

# ***Morc3 Cas9-KO Strategy***

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# Project Overview

**Project Name**

*Morc3*

**Project type**

**Cas9-KO**

**Strain background**

**C57BL/6JGpt**

# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Morc3* gene. The schematic diagram is as follows:



- The *Morc3* gene has 10 transcripts. According to the structure of *Morc3* gene, exon3-exon5 of *Morc3-206* (ENSMUST00000202261.4) transcript is recommended as the knockout region. The region contains 496bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Morc3* gene. The brief process is as follows: CRISPR/Cas9 system

- According to the existing MGI data, Mice homozygous for a null allele die at or within a day of birth.
- The non-coding transcripts 204, 210 are unaffected.
- The *Morc3* 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)

## Morc3 microrchidia 3 [Mus musculus (house mouse)]

Gene ID: 338467, updated on 31-Jan-2019

### Summary



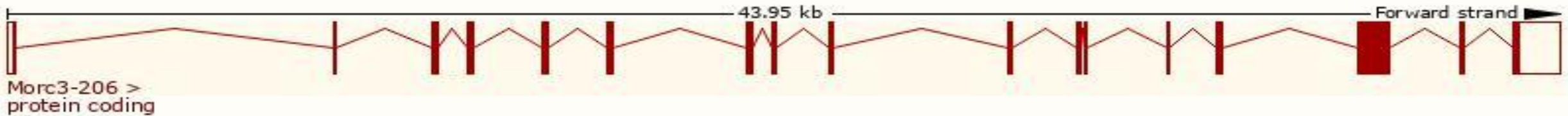
<b>Official Symbol</b>	Morc3 provided by <a href="#">MGI</a>
<b>Official Full Name</b>	microrchidia 3 provided by <a href="#">MGI</a>
<b>Primary source</b>	<a href="#">MGI:MGI:2136841</a>
<b>See related</b>	<a href="#">Ensembl:ENSMUSG000000039456</a>
<b>Gene type</b>	protein coding
<b>RefSeq status</b>	VALIDATED
<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>	1110051N18Rik, AI452146, BF318192, D16Jhu32e, NXP2, Zcwcc3
<b>Expression</b>	Ubiquitous expression in CNS E18 (RPKM 8.1), CNS E11.5 (RPKM 7.9) and 27 other tissues <a href="#">See more</a>
<b>Orthologs</b>	<a href="#">human</a> <a href="#">all</a>

