

# *Dtna* Cas9-KO Strategy

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**Reviewer:**

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# Project Overview

**Project Name**

***Dtna***

**Project type**

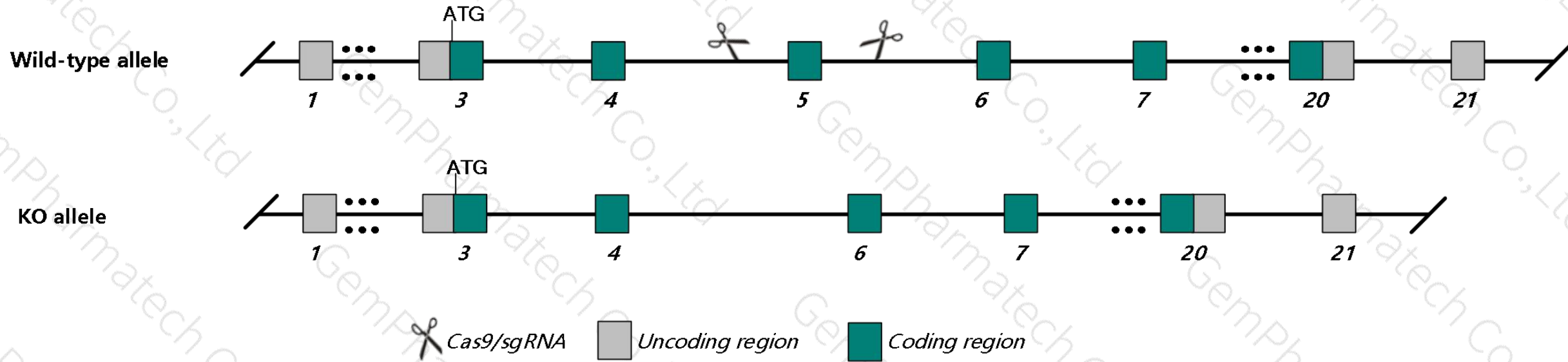
**Cas9-KO**

**Strain background**

**C57BL/6JGpt**

# Knockout strategy

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



- The *Dtna* gene has 10 transcripts. According to the structure of *Dtna* gene, exon5 of *Dtna-202* (ENSMUST00000115832.3) transcript is recommended as the knockout region. The region contains 214bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Dtna* gene. The brief process is as follows: CRISPR/Cas9 system v

- According to the existing MGI data, Homozygous targeted mutants exhibit skeletal and cardiac myopathies. Neuromuscular junctions appear to form normally, but their postnatal maturation is compromised. *Dtna* mutations do not increase the severity of *Dmd* or *Utrn* mutants whose products are also part of the dystrophin-glycoprotein complex.
- The CDS of transcript *Dtna*-208,209 are incomplete, whether they will be affected is unknown.
- The *Dtna* gene is located on the Chr18. 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 loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at existing technological level.



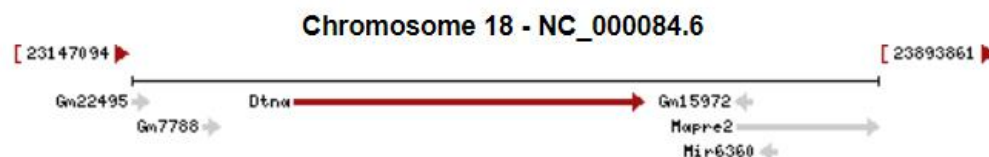
# Gene information (NCBI)

## Dtna dystrobrevin alpha [ *Mus musculus* (house mouse) ]

Gene ID: 13527, updated on 24-Oct-2019

### Summary

<b>Official Symbol</b>	Dtna provided by <a href="#">MGI</a>
<b>Official Full Name</b>	dystrobrevin alpha provided by <a href="#">MGI</a>
<b>Primary source</b>	<a href="#">MGI:MGI:106039</a>
<b>See related</b>	<a href="#">Ensembl:ENSMUSG00000024302</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>	Dtn; adbn; DTN-A; a-DB-1; Gm19389; 2210407P21Rik
<b>Expression</b>	Broad expression in cerebellum adult (RPKM 9.3), cortex adult (RPKM 8.0) and 16 other tissues <a href="#">See more</a>
<b>Orthologs</b>	<a href="#">human</a> <a href="#">all</a>

