

***Klhl40* Cas9-KO Strategy**

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Reviewer:

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Project Overview

Project Name

Klhl40

Project type

Cas9-KO

Strain background

C57BL/6JGpt

Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Klhl40* gene. The schematic diagram is as follows:



- The *Klhl40* gene has 2 transcripts. According to the structure of *Klhl40* gene, exon1-exon5 of *Klhl40-201* (ENSMUST00000098272.3) 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 *Klhl40* gene. The brief process is as follows: CRISPR/Cas9 system

- According to the existing MGI data, Homozygous disruption of this gene results in postnatal growth retardation, abnormal sarcomere morphology, skeletal muscle dysfunction, and complete postnatal lethality. Homozygotes for a null allele develop a nemaline-like myopathy.
- *Gm47108-201* will be knocked out at the same time.
- The *Klhl40* 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 the gene knockout on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.

Gene information (NCBI)

Klhl40 kelch-like 40 [*Mus musculus* (house mouse)]

Gene ID: 72330, updated on 12-Aug-2019

Summary

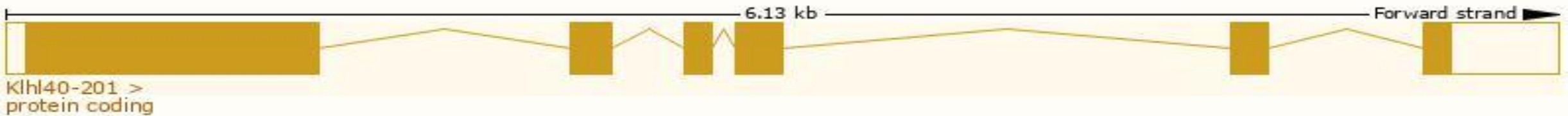
Official Symbol	Klhl40 provided by MGI
Official Full Name	kelch-like 40 provided by MGI
Primary source	MGI:MGI:1919580
See related	Ensembl:ENSMUSG000000074001
Gene type	protein coding
RefSeq status	VALIDATED
Organism	Mus musculus
Lineage	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus
Also known as	Kbtbd5; 2310024D23Rik
Expression	Biased expression in mammary gland adult (RPKM 4.8), heart adult (RPKM 3.3) and 7 other tissues See more
Orthologs	human all

Transcript information (Ensembl)