# Transcript information (Ensembl)

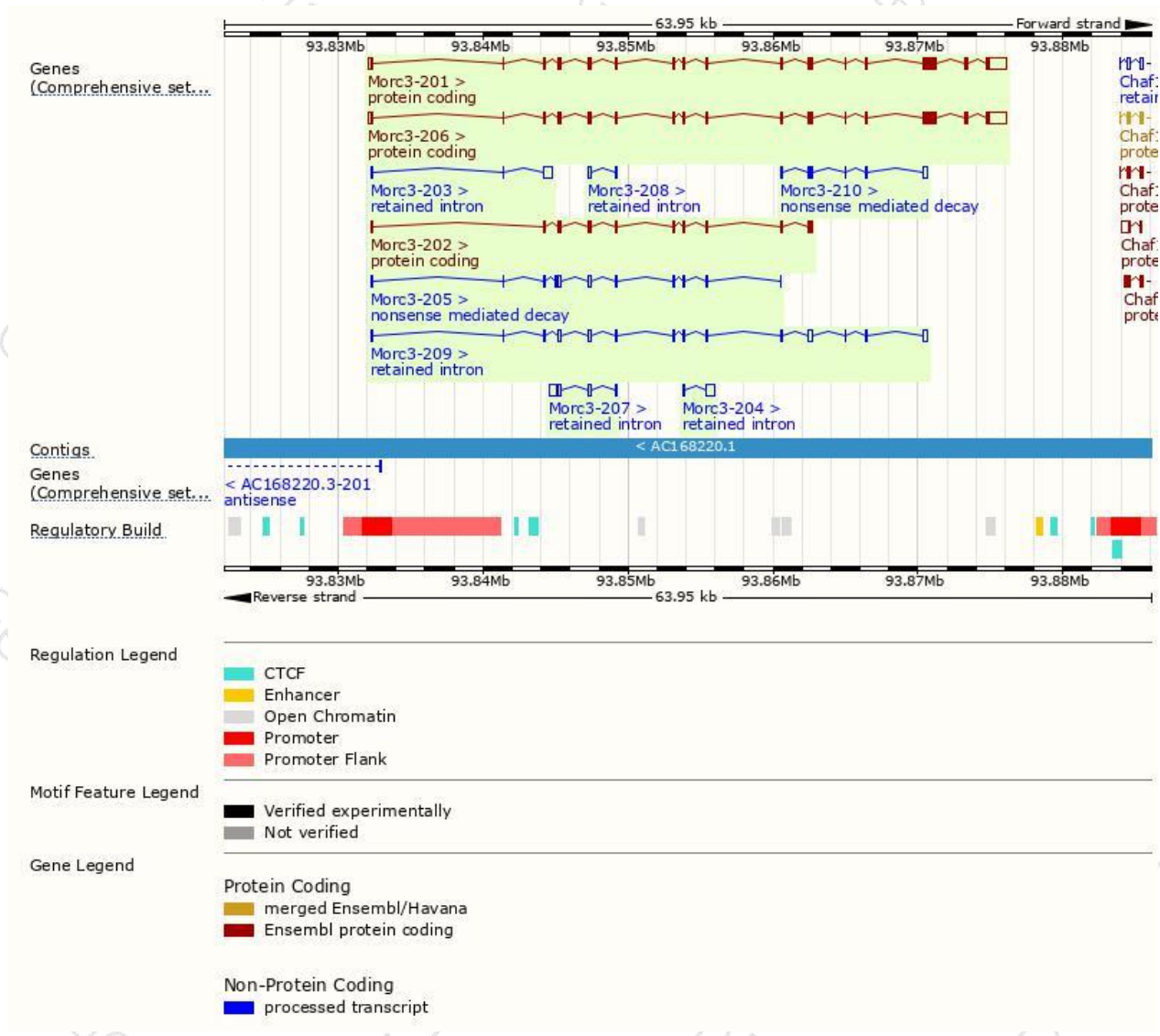
The gene has 10 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Morc3-206	<a href="#">ENSMUST00000202261.4</a>	4179	<a href="#">942aa</a>	Protein coding	<a href="#">CCDS79490</a>	<a href="#">F7BJB9</a>	TSL:1 GENCODE basic APPRIS P1
Morc3-201	<a href="#">ENSMUST00000044068.9</a>	4178	<a href="#">942aa</a>	Protein coding	-	<a href="#">F7BJB9</a>	TSL:5 GENCODE basic APPRIS P1
Morc3-202	<a href="#">ENSMUST00000201097.2</a>	1386	<a href="#">397aa</a>	Protein coding	-	<a href="#">A0A0J9YU83</a>	CDS 3' incomplete TSL:5
Morc3-205	<a href="#">ENSMUST00000201754.3</a>	1298	<a href="#">92aa</a>	Nonsense mediated decay	-	<a href="#">A0A0J9YUV3</a>	TSL:1
Morc3-210	<a href="#">ENSMUST00000232639.1</a>	726	<a href="#">58aa</a>	Nonsense mediated decay	-	<a href="#">A0A338P7B1</a>	CDS 5' incomplete
Morc3-209	<a href="#">ENSMUST00000232425.1</a>	2114	No protein	Retained intron	-	-	
Morc3-207	<a href="#">ENSMUST00000202663.1</a>	882	No protein	Retained intron	-	-	TSL:3
Morc3-203	<a href="#">ENSMUST00000201497.1</a>	735	No protein	Retained intron	-	-	TSL:2
Morc3-204	<a href="#">ENSMUST00000201706.1</a>	634	No protein	Retained intron	-	-	TSL:3
Morc3-208	<a href="#">ENSMUST00000231891.1</a>	246	No protein	Retained intron	-	-	

The strategy is based on the design of *Morc3-206* transcript,The transcription is shown below

