

# *Dot1l* Cas9-KO Strategy

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**Reviewer:**

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# Project Overview

**Project Name**

***Dot1l***

**Project type**

**Cas9-KO**

**Strain background**

**C57BL/6JGpt**

# Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Dot1l* gene. The schematic diagram is as follows:



- The *Dot1l* gene has 7 transcripts. According to the structure of *Dot1l* gene, exon2 of *Dot1l-201* (ENSMUST00000105336.8) transcript is recommended as the knockout region. The region contains 44bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Dot1l* gene. The brief process is as follows: CRISPR/Cas9 system

- According to the existing MGI data, Mice homozygous for a gene trap allele show late embryonic lethality. Mice homozygous for a null allele die by E10.5 displaying a growth arrest, abnormal yolk sac angiogenesis and heart dilation while mutant ES cells show elevated apoptosis, G2 cell cycle arrest, telomere elongation and aneuploidy.
- Transcript *Dot1l-203,206* may not be affected.
- The *Dot1l* 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)

## Dot1l DOT1-like, histone H3 methyltransferase (*S. cerevisiae*) [*Mus musculus* (house mouse)]

Gene ID: 208266, updated on 12-Feb-2019

### Summary



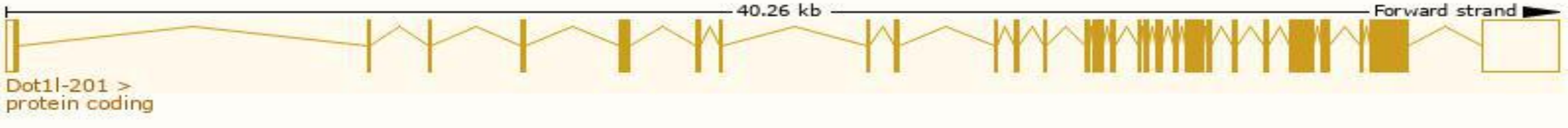
<b>Official Symbol</b>	Dot1l provided by <a href="#">MGI</a>
<b>Official Full Name</b>	DOT1-like, histone H3 methyltransferase ( <i>S. cerevisiae</i> ) provided by <a href="#">MGI</a>
<b>Primary source</b>	<a href="#">MGI:MGI:2143886</a>
<b>See related</b>	<a href="#">Ensembl:ENSMUSG00000061589</a>
<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>	A630076O07, AW907654, Dot1, KMT4, mDot1
<b>Expression</b>	Ubiquitous expression in testis adult (RPKM 34.5), adrenal adult (RPKM 27.1) and 28 other tissues <a href="#">See more</a>
<b>Orthologs</b>	<a href="#">human</a> <a href="#">all</a>

