

# *Hes1* Cas9-KO Strategy

**Designer:**

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**Design Date:**

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# Project Overview

**Project Name**

*Hes1*

**Project type**

**Cas9-KO**

**Strain background**

**C57BL/6JGpt**

# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Hes1* gene. The schematic diagram is as follows:



- The *Hes1* gene has 4 transcripts. According to the structure of *Hes1* gene, exon1-exon4 of *Hes1-201* (ENSMUST00000023171.7) transcript is recommended as the knockout region. The region contains all 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 *Hes1* gene. The brief process is as follows: CRISPR/Cas9 system v

- According to the existing MGI data, Mutants show anomalous timing in neurogenesis. Homozygotes for a null allele exhibit premature neurogenesis, severe neural tube defects, supernumerary hair cells in the inner ear, increased numbers of pulmonary neuroendocrine cells, and pancreatic hypoplasia. Death occurs in utero or neonatally.
- The *Hes1* gene is located on the Chr16. 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)

## Hes1 hes family bHLH transcription factor 1 [Mus musculus (house mouse)]

Gene ID: 15205, updated on 9-Apr-2019

### Summary



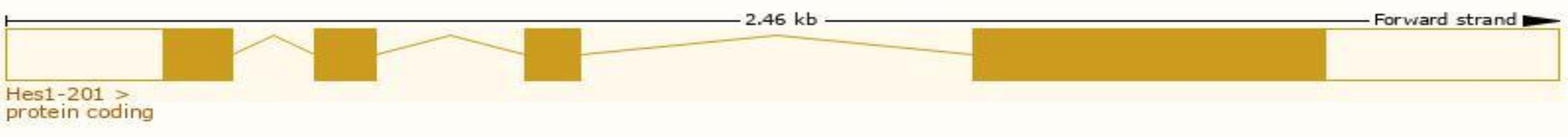
<b>Official Symbol</b>	Hes1 provided by <a href="#">MGI</a>
<b>Official Full Name</b>	hes family bHLH transcription factor 1 provided by <a href="#">MGI</a>
<b>Primary source</b>	<a href="#">MGI:MGI:104853</a>
<b>See related</b>	<a href="#">Ensembl:ENSMUSG00000022528</a>
<b>Gene type</b>	protein coding
<b>RefSeq status</b>	PROVISIONAL
<b>Organism</b>	<a href="#">Mus musculus</a>
<b>Lineage</b>	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
<b>Also known as</b>	Hry, bHLHb39
<b>Expression</b>	Broad expression in stomach adult (RPKM 82.2), colon adult (RPKM 65.6) and 21 other tissues <a href="#">See more</a>
<b>Orthologs</b>	<a href="#">human</a> <a href="#">all</a>