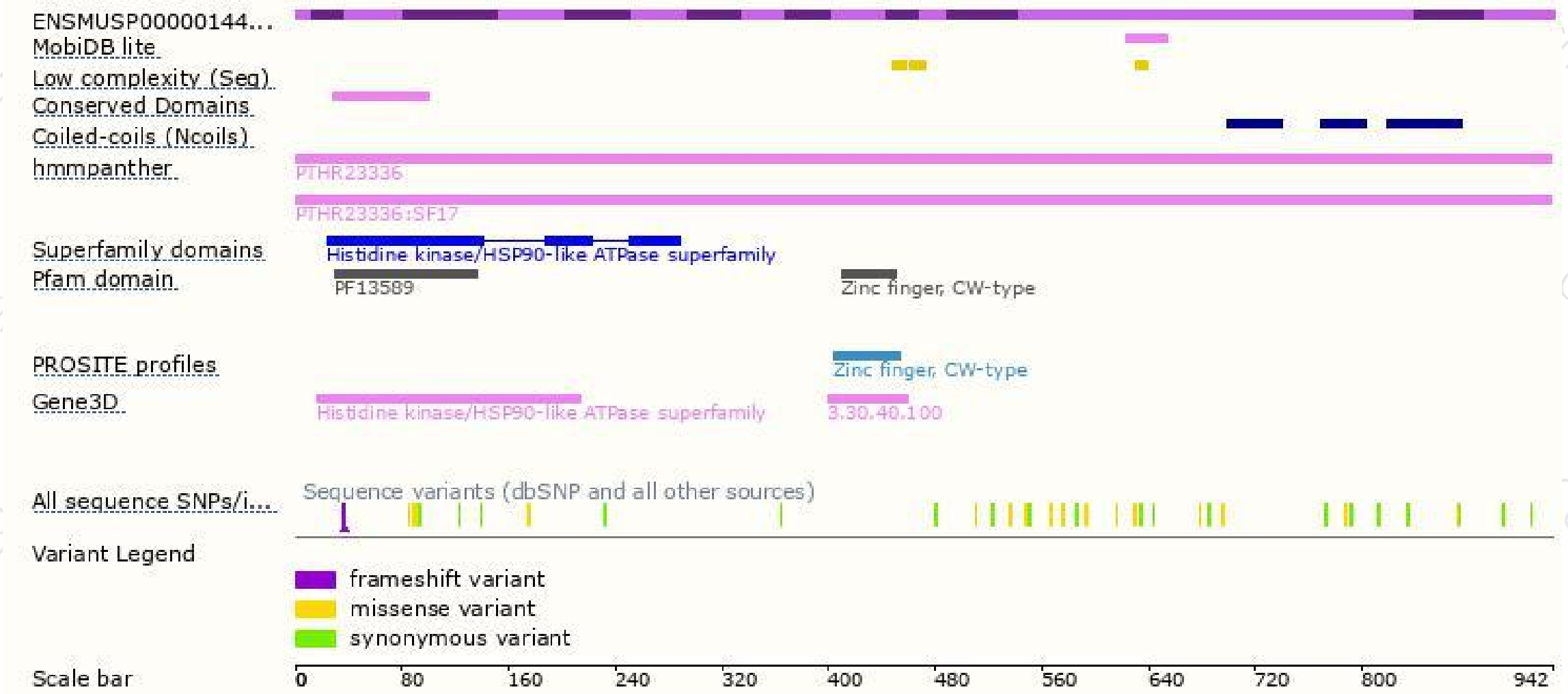


# Genomic location distribution

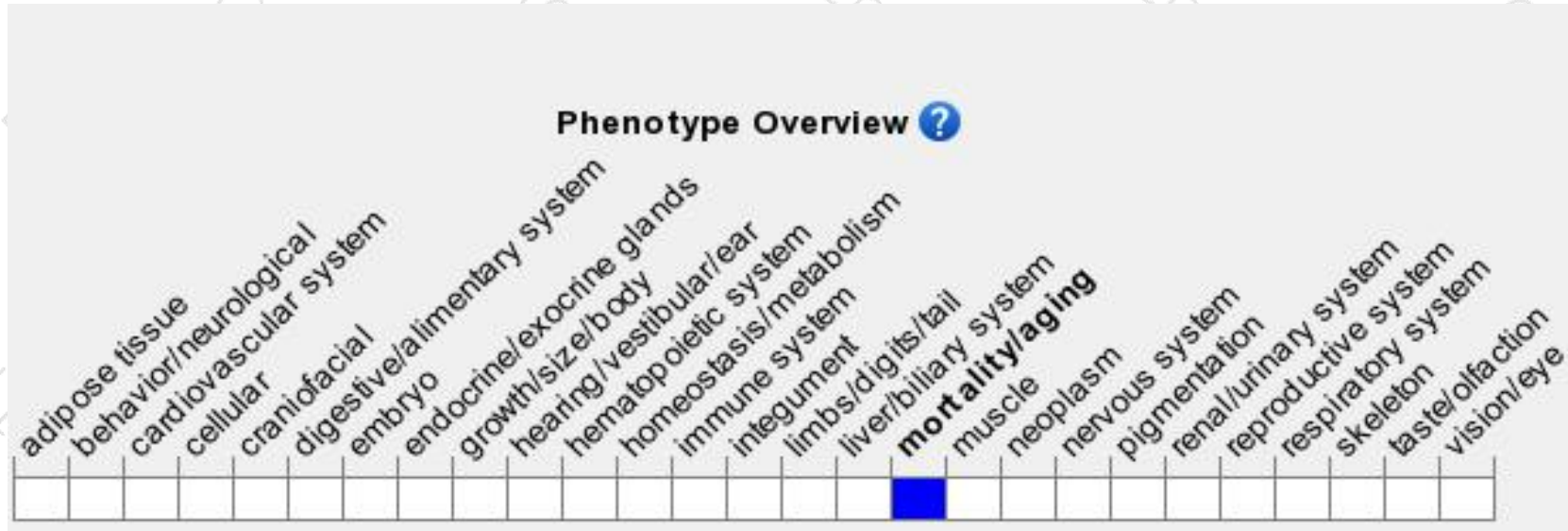




# Protein domain



# 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 null allele die at or within a day of birth.

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

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