

# *Myo1c* Cas9-CKO Strategy

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**Reviewer:**

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# Project Overview

**Project Name**

*Myo1c*

**Project type**

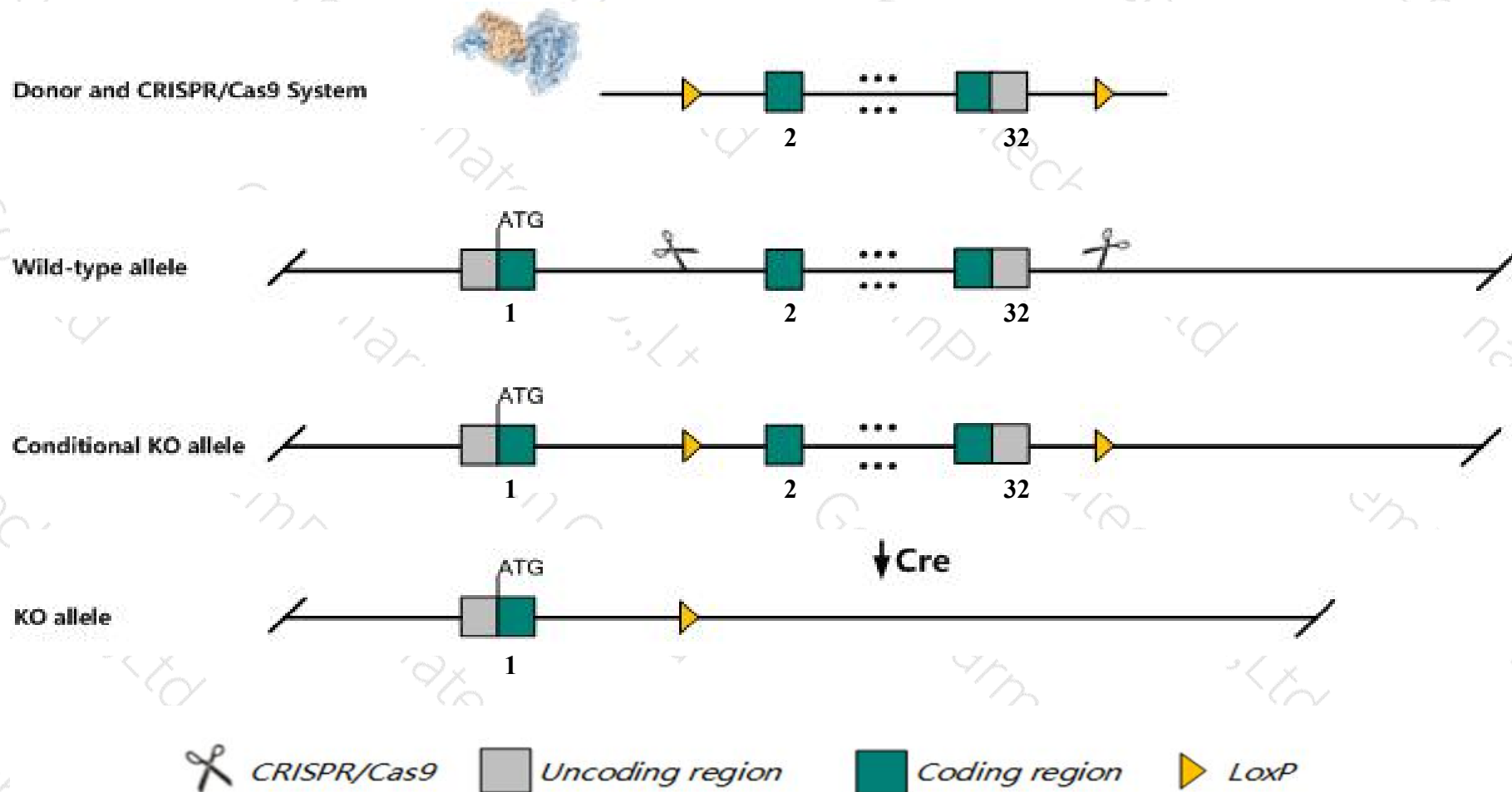
**Cas9-CKO**

**Strain background**

**C57BL/6JGpt**

# Conditional Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Myo1c* gene. The schematic diagram is as follows:



- The *Myo1c* gene has 10 transcripts. According to the structure of *Myo1c* gene, exon2-exon32 of *Myo1c*-204 (ENSMUST00000108431.2) 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 *Myo1c* gene. The brief process is as follows: CRISPR/Cas9 system and Donor 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.
- The flox mice will be knocked out after mating with mice expressing Cre recombinase, resulting in the loss of function of the target gene in specific tissues and cell types.

- According to the existing MGI data, Mice homozygous for a knock-in (Y61G) mutation that sensitizes to N6-modified ADP analogs display altered fast adaption in vestibular hair cells. Mice homozygous for a nuclear isoform-specific knock-out allele exhibit minor changes in bone marrow density and red blood cells.
