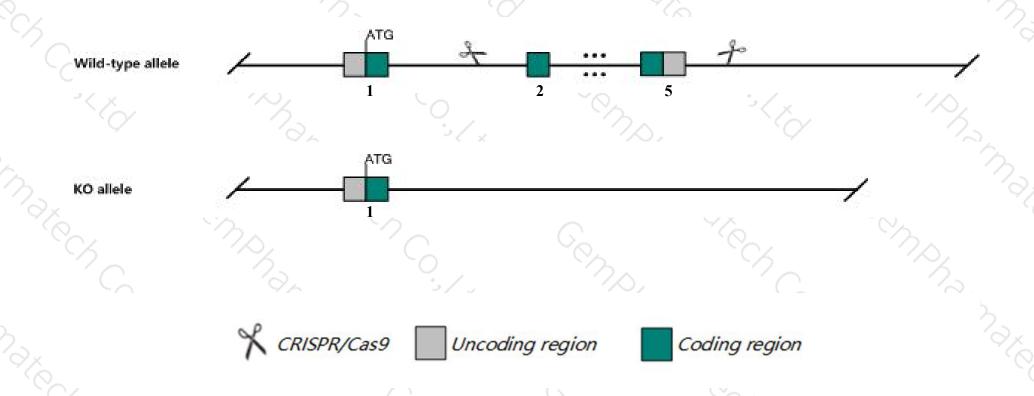
Strain background

C57BL/6JGpt

# **Knockout strategy**



This model will use CRISPR/Cas9 technology to edit the *H1f8* gene. The schematic diagram is as follows:



### **Technical routes**



- ➤ The *H1f8* gene has 5 transcripts. According to the structure of *H1f8* gene, exon2-exon5 of *H1f8-203*(ENSMUST00000161969.3) transcript is recommended as the knockout region. The region contains 641bp coding sequence.

  Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *H1f8* gene. The brief process is as follows: CRISPR/Cas9 system v

### **Notice**



- > According to the existing MGI data, Mice homozygous for a targeted mutation exhibit no detectable abnormalities. Oocytes develop normally and no defects in fertility or litter sizes are observed.
- The *H1f8* gene is located on the Chr6. 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)



#### H1f8 H1.8 linker histone [ Mus musculus (house mouse) ]

Gene ID: 171506, updated on 12-Aug-2019

#### Summary

2

Official Symbol H1f8 provided by MGI

Official Full Name H1.8 linker histone provided by MGI

Primary source MGI:MGI:2176207

See related Ensembl: ENSMUSG00000042279

Gene type protein coding
RefSeq status REVIEWED
Organism Mus musculus

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

Muroidea; Muridae; Murinae; Mus; Mus

Also known as H1-8; H1.8; H1fo; H1oo; H1foo; C86609

Summary Histories are basic nuclear proteins that are responsible for the nucleosome structure of the chromosomal fiber in eukaryotes.

Nucleosomes consist of approximately 146 bp of DNA wrapped around a histone octamer composed of pairs of each of the four core histones (H2A, H2B, H3, and H4). The chromatin fiber is further compacted through the interaction of a linker histone, H1, with the DNA between the nucleosomes to form higher order chromatin structures. The protein encoded is a replication-independent histone that is a member of the histone H1 family. This gene contains introns, unlike most histone genes and the encoded protein is expressed only in

oocytes. [provided by RefSeq, Oct 2015]

Expression Restricted expression toward ovary adult (RPKM 8.4) See more

Orthologs human all

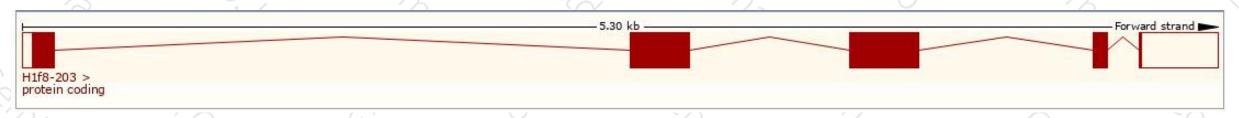
# Transcript information (Ensembl)