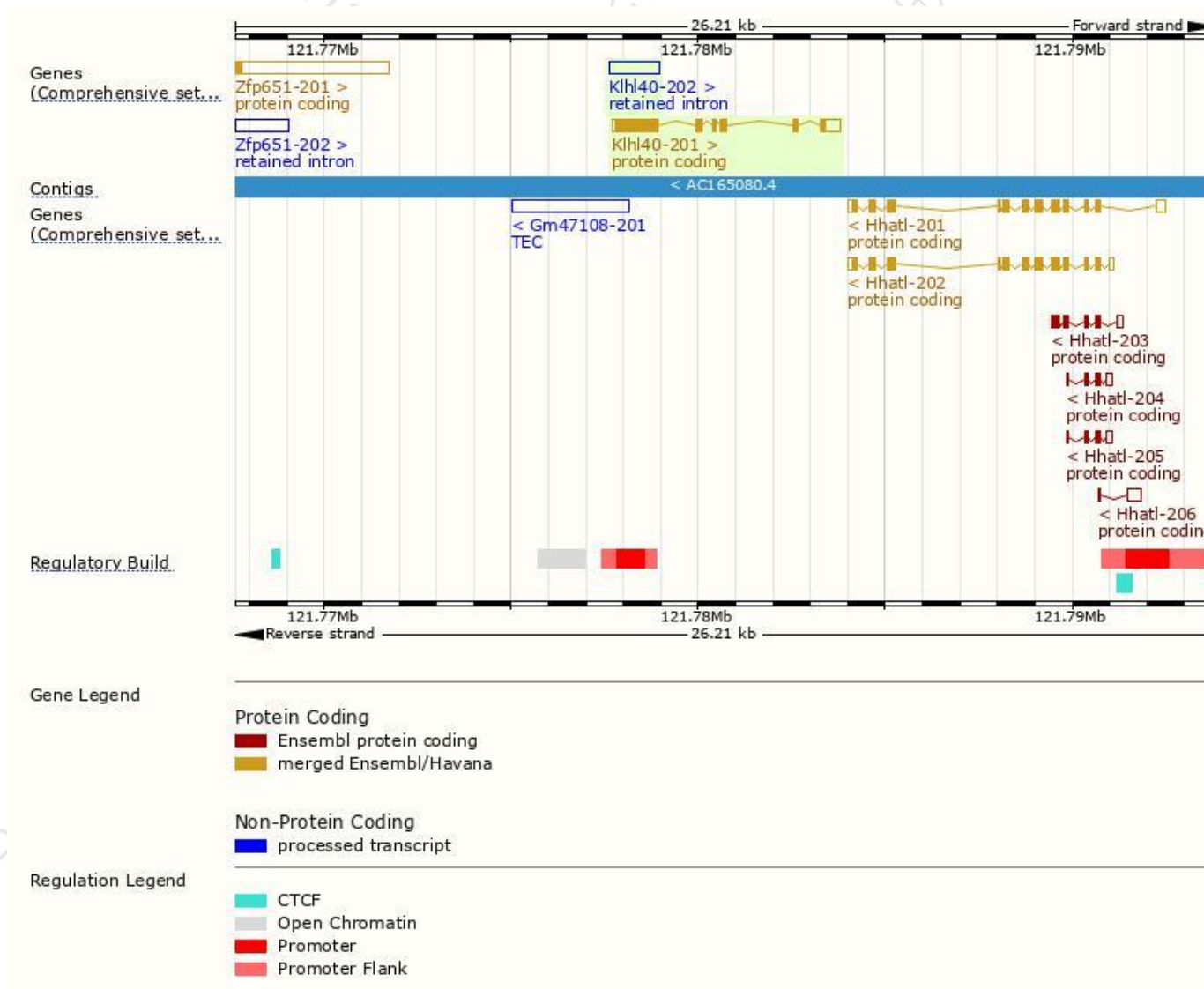
The gene has 2 transcripts,all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Klhl40-201	ENSMUST00000098272.3	2370	621aa	Protein coding	CCDS23637	A0A0R4J166	TSL:1 GENCODE basic APPRIS P1
Klhl40-202	ENSMUST00000216358.1	1368	No protein	Retained intron	-	-	TSL:NA

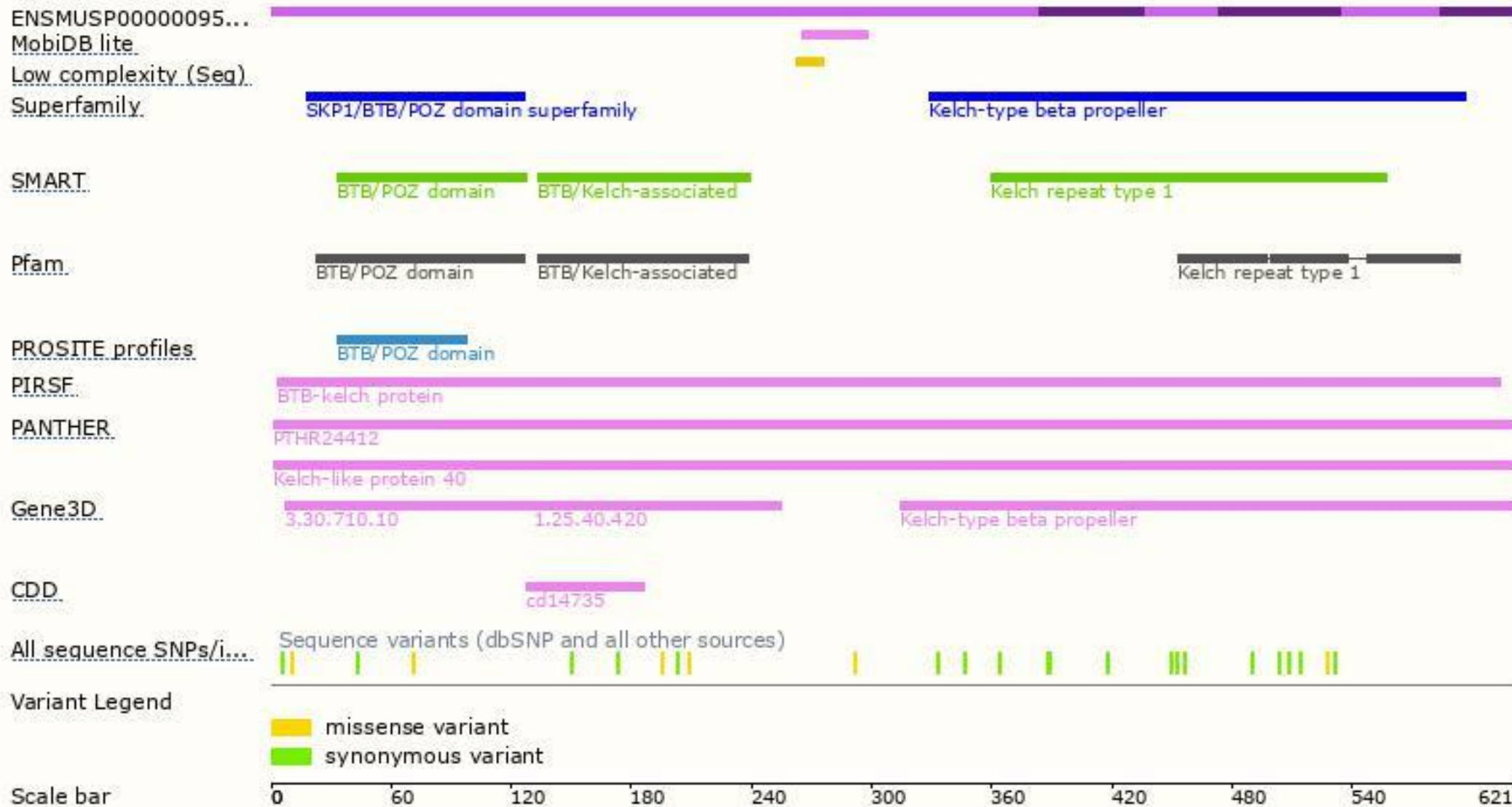
The strategy is based on the design of *Klhl40-201* transcript,The transcription is shown below