# Transcript information (Ensembl)

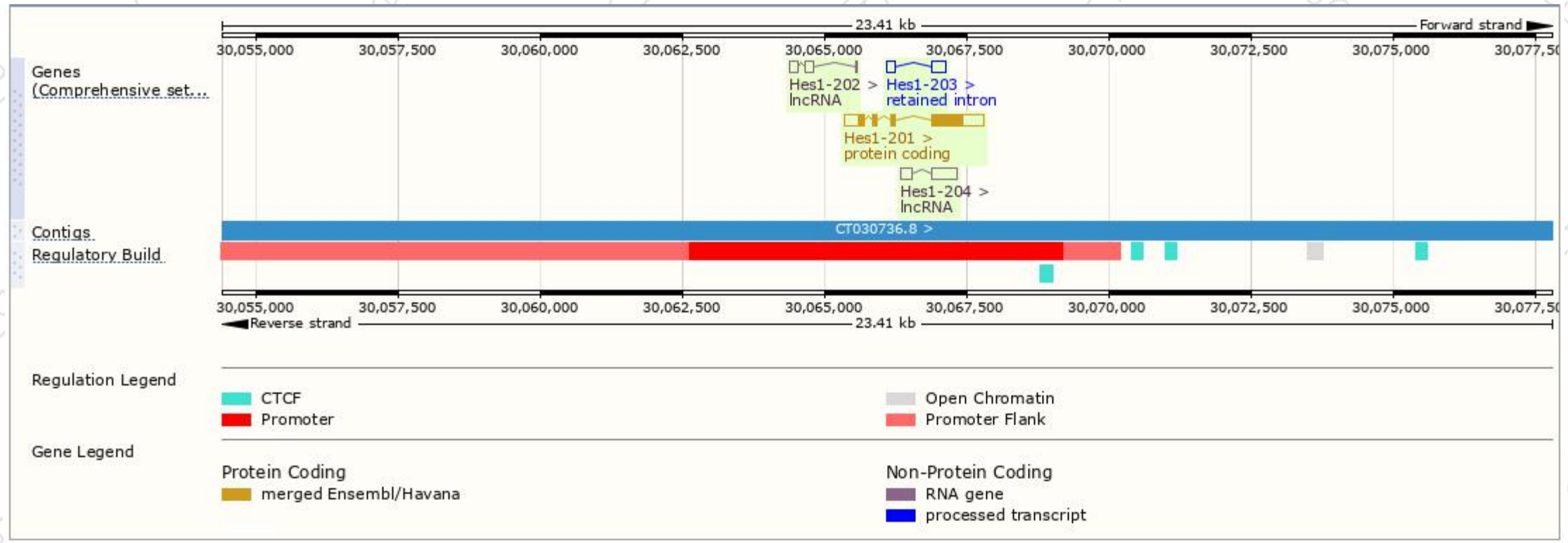
The gene has 4 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Hes1-201	<a href="#">ENSMUST00000023171.7</a>	1468	<a href="#">282aa</a>	Protein coding	<a href="#">CCDS28098</a>	<a href="#">P35428</a> <a href="#">Q3UZZ2</a>	TSL:1 Gencode basic APPRIS P1
Hes1-203	<a href="#">ENSMUST00000161676.1</a>	406	No protein	Retained intron	-	-	TSL:2
Hes1-204	<a href="#">ENSMUST00000161839.1</a>	653	No protein	lncRNA	-	-	TSL:3
Hes1-202	<a href="#">ENSMUST00000160592.2</a>	344	No protein	lncRNA	-	-	TSL:2

The strategy is based on the design of *Hes1-201* transcript,The transcription is shown below

