

Dkc1 Cas9-KO Strategy

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Reviewer: JiaYu

Project Overview



Project Name

Dkc1

Project type

Cas9-KO

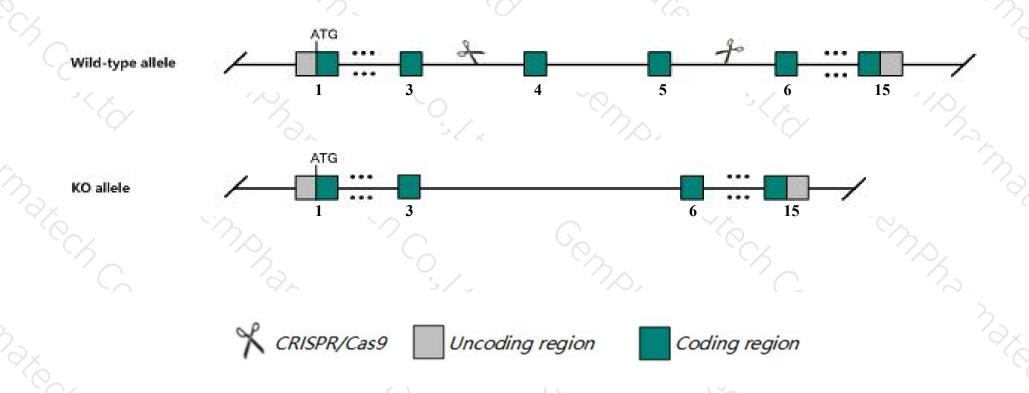
Strain background

C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Dkc1* gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Dkc1* gene has 12 transcripts. According to the structure of *Dkc1* gene, exon4-exon5 of *Dkc1-201*(ENSMUST00000033776.14) transcript is recommended as the knockout region. The region contains 277bp coding sequence Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Dkc1* gene. The brief process is as follows: CRISPR/Cas9 system

Notice



- ➤ According to the existing MGI data, Early generation male mice hemizygous for a hypomorphic allele exhibit bone marrow failure, dyskeratosis, extramedullary hematopoieis, splenomegaly, lung and kidney abnormalities, increased tumor incidence, and altered ribosome function; decreased telomere length is noted only in later generations.
- The *Dkc1* gene is located on the ChrX. 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)



Dkc1 dyskeratosis congenita 1, dyskerin [Mus musculus (house mouse)]

Gene ID: 245474, updated on 12-Aug-2019

Summary

Official Symbol Dkc1 provided by MGI

Official Full Name dyskeratosis congenita 1, dyskerin provided by MGI

Primary source MGI:MGI:1861727

See related Ensembl: ENSMUSG00000031403

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 BC068171

Expression Broad expression in liver E14 (RPKM 19.1), CNS E11.5 (RPKM 18.3) and 22 other tissues See more

Orthologs human all

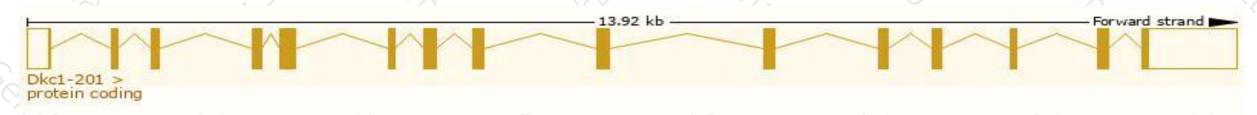
Transcript information (Ensembl)



The gene has 12 transcripts, all transcripts are shown below:

