



Trim71 Cas9-CKO Strategy

Designer:

Reviewer:

Design Date:

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2018-6-20

Project Overview

Project Name

Trim71

Project type

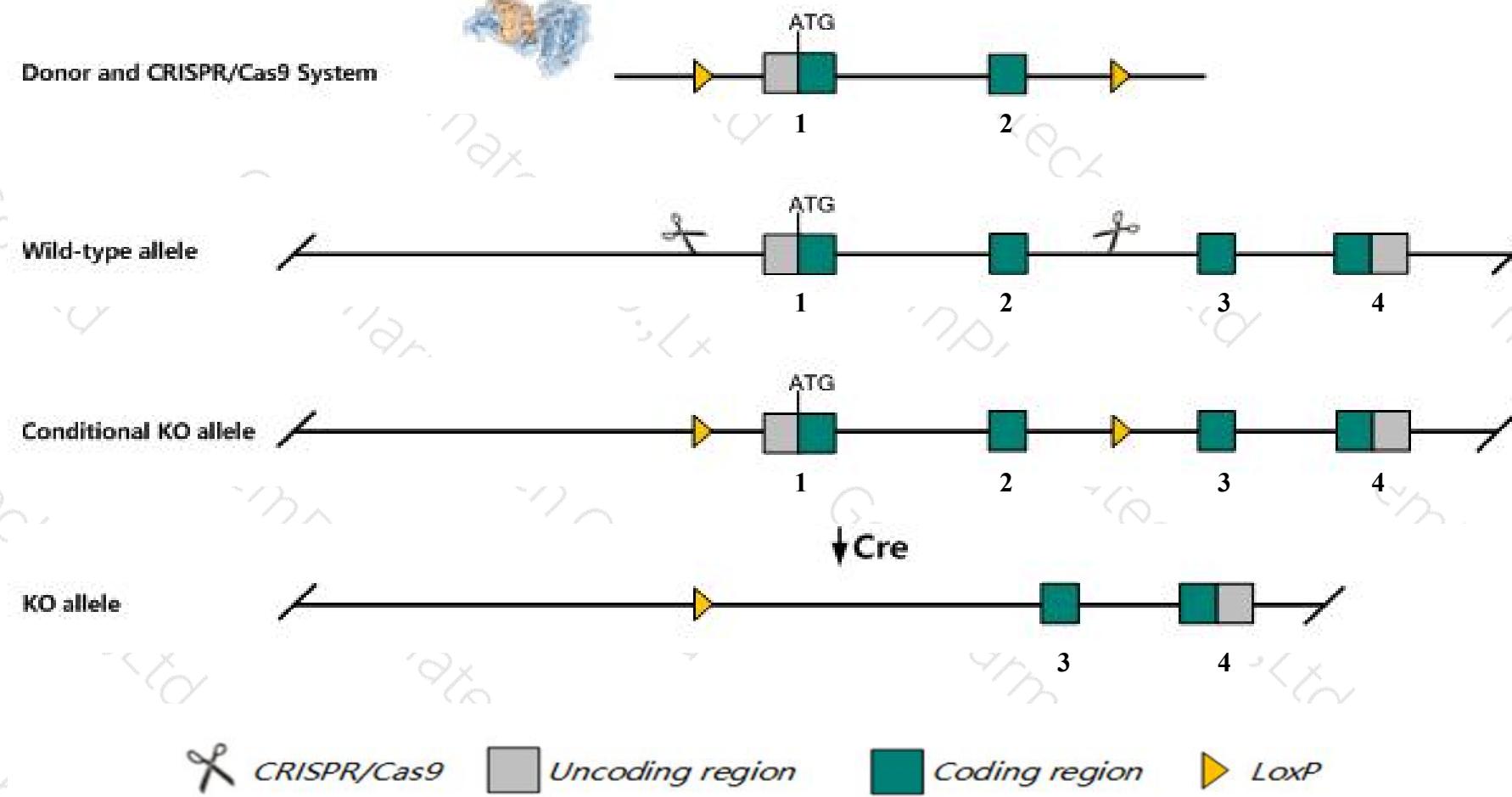
Cas9-CKO

Strain background

C57BL/6JGpt

Conditional Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Trim71* gene. The schematic diagram is as follows:



Technical routes

- The *Trim71* gene has 1 transcript. According to the structure of *Trim71* gene, exon1-exon2 of *Trim71-201* (ENSMUST00000111816.2) transcript is recommended as the knockout region. The region contains start codon ATG. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Trim71* 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.



