

Hivep2 Cas9-KO Strategy

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Reviewer: Daohua Xu

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Project Overview



Project Name

Hivep2

Project type

Cas9-KO

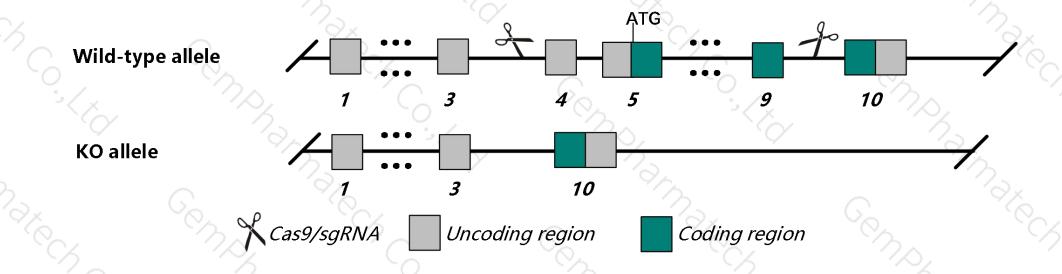
Strain background

C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Hivep2* gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Hivep2* gene has 6 transcripts. According to the structure of *Hivep2* gene, exon4-exon9 of *Hivep2-205*(ENSMUST00000191138.6) transcript is recommended as the knockout region. The region contains most of the coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Hivep2* gene. The brief process is as follows: CRISPR/Cas9 system 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 knock-out allele display abnormal thymus anatomy, severely defective positive selection of CD4+ and CD8+ cells, and enhanced T-helper 2 cell differentiation.
- ➤ Transcript *Hivep2*-202&206 may not be affected.
- > The *Hivep2* 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)



Hivep2 human immunodeficiency virus type I enhancer binding protein 2 [Mus musculus (house mouse)]

Gene ID: 15273, updated on 13-Mar-2020

Summary

☆ ?

Official Symbol Hivep2 provided by MGI

Official Full Name human immunodeficiency virus type I enhancer binding protein 2 provided by MGI

Primary source MGI:MGI:1338076

See related Ensembl: ENSMUSG00000015501

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 Gm20114, MIBP-1, MIBP1, Schnurri-2, Shn-2

Expression Broad expression in cortex adult (RPKM 18.2), frontal lobe adult (RPKM 17.6) and 20 other tissuesSee more

Orthologs <u>human all</u>

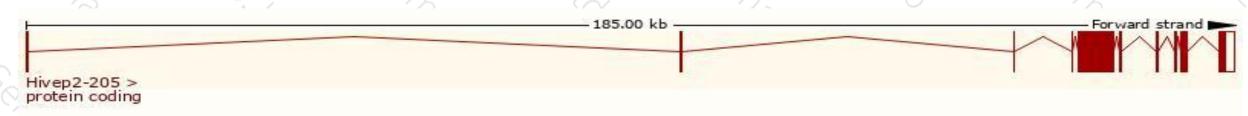
Transcript information (Ensembl)



The gene has 6 transcripts, all transcripts are shown below:

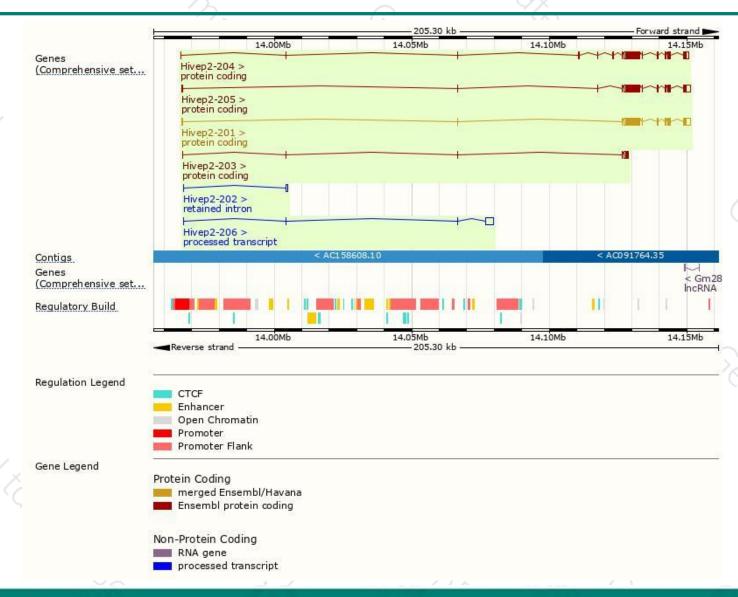
| Transcript ID | bp | Protein | Biotype | CCDS | UniProt | Flags |
|-----------------------|---|--|--|---|---|---|
| ENSMUST00000191138.6 | 9846 | 2430aa | Protein coding | CCDS23704 | Q3UHF7 | TSL:1 GENCODE basic APPRIS is a system to annotate alternatively spliced transcripts based on a range of computational methods to identify the most functionally important transcript(s) of a gene. APPRIS P1 |
| ENSMUST00000015645.10 | 9763 | 2430aa | Protein coding | CCDS23704 | Q3UHF7 | TSL:1 GENCODE basic APPRIS is a system to annotate alternatively spliced transcripts based on a range of computational methods to identify the most functionally important transcript(s) of a gene. APPRIS P1 |
| ENSMUST00000187083.6 | 9268 | 2430aa | Protein coding | CCDS23704 | Q3UHF7 | TSL:1 GENCODE basic APPRIS is a system to annotate alternatively spliced transcripts based on a range of computational methods to identify the most functionally important transcript(s) of a gene. APPRIS P1 |
| ENSMUST00000186989.6 | 1762 | 373aa | Protein coding | | A0A087WQF9 | CDS 3' incomplete TSL:1 |
| ENSMUST00000191464.1 | 2899 | No protein | Processed transcript | | - | TSL:1 |
| ENSMUST00000185392.1 | 758 | No protein | Retained intron | - | - | TSL:5 |
| | ENSMUST00000191138.6 ENSMUST00000015645.10 ENSMUST00000187083.6 ENSMUST00000186989.6 ENSMUST00000191464.1 | ENSMUST00000191138.6 9846 ENSMUST00000015645.10 9763 ENSMUST00000187083.6 9268 ENSMUST00000186989.6 1762 ENSMUST00000191464.1 2899 | ENSMUST00000191138.6 9846 2430aa ENSMUST00000015645.10 9763 2430aa ENSMUST00000187083.6 9268 2430aa ENSMUST00000186989.6 1762 373aa ENSMUST00000191464.1 2899 No protein | ENSMUST00000191138.6 9846 2430aa Protein coding ENSMUST00000015645.10 9763 2430aa Protein coding ENSMUST00000187083.6 9268 2430aa Protein coding ENSMUST00000186989.6 1762 373aa Protein coding ENSMUST00000191464.1 2899 No protein Processed transcript | ENSMUST00000191138.6 9846 2430aa Protein coding CCDS23704 ENSMUST00000015645.10 9763 2430aa Protein coding CCDS23704 ENSMUST00000187083.6 9268 2430aa Protein coding CCDS23704 ENSMUST00000186989.6 1762 373aa Protein coding - ENSMUST00000191464.1 2899 No protein Processed transcript - | ENSMUST00000191138.6 9846 2430aa Protein coding CCDS23704 Q3UHF7 ENSMUST00000015645.10 9763 2430aa Protein coding CCDS23704 Q3UHF7 ENSMUST00000187083.6 9268 2430aa Protein coding CCDS23704 Q3UHF7 ENSMUST00000186989.6 1762 373aa Protein coding - A0A087WQF9 ENSMUST00000191464.1 2899 No protein Processed transcript - - |