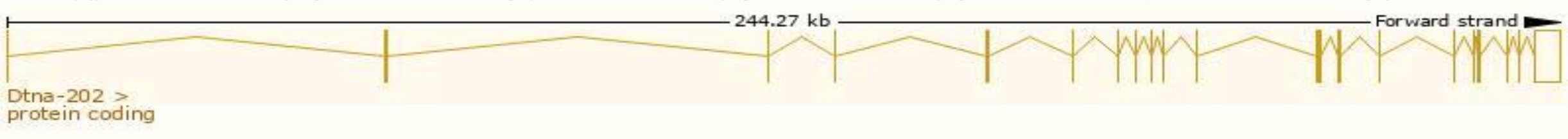


# Transcript information (Ensembl)

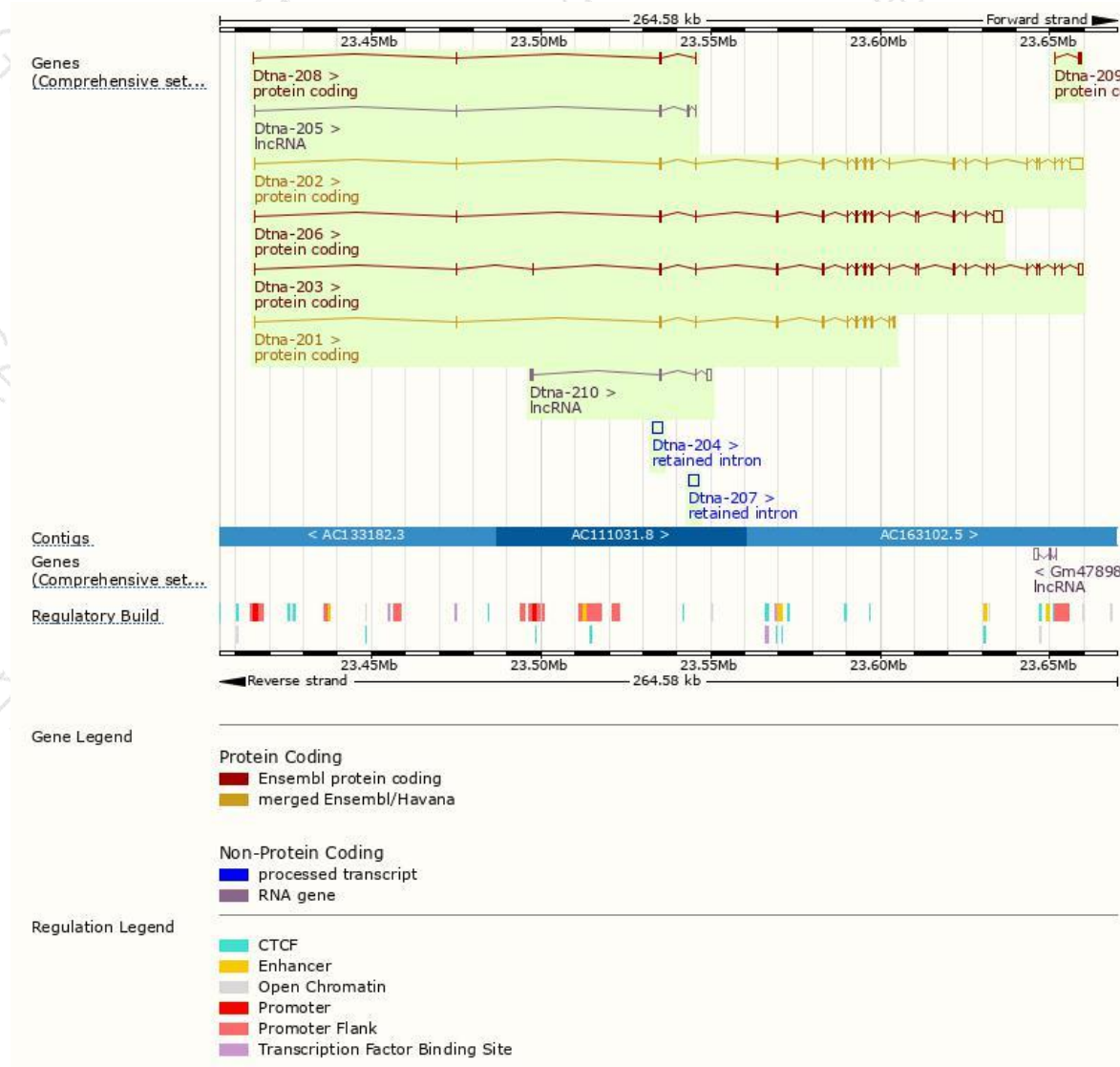
The gene has 10 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Dtna-202	<a href="#">ENSMUST00000115832.3</a>	6295	<a href="#">682aa</a>	Protein coding	<a href="#">CCDS50236</a>	<a href="#">Q8CFR5</a>	TSL:1 GENCODE basic APPRIS ALT 1
Dtna-201	<a href="#">ENSMUST00000047954.14</a>	1596	<a href="#">371aa</a>	Protein coding	<a href="#">CCDS29094</a>	<a href="#">Q8BTD9 Q9D2N4</a>	TSL:1 GENCODE basic APPRIS P3
Dtna-206	<a href="#">ENSMUST00000221880.1</a>	4490	<a href="#">567aa</a>	Protein coding	-	<a href="#">A0A1Y7VJN9</a>	TSL:2 GENCODE basic APPRIS ALT 1
Dtna-203	<a href="#">ENSMUST00000220904.1</a>	3860	<a href="#">746aa</a>	Protein coding	-	<a href="#">A0A1Y7VL34</a>	TSL:1 GENCODE basic
Dtna-209	<a href="#">ENSMUST00000222726.1</a>	863	<a href="#">73aa</a>	Protein coding	-	<a href="#">A0A1Y7VLZ6</a>	CDS 5' incomplete TSL:5
Dtna-208	<a href="#">ENSMUST00000222515.1</a>	359	<a href="#">35aa</a>	Protein coding	-	<a href="#">A0A1Y7VK79</a>	CDS 3' incomplete TSL:2
Dtna-207	<a href="#">ENSMUST00000222351.1</a>	3086	No protein	Retained intron	-	-	TSL:NA
Dtna-204	<a href="#">ENSMUST00000220967.1</a>	2941	No protein	Retained intron	-	-	TSL:NA
Dtna-210	<a href="#">ENSMUST00000223441.1</a>	1489	No protein	lncRNA	-	-	TSL:1
Dtna-205	<a href="#">ENSMUST00000221471.1</a>	723	No protein	lncRNA	-	-	TSL:3