- The *Myo1c* gene is located on the Chr11. 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)

## Myo1c myosin IC [Mus musculus (house mouse)]

Gene ID: 17913, updated on 3-Feb-2019

### Summary



**Official Symbol** Myo1c provided by [MGI](#)

**Official Full Name** myosin IC provided by [MGI](#)

**Primary source** [MGI:MGI:106612](#)

**See related** [Ensembl:ENSMUSG00000017774](#)

**Gene type** protein coding

**RefSeq status** REVIEWED

**Organism** [Mus musculus](#)

**Lineage** Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus

**Also known as** C80397, MM1b, MYO1E, NMI, mm1beta, myr2

**Summary** This gene encodes a member of the unconventional myosin protein family, which are actin-based molecular motors. The protein is found in the cytoplasm, and one isoform with a unique N-terminus is also found in the nucleus. The protein functions in intracellular vesicle transport to the plasma membrane. The nuclear isoform associates with RNA polymerase I and II and functions in transcription initiation. Multiple transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Jul 2008]

**Expression** Ubiquitous expression in lung adult (RPKM 70.4), subcutaneous fat pad adult (RPKM 51.1) and 24 other tissues [See more](#)

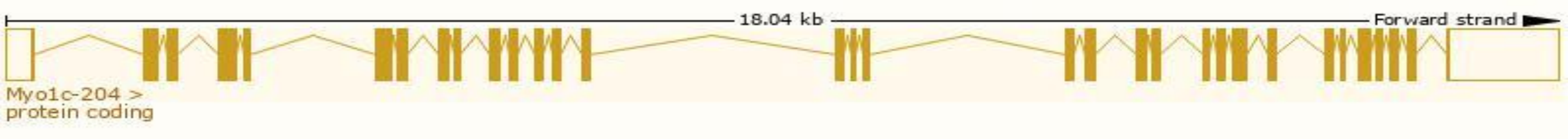
**Orthologs** [human](#) [all](#)