# Transcript information (Ensembl)

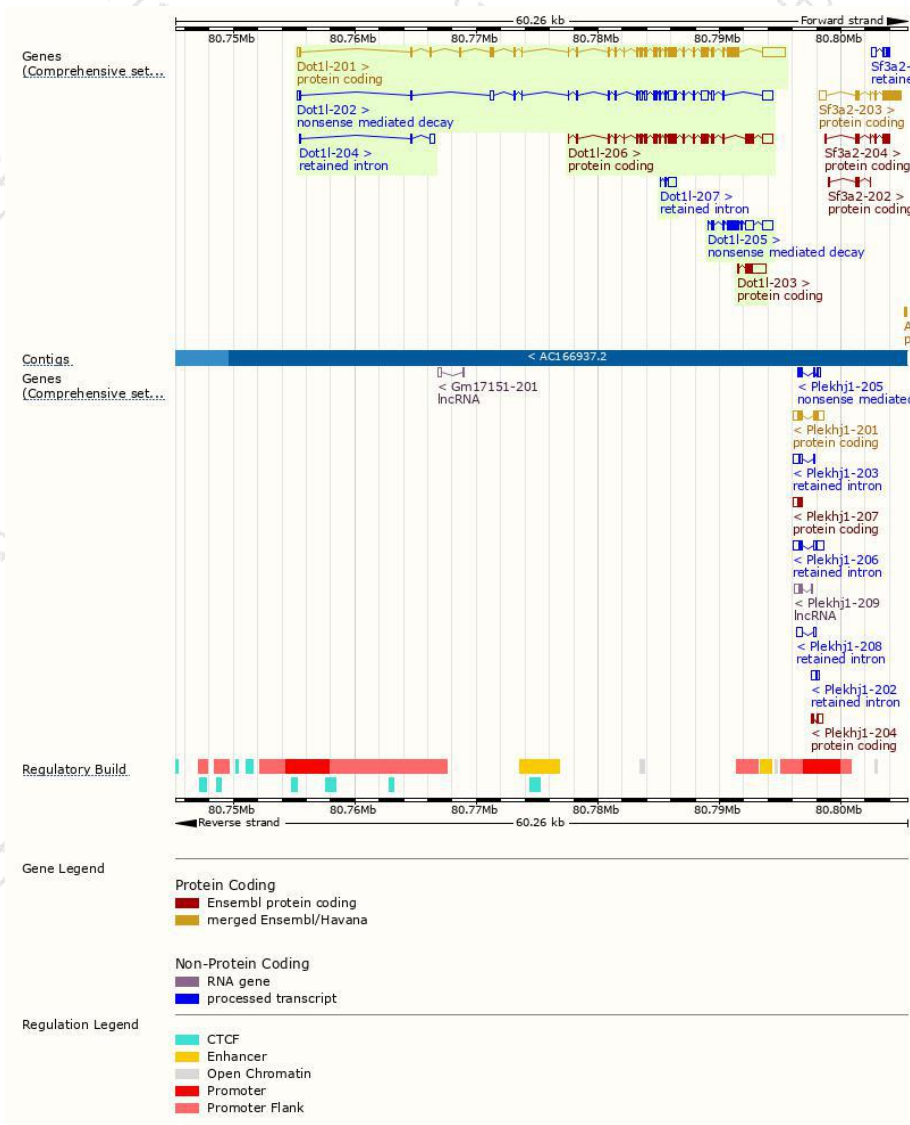
The gene has 7 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Dot1l-201	<a href="#">ENSMUST00000105336.8</a>	6808	<a href="#">1540aa</a>	Protein coding	<a href="#">CCDS35985</a>	<a href="#">Q6XZL8</a>	TSL:1 GENCODE basic APPRIS P1
Dot1l-206	<a href="#">ENSMUST00000150338.7</a>	4529	<a href="#">1202aa</a>	Protein coding	-	<a href="#">F7CVL0</a>	CDS 5' incomplete TSL:1
Dot1l-203	<a href="#">ENSMUST00000138505.1</a>	1784	<a href="#">234aa</a>	Protein coding	-	<a href="#">F6S070</a>	CDS 5' incomplete TSL:1
Dot1l-202	<a href="#">ENSMUST00000127740.7</a>	4600	<a href="#">49aa</a>	Nonsense mediated decay	-	<a href="#">E9Q6I8</a>	TSL:1
Dot1l-205	<a href="#">ENSMUST00000149394.1</a>	2820	<a href="#">424aa</a>	Nonsense mediated decay	-	<a href="#">F6SJX8</a>	CDS 5' incomplete TSL:1
Dot1l-207	<a href="#">ENSMUST00000163526.1</a>	790	No protein	Retained intron	-	-	TSL:3
Dot1l-204	<a href="#">ENSMUST00000147579.1</a>	531	No protein	Retained intron	-	-	TSL:2

The strategy is based on the design of *Dot1l-201* transcript,The transcription is shown below