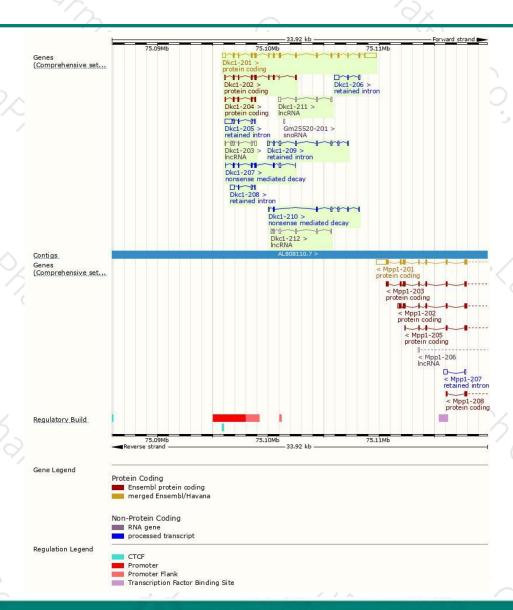
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Name A	Transcript ID 🗼	bp 🌲	Protein	Biotype	CCDS .	UniProt 4	Flags	A
Dkc1-201	ENSMUST00000033776.14	2804	<u>509aa</u>	Protein coding	CCDS41029₽	Q9ESX5₽	TSL:1 GENCODE basic	APPRIS P1
Dkc1-202	ENSMUST00000131155.7	839	260aa	Protein coding	8.70	B7ZCL7₽	CDS 3' incomplete	TSL:5
Dkc1-203	ENSMUST00000131867.7	541	No protein	IncRNA	870	-	TSL:3	
Dkc1-204	ENSMUST00000132000.7	462	145aa	Protein coding	8.70	A2AN81₽	CDS 3' incomplete	TSL:3
Dkc1-205	ENSMUST00000135389.7	898	No protein	Retained intron			TSL:3	
Dkc1-206	ENSMUST00000137286.1	610	No protein	Retained intron	8-5		TSL:2	
Dkc1-207	ENSMUST00000140609.7	754	<u>184aa</u>	Nonsense mediated decay	8.5	F6YUI5₽	CDS 5' incomplete	TSL:5
Dkc1-208	ENSMUST00000145222.1	592	No protein	Retained intron			TSL:2	
Dkc1-209	ENSMUST00000151327.7	963	No protein	Retained intron	8-5	5	TSL:5	
Dkc1-210	ENSMUST00000153844.7	657	74aa	Nonsense mediated decay	8.70	F6S1S5 €	CDS 5' incomplete	TSL:5
Dkc1-211	ENSMUST00000154422.1	551	No protein	IncRNA	8-		TSL:2	
Dkc1-212	ENSMUST00000154936.7	668	No protein	IncRNA	8.50	-	TSL:3	
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The strategy is based on the design of *Dkc1-201* transcript, The transcription is shown below



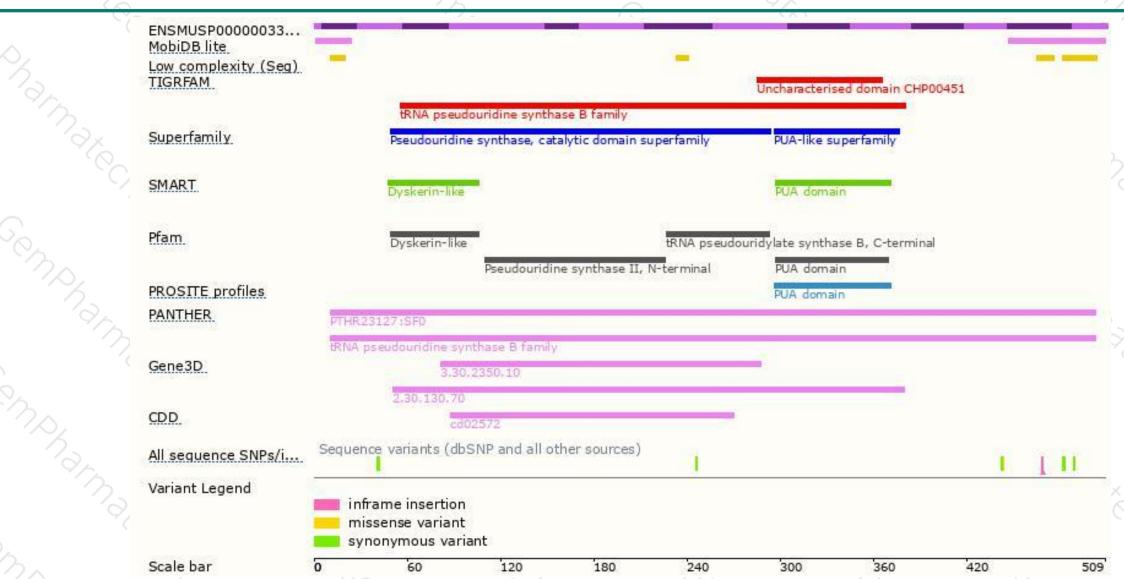
Genomic location distribution





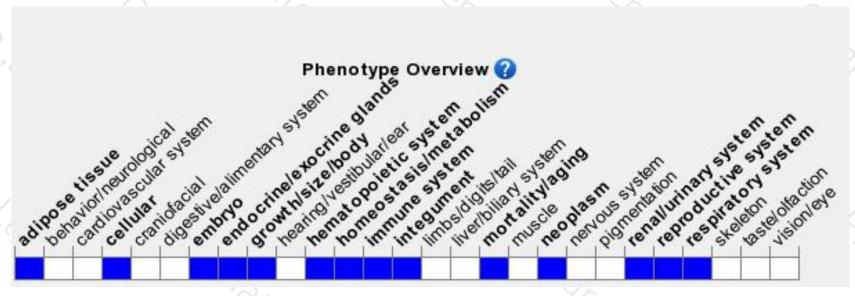
Protein domain





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, Early generation male mice hemizygous for a hypomorphic allele exhibit bone marrow failure, dyskeratosis, extramedullary hematopoieis, splenomegaly, lung and kidney abnormalities, increased tumor incidence, and altered ribosome function; decreased telomere length is noted only in later generations.



If you have any questions, you are welcome to inquire. Tel: 400-9660890