# Transcript information (Ensembl)

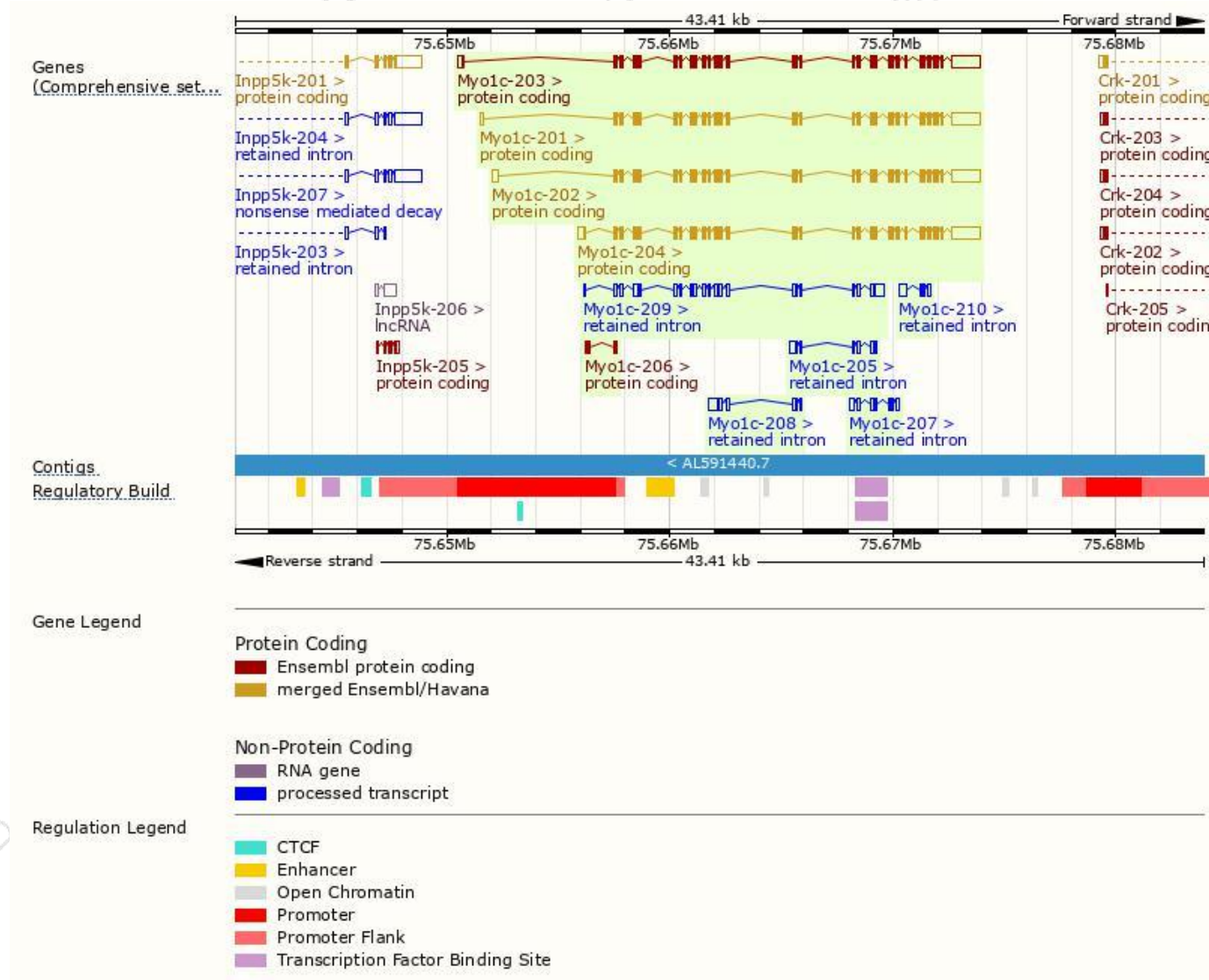
The gene has 10 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Myo1c-204	<a href="#">ENSMUST00000108431.2</a>	4718	<a href="#">1044aa</a>	Protein coding	<a href="#">CCDS36228</a>	<a href="#">Q9WTI7</a>	TSL:1 GENCODE basic APPRIS ALT 1
Myo1c-202	<a href="#">ENSMUST00000102504.9</a>	4659	<a href="#">1028aa</a>	Protein coding	<a href="#">CCDS25054</a>	<a href="#">Q9WTI7</a>	TSL:1 GENCODE basic APPRIS P3
Myo1c-201	<a href="#">ENSMUST00000069057.12</a>	4547	<a href="#">1028aa</a>	Protein coding	<a href="#">CCDS25054</a>	<a href="#">Q9WTI7</a>	TSL:1 GENCODE basic APPRIS P3
Myo1c-203	<a href="#">ENSMUST00000102505.9</a>	4652	<a href="#">1063aa</a>	Protein coding	-	<a href="#">Q9WTI7</a>	TSL:5 GENCODE basic APPRIS ALT 1
Myo1c-206	<a href="#">ENSMUST00000136935.1</a>	351	<a href="#">80aa</a>	Protein coding	-	<a href="#">Q5ND45</a>	CDS 3' incomplete TSL:2
Myo1c-209	<a href="#">ENSMUST00000151174.7</a>	2503	No protein	Retained intron	-	-	TSL:1
Myo1c-208	<a href="#">ENSMUST00000148659.7</a>	783	No protein	Retained intron	-	-	TSL:5
Myo1c-207	<a href="#">ENSMUST00000146419.1</a>	780	No protein	Retained intron	-	-	TSL:3
Myo1c-205	<a href="#">ENSMUST00000123064.7</a>	732	No protein	Retained intron	-	-	TSL:5
Myo1c-210	<a href="#">ENSMUST00000155027.1</a>	642	No protein	Retained intron	-	-	TSL:3