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Notice

- According to the existing MGI data, Homozygous gene trap mutations of this gene result in failure of cranial neural tube closure and embryonic lethality. Homozygotes for a gene trap allele exhibit exencephaly, abnormal nasal process and facial prominence, reduced brain size, and embryonic or fetal lethality.
- The *Trim71* gene is located on the Chr9. 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.



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Gene information (NCBI)

Trim71 tripartite motif-containing 71 [*Mus musculus* (house mouse)]

Gene ID: 636931, updated on 12-Aug-2019

Summary



Official Symbol Trim71 provided by [MGI](#)

Official Full Name tripartite motif-containing 71 provided by [MGI](#)

Primary source [MGI:MGID:2685973](#)

See related [Ensembl:ENSMUSG00000079259](#)

Gene type protein coding

RefSeq status VALIDATED

Organism [Mus musculus](#)

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

Also known as Lin41; Gm1127; lin-41; mLin41; mlin-41; AL022943; 2610206G21Rik

Expression Biased expression in placenta adult (RPKM 2.7), CNS E11.5 (RPKM 0.9) and 4 other tissues [See more](#)

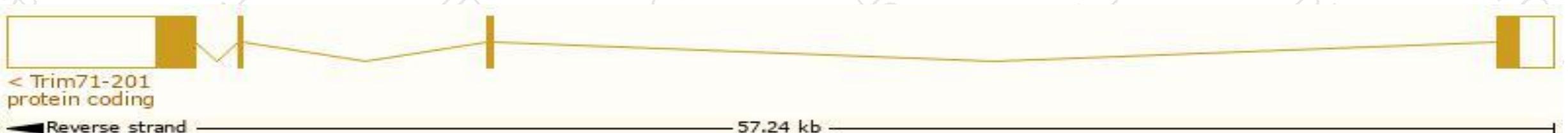
Orthologs [human](#) [all](#)

Transcript information (Ensembl)

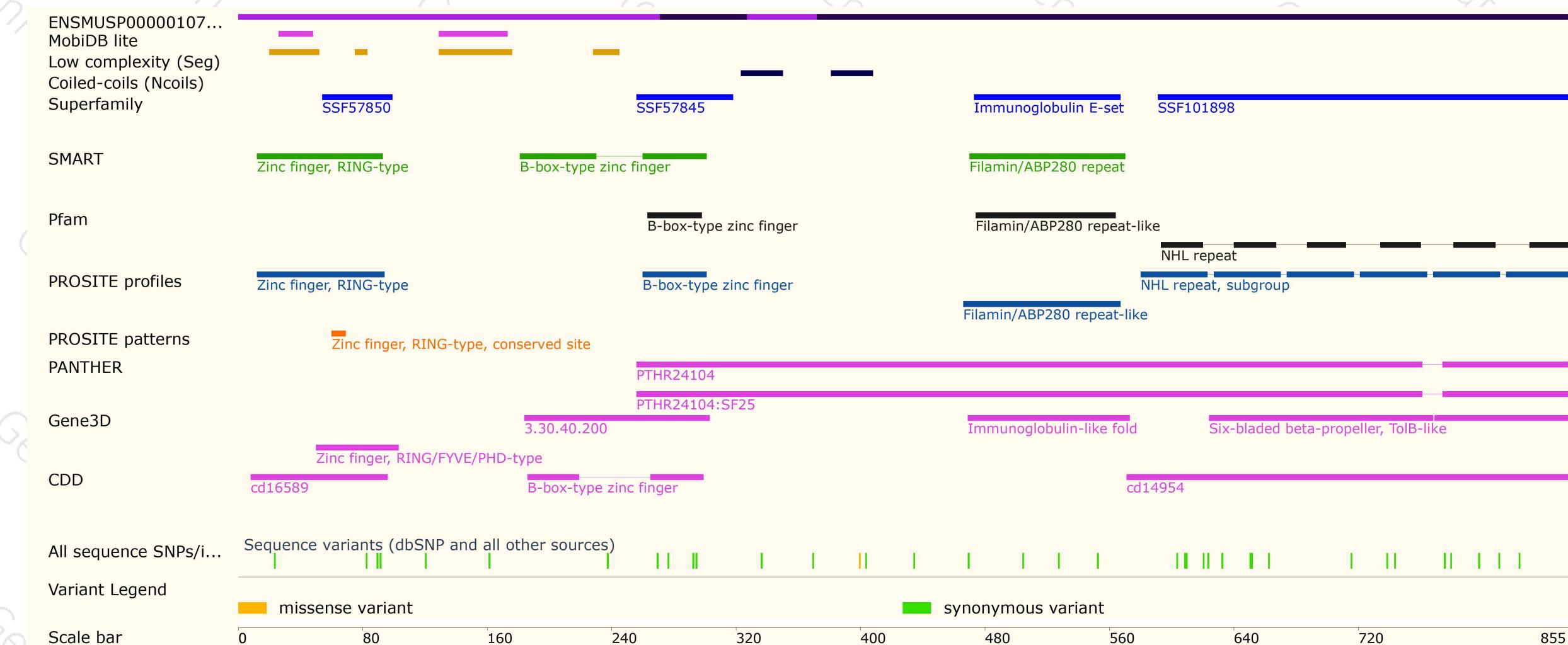
The gene has 1 transcript, and the transcript is shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Trim71-201	ENSMUST00000111816.2	9332	855aa	Protein coding	CCDS40793	Q1PSW8	TSL:1 GENCODE basic APPRIS P1