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

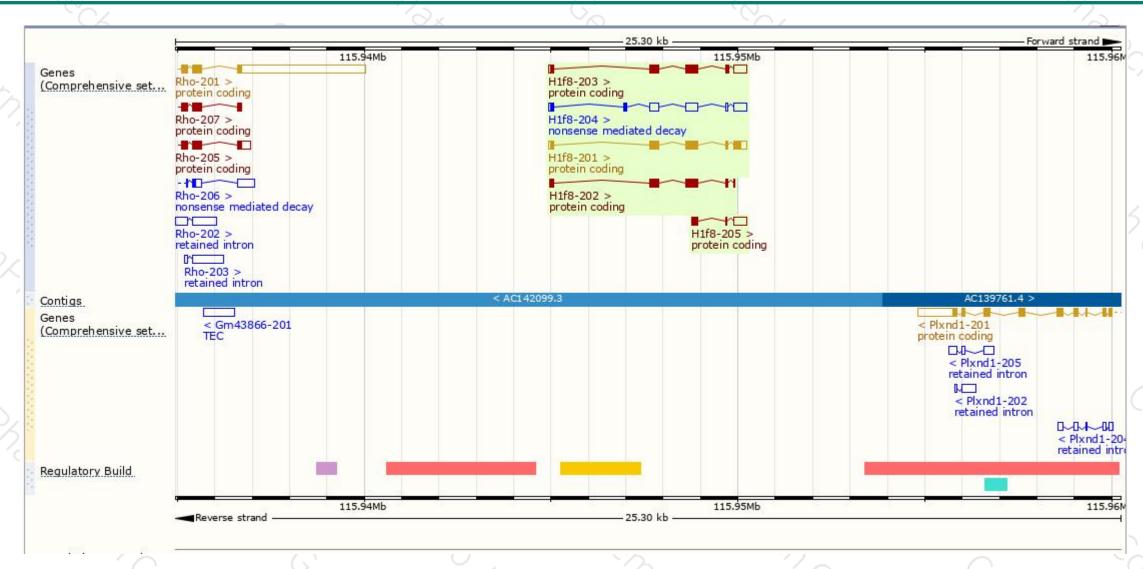
Name 🌲	Transcript ID	bp 🌲	Protein 🌲	Biotype	CCDS 🍦	UniProt	Flags
H1f8-203	ENSMUST00000161969.3	1122	246aa	Protein coding	CCDS85133 ₪	Q8VIK3₽	TSL:1 GENCODE basic APPRIS ALT2
H1f8-201	ENSMUST00000037831.13	1113	304aa	Protein coding	CCDS20447 ₺	Q8VIK3₽	TSL:1 GENCODE basic APPRIS P3
H1f8-202	ENSMUST00000161617.6	785	<u>251aa</u>	Protein coding		E0CZ52₺	CDS 3' incomplete TSL:3
H1f8-205	ENSMUST00000205177.1	520	<u>64aa</u>	Protein coding	323	A0A0N4SV54 ₺	CDS 5' incomplete TSL:3
H1f8-204	ENSMUST00000162084.7	1192	<u>46aa</u>	Nonsense mediated decay	755	E0CYL2₽	TSL:1

The strategy is based on the design of *H1f8-203* transcript, The transcription is shown below



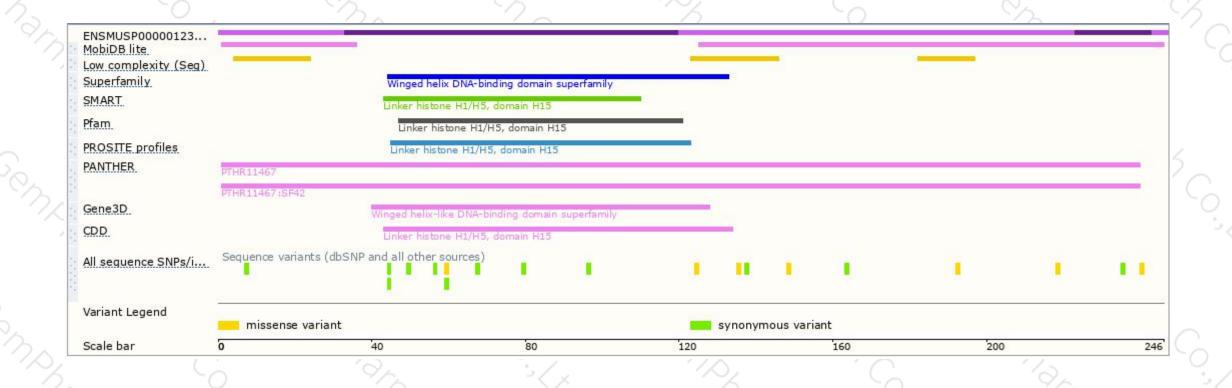
### Genomic location distribution





### 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 targeted mutation exhibit no detectable abnormalities. Oocytes develop normally and no defects in fertility or litter sizes are observed.



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