The strategy is based on the design of *Hivep2-205* transcript, the transcription is shown below:



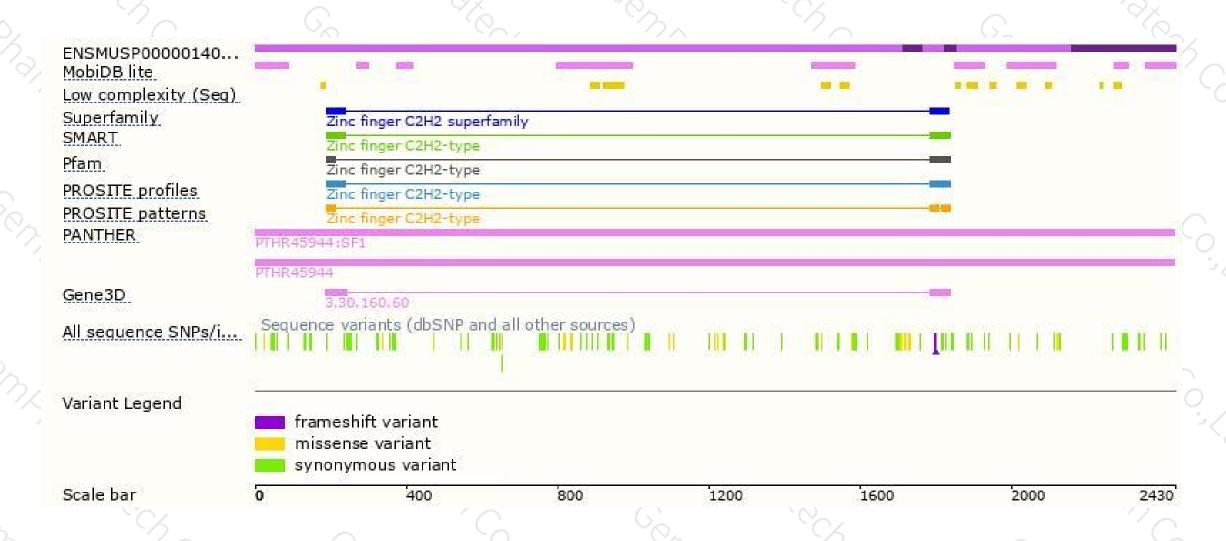
Genomic location distribution





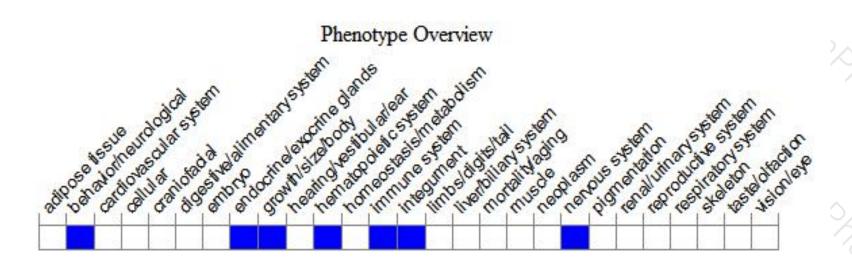
Protein domain





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 knock-out allele display abnormal thymus anatomy, severely defective positive selection of CD4+ and CD8+ cells, and enhanced T-helper 2 cell differentiation.



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