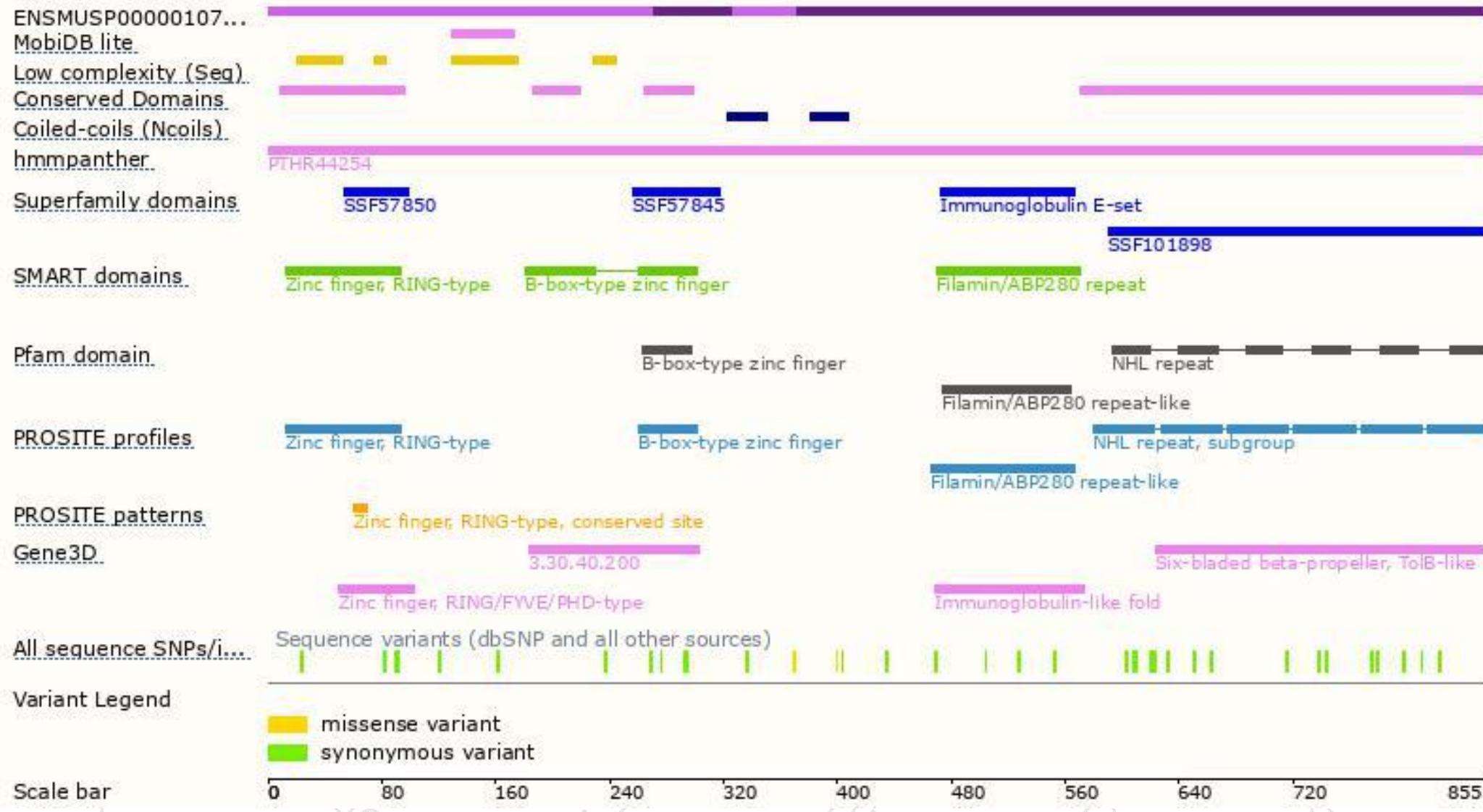
The strategy is based on the design of *Trim71-201* transcript, The transcription is shown below