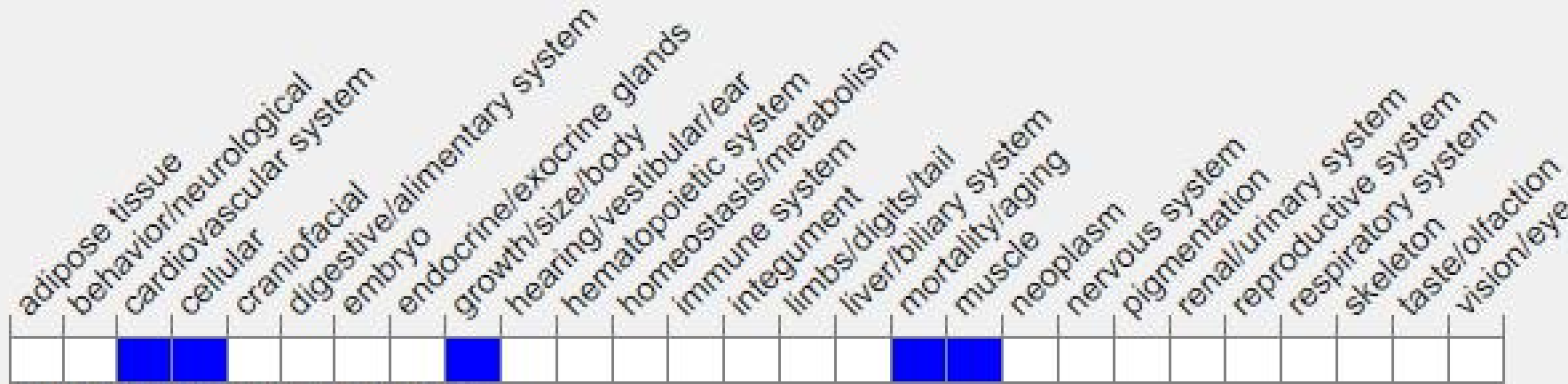
Genomic location distribution



Protein domain



集萃药康
GemPharmatech



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 disruption of this gene results in postnatal growth retardation, abnormal sarcomere morphology, skeletal muscle dysfunction, and complete postnatal lethality. Homozygotes for a null allele develop a nemaline-like myopathy.

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

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