

Zap70 Cas9-KO Strategy

Designer: Daohua Xu Reviewer: Huimin Su

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Project Overview



Project Name

Zap 70

Project type

Cas9-KO

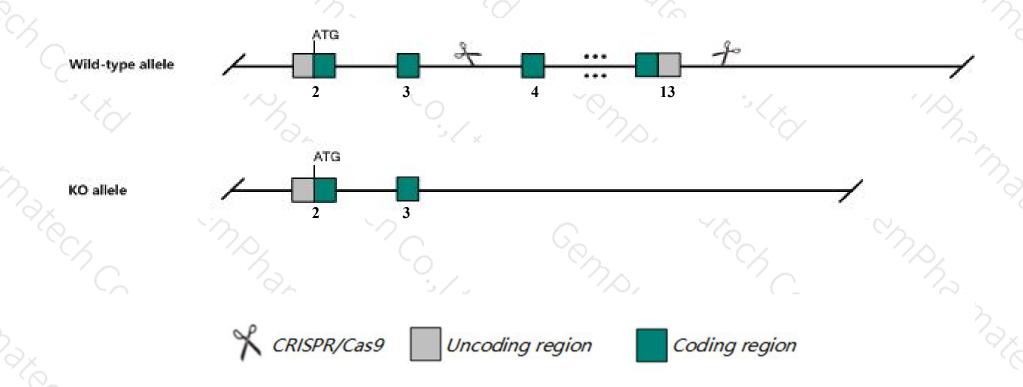
Strain background

C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Zap70 gene. The schematic diagram is as follows:



Technical routes



- ➤ The Zap70 gene has 4 transcripts. According to the structure of Zap70 gene, exon4-exon13 of Zap70-201 (ENSMUST00000027291.6) transcript is recommended as the knockout region. The region contains 1294bp coding sequence Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify Zap70 gene. The brief process is as follows: CRISPR/Cas9 system

Notice



- ➤ According to the existing MGI data, Mutant mice show T cell defects. Null mutants lack alpha-beta T cells in the thymus and have fewer T cells in dendritic and intestinal epithelium. Spontaneous and knock-in missense mutations affect T cell receptor signaling, one of the former resulting in severe chronic arthritis.
- > The Zap70 gene is located on the Chr1. 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)



Zap70 zeta-chain (TCR) associated protein kinase [Mus musculus (house mouse)]

Gene ID: 22637, updated on 5-Mar-2019

Summary

☆ ?

Official Symbol Zap70 provided by MGI

Official Full Name zeta-chain (TCR) associated protein kinase provided by MGI

Primary source MGI:MGI:99613

See related Ensembl: ENSMUSG00000026117

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 Srk, ZAP-70, mrtle, mur

Summary This gene encodes a member of the protein tyrosine kinase family. The encoded protein is essential for development of T lymphocytes and

thymocytes, and functions in the initial step of T lymphocyte receptor-mediated signal transduction. A mutation in this gene causes chronic autoimmune arthritis, similar to rheumatoid arthritis in humans. Mice lacking this gene are deficient in alpha-beta T lymphocytes in the thymus. In humans, mutations in this gene cause selective T-cell defect, a severe combined immunodeficiency disease characterized by a

selective absence of CD8-positive T lymphocytes. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Jan 2014]

Expression Biased expression in thymus adult (RPKM 70.4), spleen adult (RPKM 17.5) and 3 other tissuesSee more

Orthologs <u>human</u> all

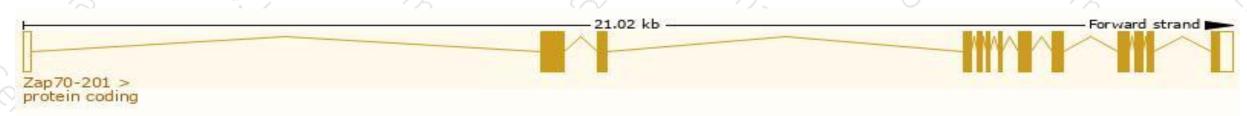
Transcript information (Ensembl)



The gene has 4 transcripts, all transcripts are shown below:

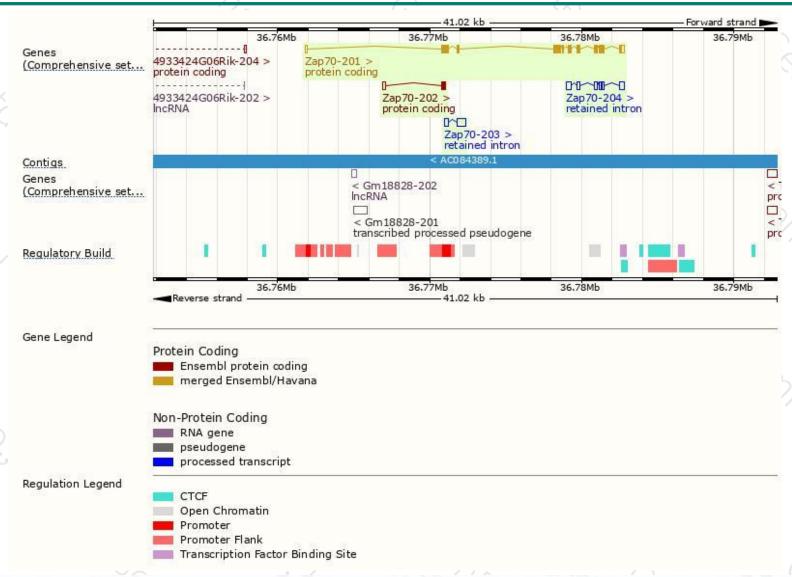
Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
ENSMUST00000027291.6	2245	618aa	Protein coding	CCDS14888	P43404	TSL:1 GENCODE basic APPRIS P1
ENSMUST00000185871.1	454	<u>85aa</u>	Protein coding	e	A0A087WQ05	CDS 3' incomplete TSL:2
ENSMUST00000190128.1	1338	No protein	Retained intron	9	20	TSL:1
ENSMUST00000186624.1	808	No protein	Retained intron		20	TSL:2
	ENSMUST000000185871.1 ENSMUST00000190128.1	ENSMUST000000185871.1 454 ENSMUST00000190128.1 1338	ENSMUST00000027291.6 2245 618aa ENSMUST00000185871.1 454 85aa ENSMUST00000190128.1 1338 No protein	ENSMUST00000027291.6 2245 618aa Protein coding ENSMUST00000185871.1 454 85aa Protein coding ENSMUST00000190128.1 1338 No protein Retained intron	ENSMUST00000027291.6 2245 618aa Protein coding CCDS14888 ENSMUST00000185871.1 454 85aa Protein coding - ENSMUST00000190128.1 1338 No protein Retained intron -	ENSMUST00000027291.6 2245 618aa Protein coding CCDS14888 P43404 ENSMUST00000185871.1 454 85aa Protein coding - A0A087WQ05 ENSMUST00000190128.1 1338 No protein Retained intron - -

The strategy is based on the design of Zap70-201 transcript, The transcription is shown below



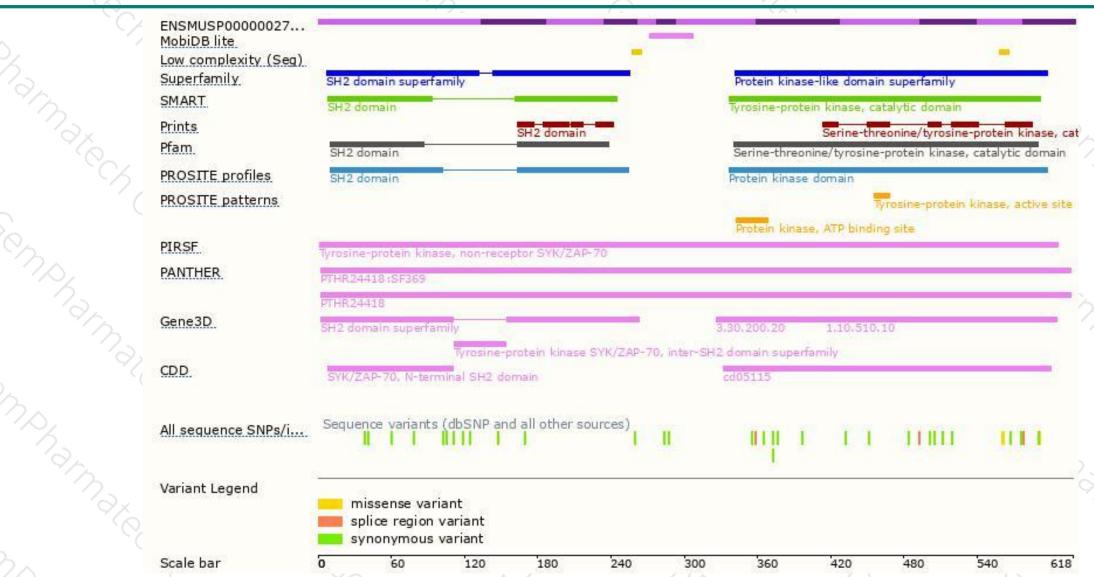
Genomic location distribution





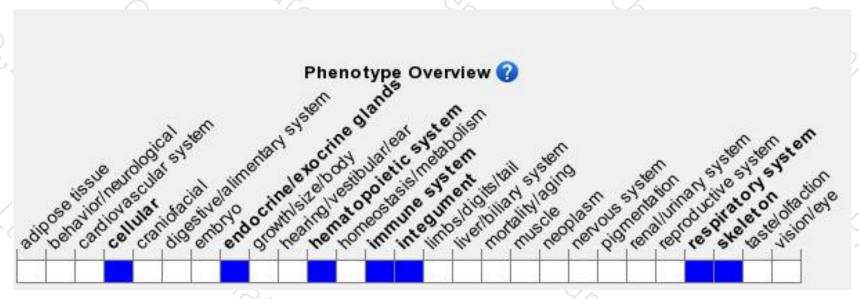
Protein domain





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, Mutant mice show T cell defects. Null mutants lack alpha-beta T cells in the thymus and have fewer T cells in dendritic and intestinal epithelium. Spontaneous and knock-in missense mutations affect T cell receptor signaling, one of the former resulting in severe chronic arthritis.



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