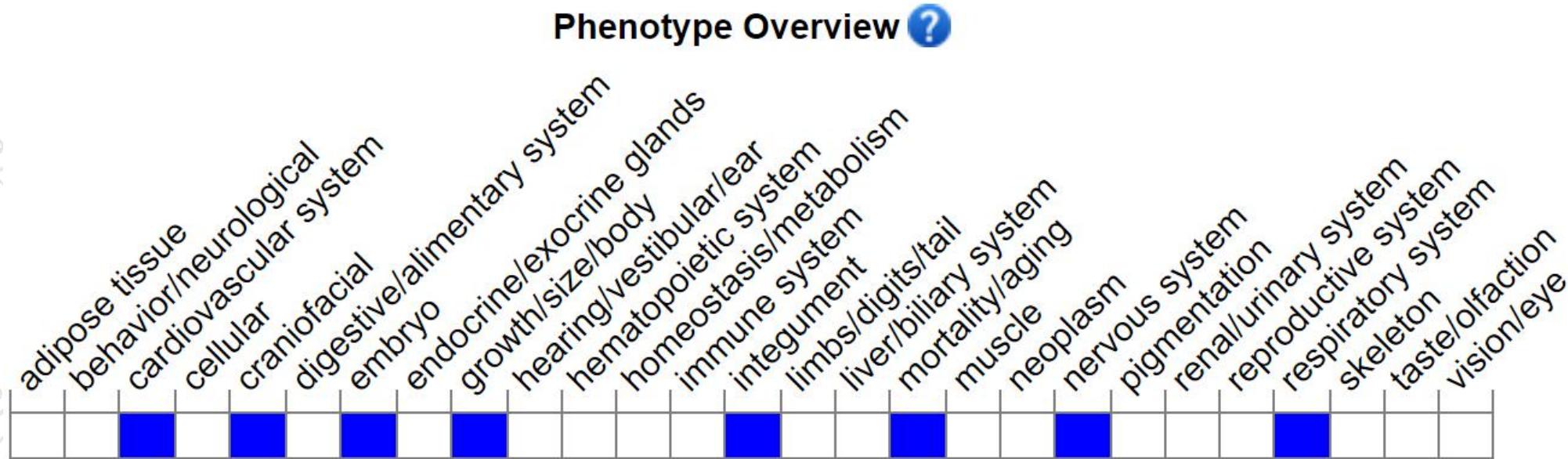
Genomic location distribution



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, Homozygous gene trap mutations of this gene result in failure of cranial neural tube closure and embryonic lethality. Homozygotes for a gene trap allele exhibit exencephaly, abnormal nasal process and facial prominence, reduced brain size, and embryonic or fetal lethality.



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

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