The strategy is based on the design of *Myo1c-204* transcript,The transcription is shown below

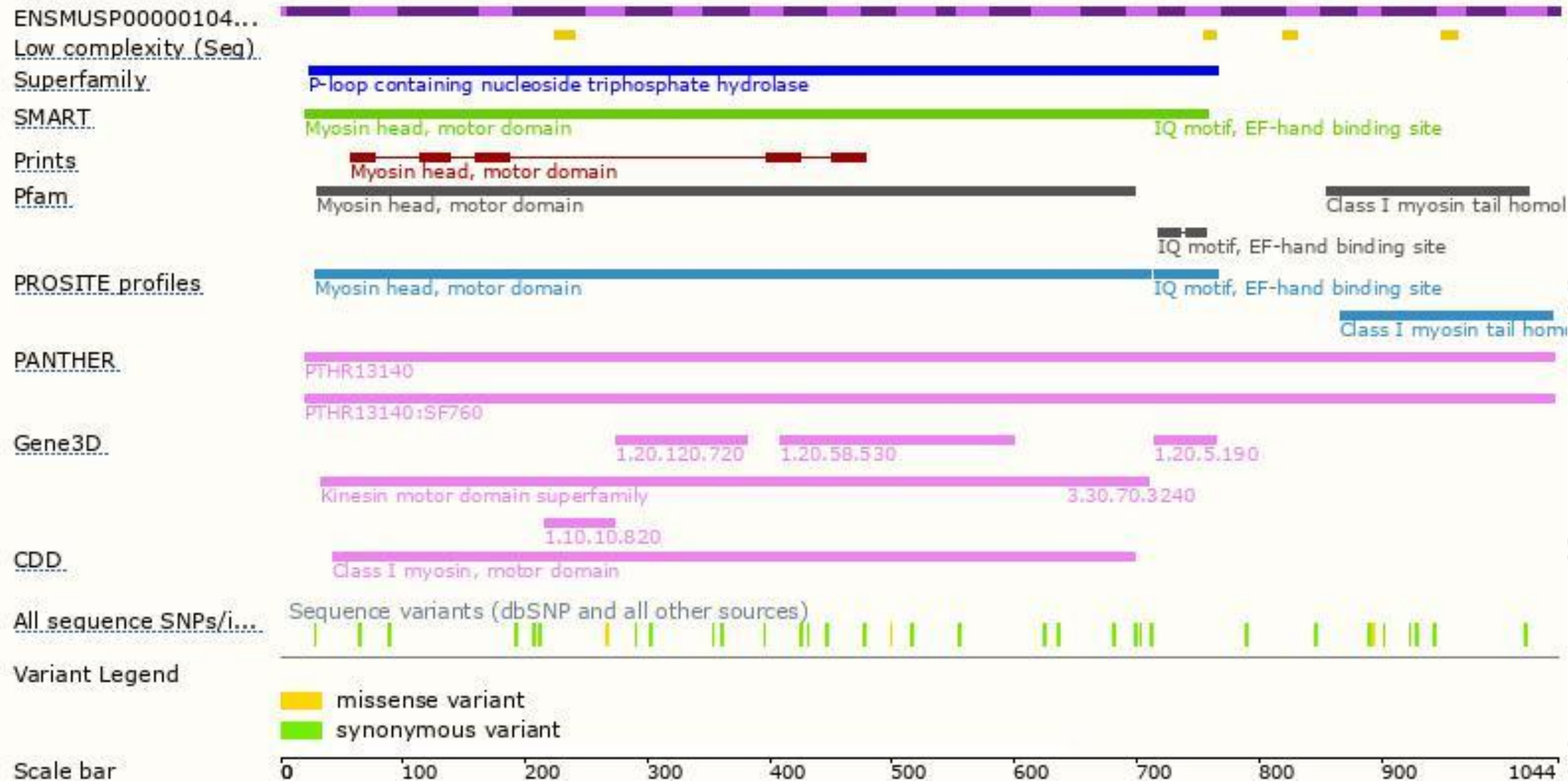


# Genomic location distribution

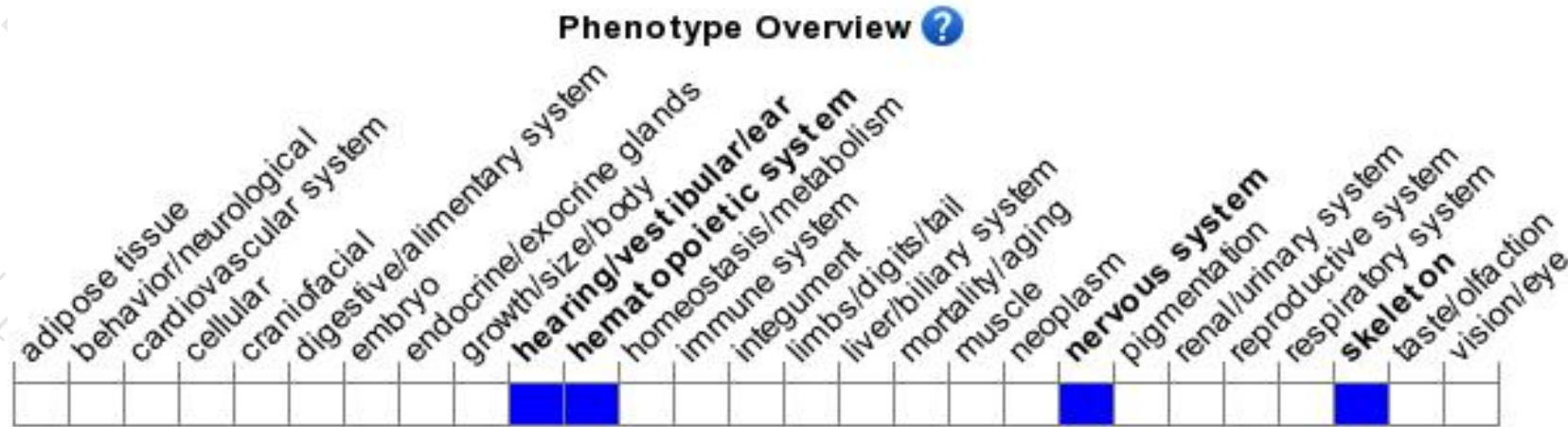




# Protein domain



# 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-in (Y61G) mutation that sensitizes to N6-modified ADP analogs display altered fast adaption in vestibular hair cells. Mice homozygous for a nuclear isoform-specific knock-out allele exhibit minor changes in bone marrow density and red blood cells.

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

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