

Fanca Cas9-KO Strategy

Designer: Huimin Su

Reviewer: Ruirui Zhang

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Project Overview



Project Name

Fanca

Project type

Cas9-KO

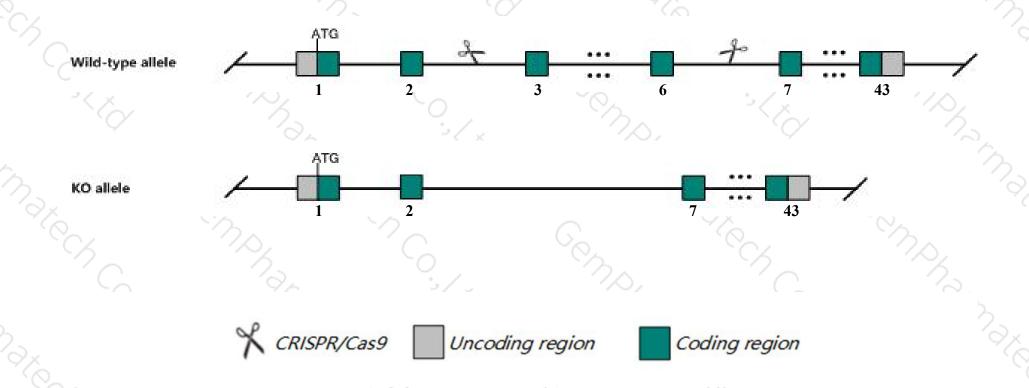
Strain background

C57BL/6JGpt

Knockout strategy



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



Technical routes



- ➤ The Fanca gene has 14 transcripts. According to the structure of Fanca gene, exon3-exon6 of Fanca-201 (ENSMUST00000035495.14) transcript is recommended as the knockout region. The region contains 398bp coding sequence Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Fanca* gene. The brief process is as follows: CRISPR/Cas9 system

Notice



- ➤ According to the existing MGI data, Mutants show variably: growth retardation, microphthalmia, craniofacial malformations and hematological changes, depending on allele and strain background. Both sexes show hypogonadism, including diminished primordial germ cells and impaired fertility.
- > Fanca is located in intron of Gm20388, so part intorn of Gm20388 will be deleted together.
- > The Fanca gene is located on the Chr8. 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)



Fanca Fanconi anemia, complementation group A [Mus musculus (house mouse)]

Gene ID: 14087, updated on 10-Oct-2019

Summary

☆ ?

Official Symbol Fanca provided by MGI

Official Full Name Fanconi anemia, complementation group A provided by MGI

Primary source MGI:MGI:1341823

See related Ensembl:ENSMUSG00000032815

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 FACA; AW208693

Expression Ubiquitous expression in genital fat pad adult (RPKM 12.2), CNS E11.5 (RPKM 9.2) and 27 other tissues See more

Orthologs <u>human</u> all

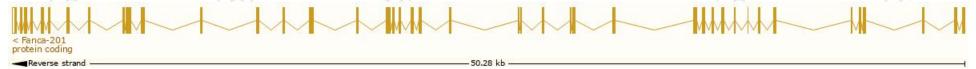
Transcript information (Ensembl)



The gene has 14 transcripts, all transcripts are shown below:

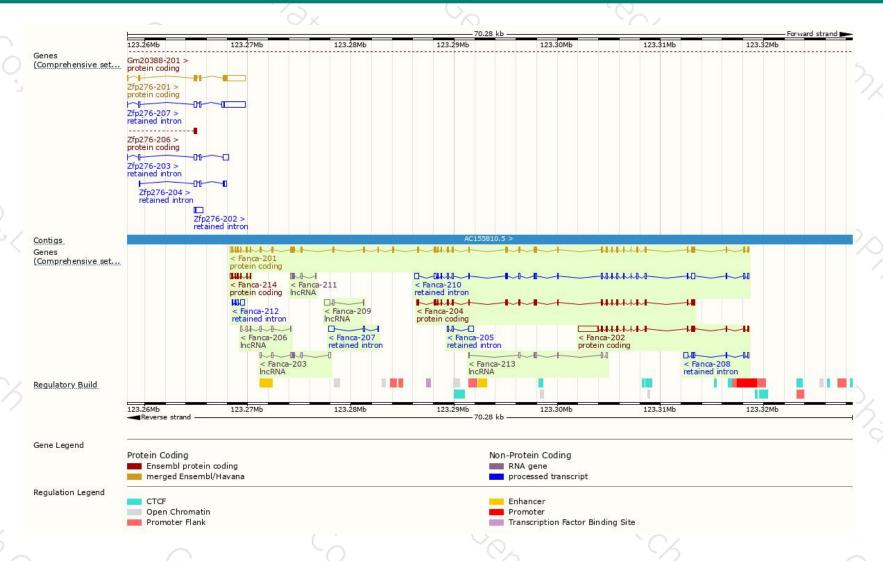
Name	Transcript ID #	bp 🛊	Protein 🍦	Biotype	CCDS	UniProt #	Flags
Fanca-201	ENSMUST00000035495.14	4485	1439aa	Protein coding	CCDS22753₽	Q9JL70 ₽	TSL:1 GENCODE basic APPRIS P1
Fanca-202	ENSMUST00000118395.1	3274	481aa	Protein coding	170	D3Z6Y5@	TSL:1 GENCODE basic
Fanca-204	ENSMUST00000127904.7	2355	<u>771aa</u>	Protein coding		F7CK17₽	CDS 5' incomplete TSL:1
Fanca-214	ENSMUST00000213090.1	650	<u>169aa</u>	Protein coding	- 170	A0A1D5RLC1₽	CDS 5' incomplete TSL:5
Fanca-210	ENSMUST00000155488.7	2994	No protein	Retained intron	-	15-	TSL:1
Fanca-208	ENSMUST00000150318.1	895	No protein	Retained intron	-	-	TSL:1
anca-205	ENSMUST00000133950.1	716	No protein	Retained intron	-	-	TSL:1
anca-212	ENSMUST00000211934.1	701	No protein	Retained intron	-	15	TSL:2
anca-207	ENSMUST00000146687.1	668	No protein	Retained intron	-	-	TSL:3
Fanca-203	ENSMUST00000126834.7	918	No protein	IncRNA	-	15-	TSL:5
anca-209	ENSMUST00000155279.1	729	No protein	IncRNA	-	-	TSL:3
anca-213	ENSMUST00000212953.1	709	No protein	IncRNA	-		TSL:5
anca-206	ENSMUST00000135702.1	667	No protein	IncRNA ■	-	157	TSL:5
Fanca-211	ENSMUST00000155510.1	483	No protein	IncRNA	-	157.0	TSL:3

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



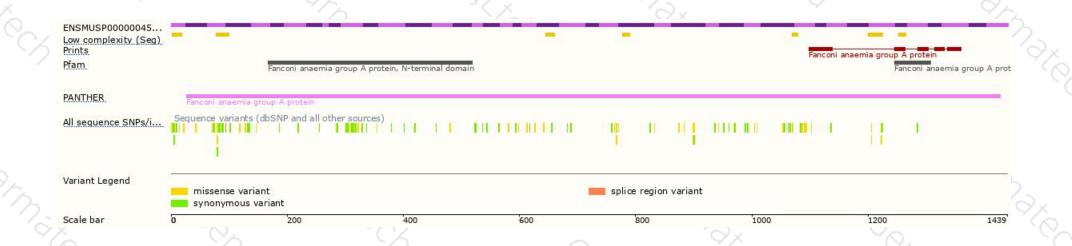
Genomic location distribution





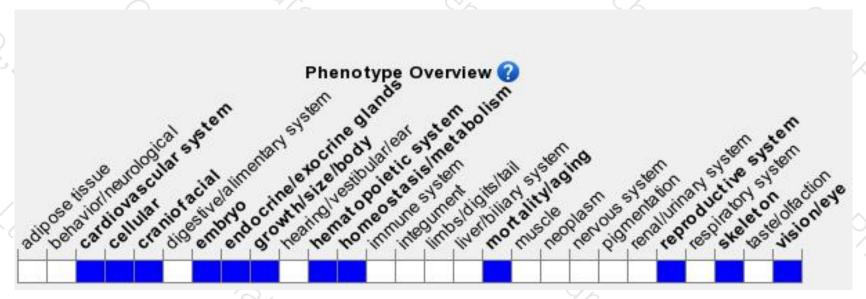
Protein domain





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, Mutants show variably: growth retardation, microphthalmia, craniofacial malformations and hematological changes, depending on allele and strain background. Both sexes show hypogonadism, including diminished primordial germ cells and impaired fertility.



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





