

Katnal2 Cas9-CKO Strategy

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Reviewer:

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Project Overview



Project Name

Katnal2

Project type

Cas9-CKO

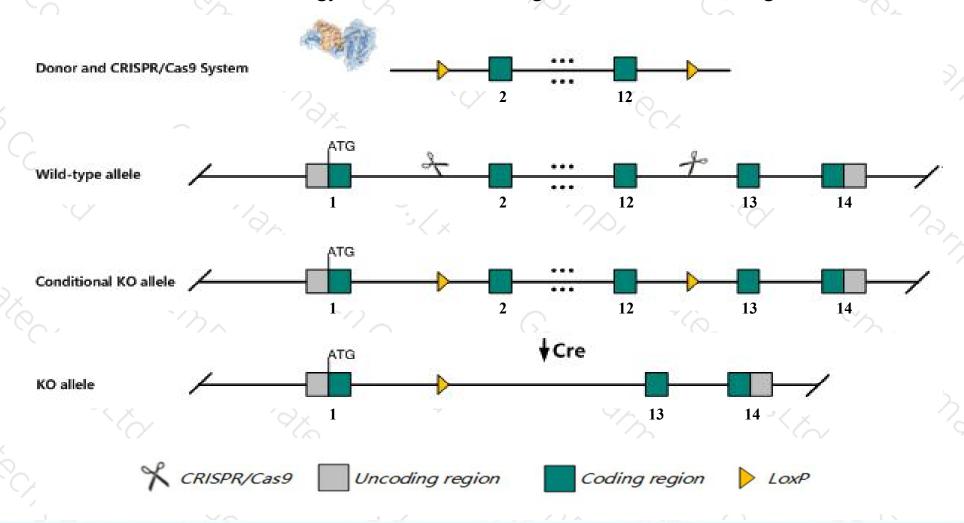
Strain background

C57BL/6JGpt

Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Katnal2* gene. The schematic diagram is as follows:



Technical routes



- The *Katnal2* gene has 11 transcripts. According to the structure of *Katnal2* gene, exon2-exon12 of *Katnal2-201* (ENSMUST00000026486.12) transcript is recommended as the knockout region. The region contains 1052bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Katnal2* 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.

Notice



- According to the existing MGI data, Homozygous null mutations in this gene result in male sterility associated with defects in several aspects of spermatogenesis, including abnormalities in the initiation of sperm tail growth from the basal body, sperm head shaping, manchette movement and dissolution, acrosome attachment to the nucleus, and sperm release via spermiation.
- The *Katnal2* gene is located on the Chr18. 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)



Katnal2 katanin p60 subunit A-like 2 [Mus musculus (house mouse)]

Gene ID: 71206, updated on 21-Aug-2019

Summary

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Official Symbol Katnal2 provided by MGI

Official Full Name katanin p60 subunit A-like 2 provided by MGI

Primary source MGI:MGI:1924234

See related Ensembl: ENSMUSG00000025420

Gene type protein coding
RefSeq status VALIDATED
Organism <u>Mus musculus</u>

Lineage Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;

Myomorpha; Muroidea; Muridae; Murinae; Mus; Mus

Also known as 3110023G01Rik; 4933439B08Rik

Expression Biased expression in testis adult (RPKM 8.9), CNS E11.5 (RPKM 2.5) and 5 other tissues See more

Orthologs human all

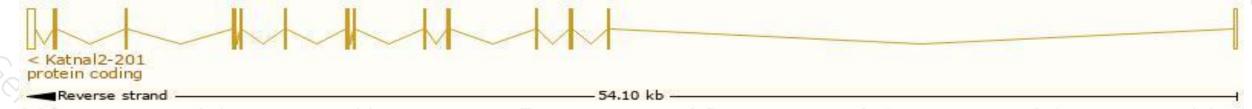
Transcript information (Ensembl)



The gene has 11 transcripts, all transcripts are shown below:

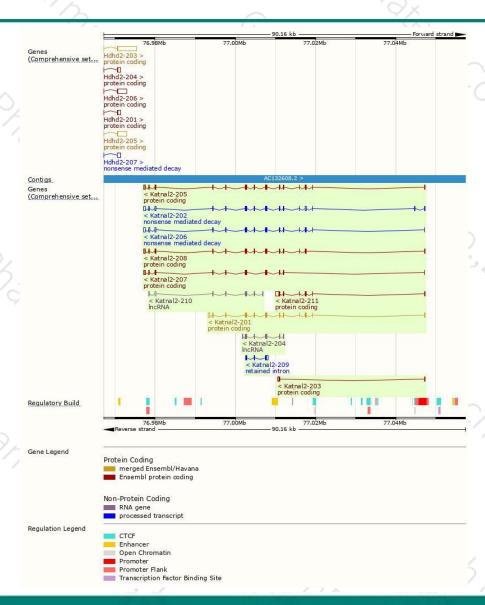
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Katnal2-201	ENSMUST00000026486.12	1674	409aa	Protein coding	CCDS37865	Q9D3R6	TSL:1 GENCODE basic
Katnal2-205	ENSMUST00000126153.7	1942	<u>539aa</u>	Protein coding	-8	Q9D3R6	TSL:1 GENCODE basic APPRIS P
Katnal2-208	ENSMUST00000137498.7	1824	495aa	Protein coding	20	D3Z4J2	TSL:5 GENCODE basic
Katnal2-211	ENSMUST00000154665.1	1515	217aa	Protein coding	29	Q9D3R6	TSL:1 GENCODE basic
Katnal2-207	ENSMUST00000137354.7	1482	277aa	Protein coding	5.6	D3Z0U5	TSL:5 GENCODE basic
Katnal2-203	ENSMUST00000123650.1	638	<u>85aa</u>	Protein coding	-83	D3Z087	TSL:1 GENCODE basic
Katnal2-202	ENSMUST00000122984.7	2039	<u>39aa</u>	Nonsense mediated decay	20	D6RI39	TSL:5
Katnal2-206	ENSMUST00000135029.7	2003	372aa	Nonsense mediated decay	29	D6RGM7	TSL:2
Katnal2-209	ENSMUST00000138336.1	668	No protein	Retained intron	50	55	TSL:2
Katnal2-210	ENSMUST00000154053.7	792	No protein	IncRNA	-	89	TSL:3
Katnal2-204	ENSMUST00000125744.7	768	No protein	IncRNA	20	84	TSL:5
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The strategy is based on the design of Katnal2-201 transcript, The transcription is shown below



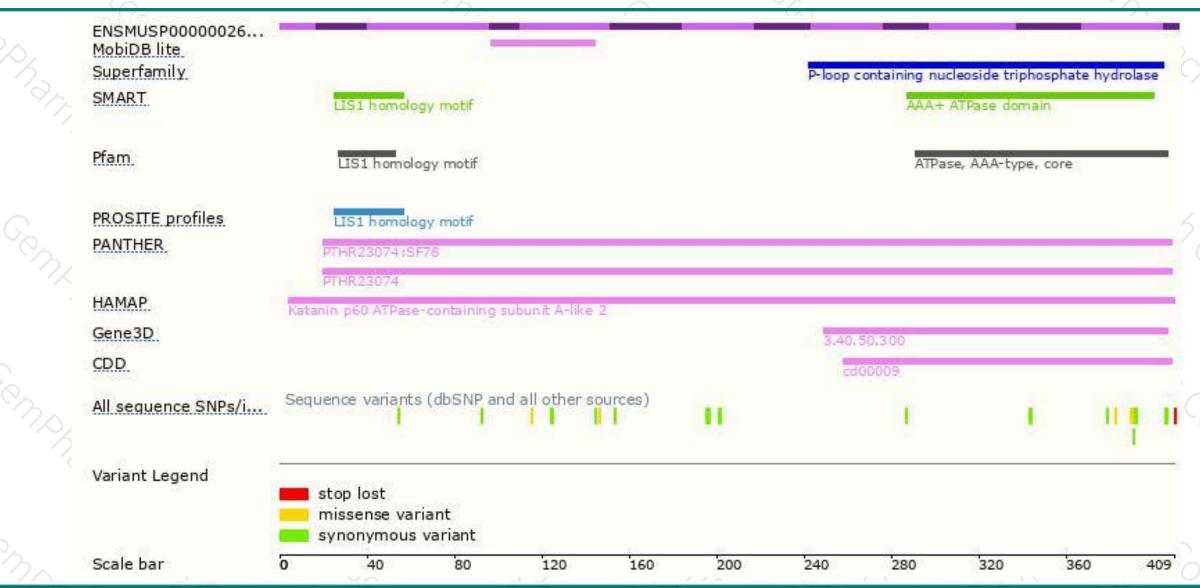
Genomic location distribution





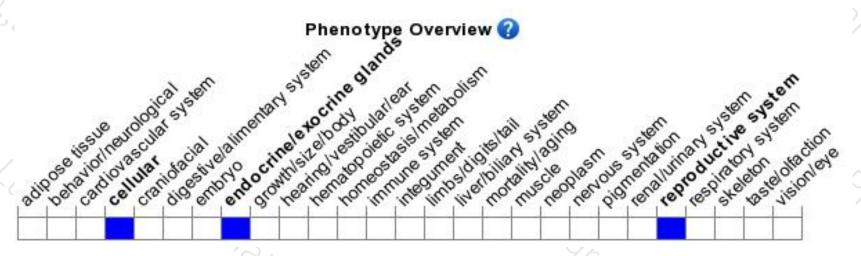
Protein domain





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 null mutations in this gene result in male sterility associated with defects in several aspects of spermatogenesis, including abnormalities in the initiation of sperm tail growth from the basal body, sperm head shaping, manchette movement and dissolution, acrosome attachment to the nucleus, and sperm release via spermiation.



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