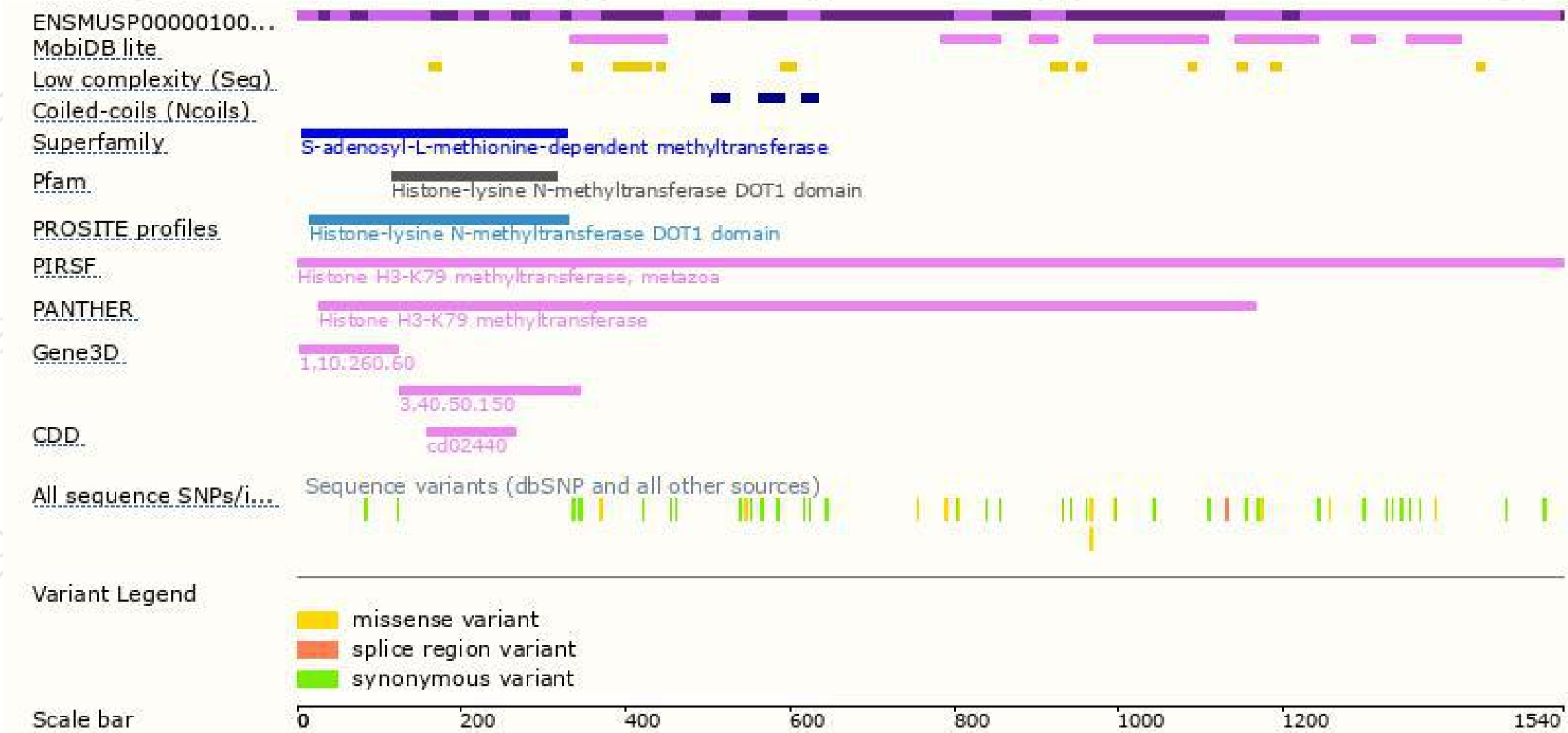


# Genomic location distribution

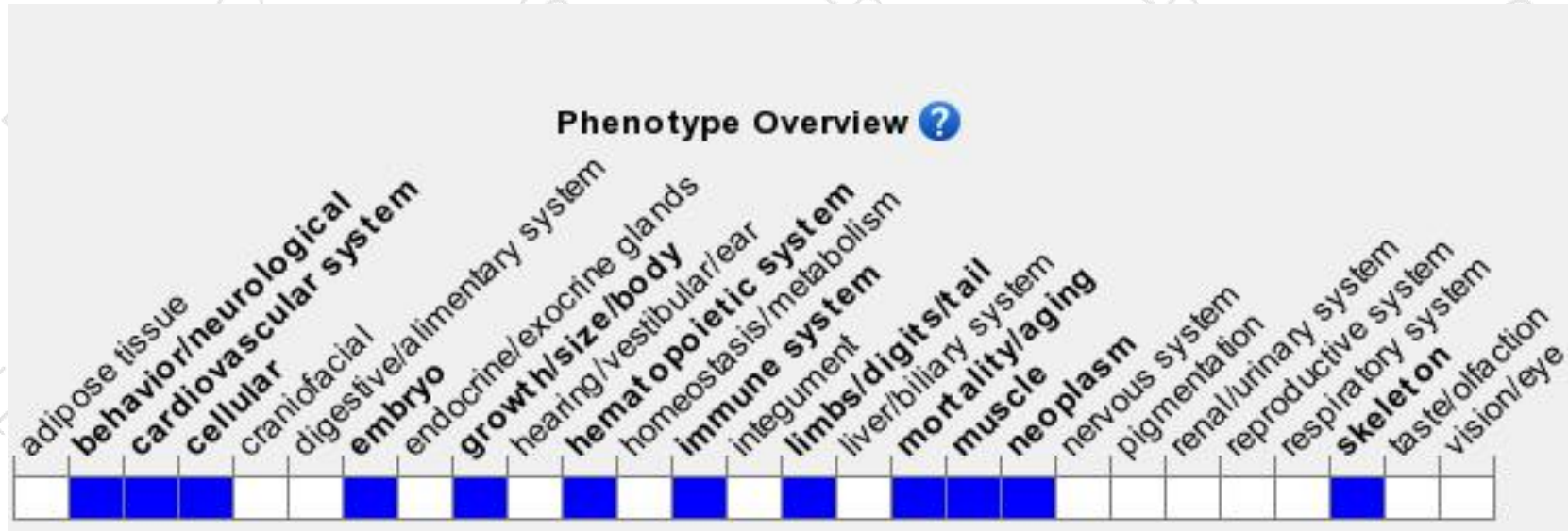




# Protein domain



# 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, Mice homozygous for a gene trap allele show late embryonic lethality. Mice homozygous for a null allele die by E10.5 displaying a growth arrest, abnormal yolk sac angiogenesis and heart dilation while mutant ES cells show elevated apoptosis, G2 cell cycle arrest, telomere elongation and aneuploidy.

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

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