The strategy is based on the design of *Dtna-202* transcript,The transcription is shown below

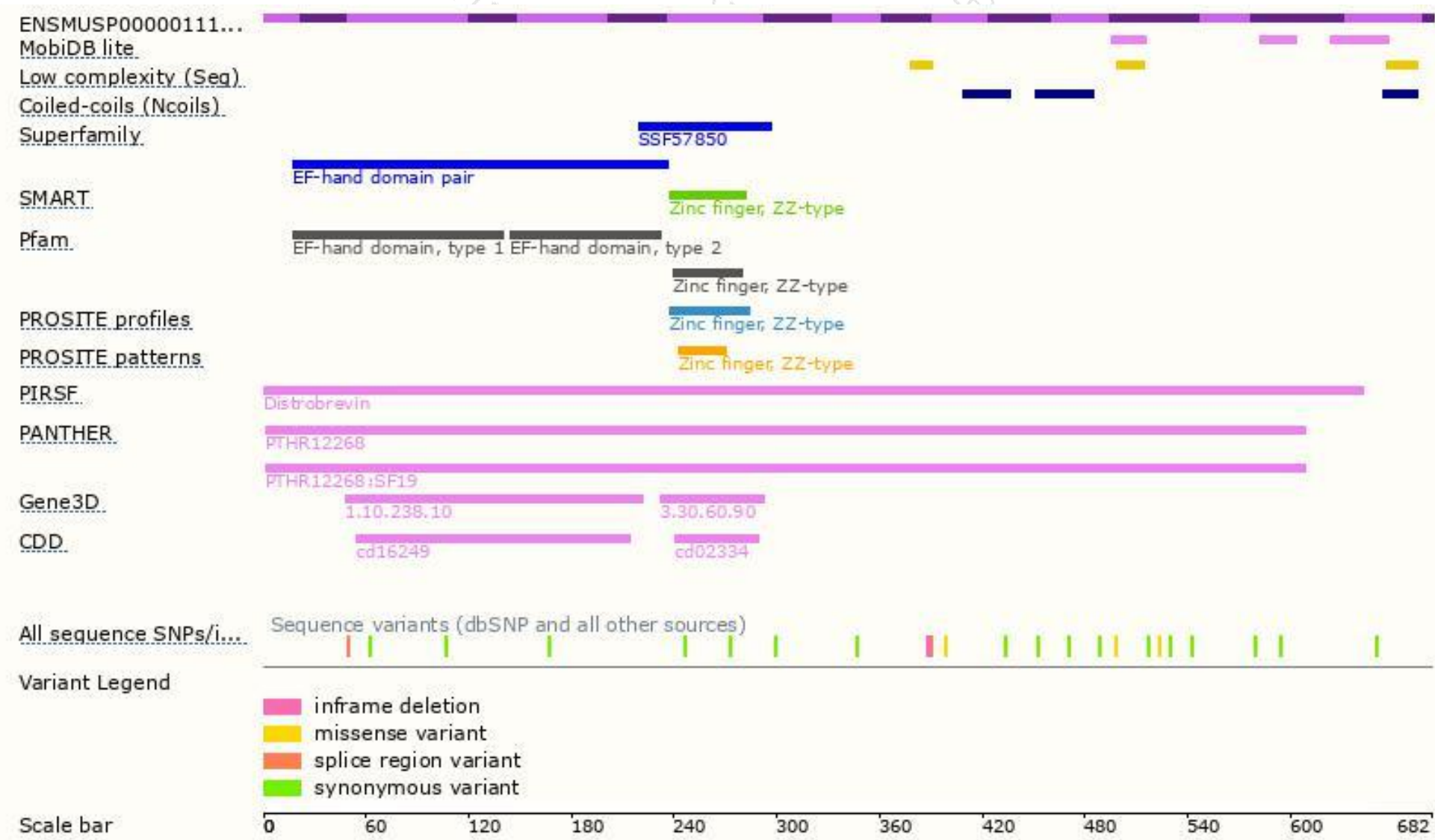


# Genomic location distribution

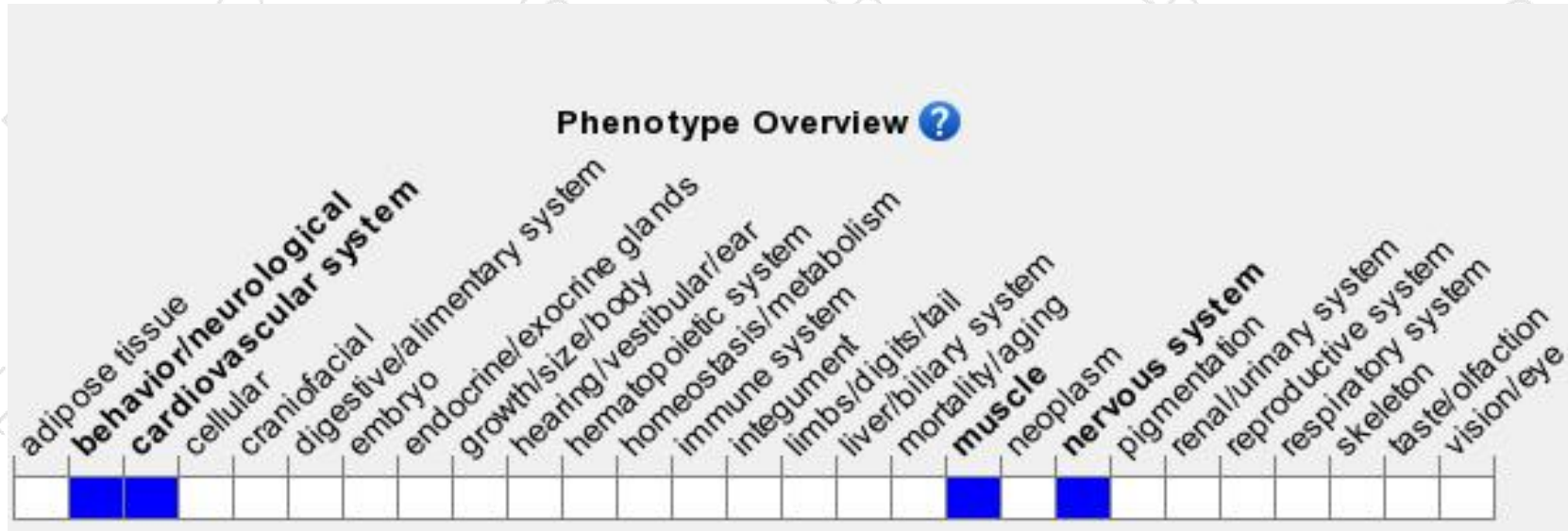




# Protein domain



# 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, Homozygous targeted mutants exhibit skeletal and cardiac myopathies. Neuromuscular junctions appear to form normally, but their postnatal maturation is compromised. Dtna mutations do not increase the severity of Dmd or Utrn mutants whose products are also part of the dystrophin-glycoprotein complex.

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

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