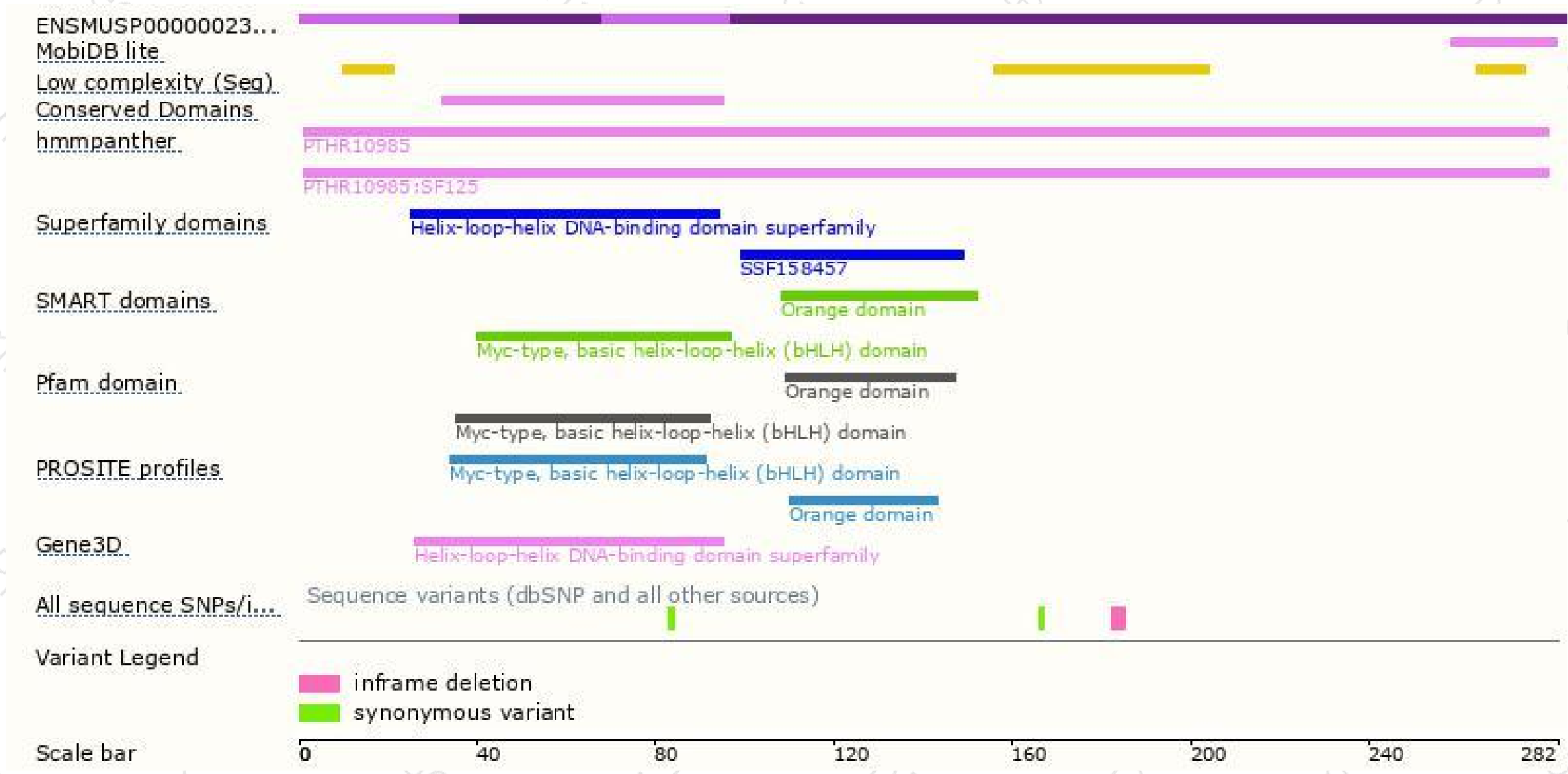


# Genomic location distribution

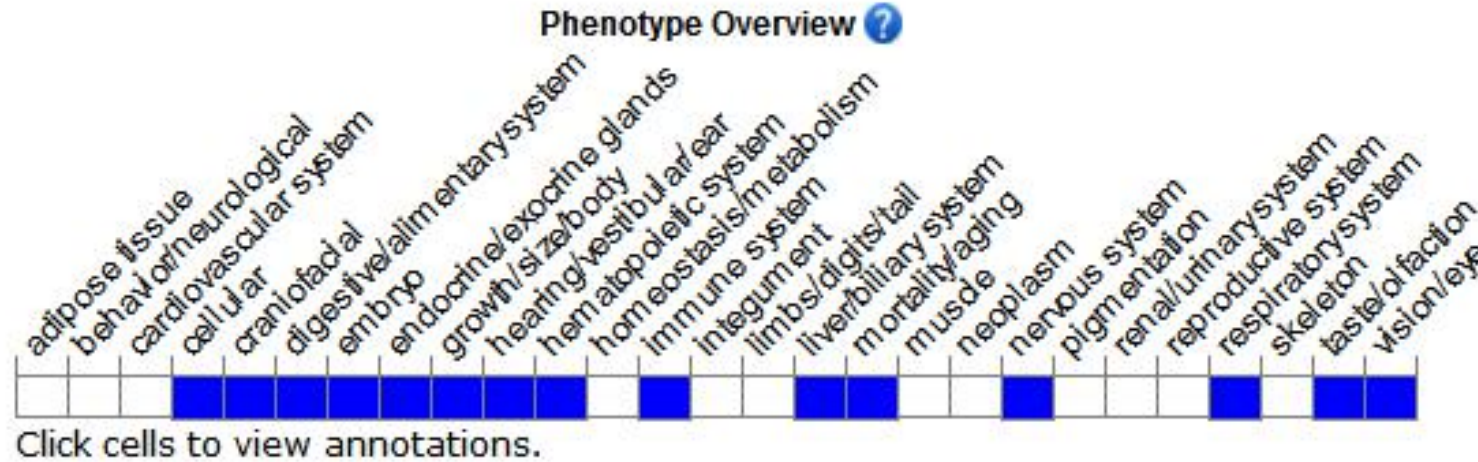




# Protein domain



# 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, Mutants show anomalous timing in neurogenesis. Homozygotes for a null allele exhibit premature neurogenesis, severe neural tube defects, supernumerary hair cells in the inner ear, increased numbers of pulmonary neuroendocrine cells, and pancreatic hypoplasia. Death occurs in utero or neonatally.

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

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