

# Fos Cas9-CKO Strategy

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# **Project Overview**



Project Name Fos

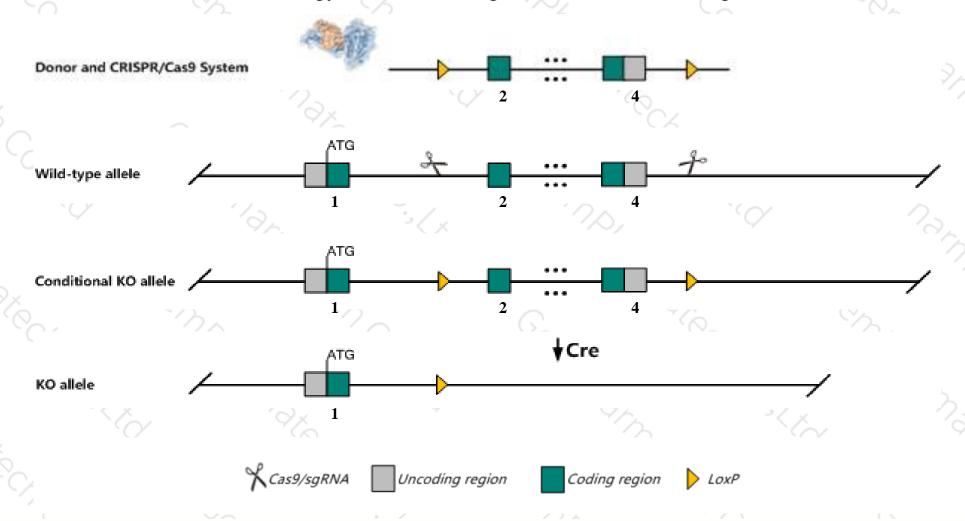
Project type Cas9-CKO

Strain background C57BL/6JGpt

# **Conditional Knockout strategy**



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



### **Technical routes**



- ➤ The Fos gene has 4 transcripts. According to the structure of Fos gene, exon2-exon4 of Fos-201 (ENSMUST00000021674.6) transcript is recommended as the knockout region. The region contains most of the coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Fos* gene. The brief process is as follows:sgRNA was transcribed in vitro, donor vector was constructed.Cas9, sgRNA 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 was 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, Null mutants are growth-retarded, most dying perinatally. Survivors have osteopetrosis and abnormal tooth eruption, gametogenesis, hemopoiesis, behavior and photoreceptor apoptosis. Hippocampal-specific mutants have seizures and highly excitable neurons.
- ➤ The *Fos* gene is located on the Chr12. 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)



#### Fos FBJ osteosarcoma oncogene [Mus musculus (house mouse)]

Gene ID: 14281, updated on 19-Mar-2019

#### Summary

☆ ?

Official Symbol Fos provided by MGI

Official Full Name FBJ osteosarcoma oncogene provided by MGI

Primary source MGI:MGI:95574

See related Ensembl:ENSMUSG00000021250

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 D12Rfj1, c-fos, cFos

Expression Biased expression in duodenum adult (RPKM 93.1), small intestine adult (RPKM 77.3) and 11 other tissuesSee more

Orthologs <u>human</u> all

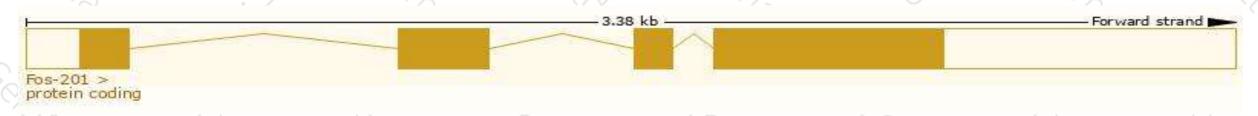
# Transcript information (Ensembl)



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

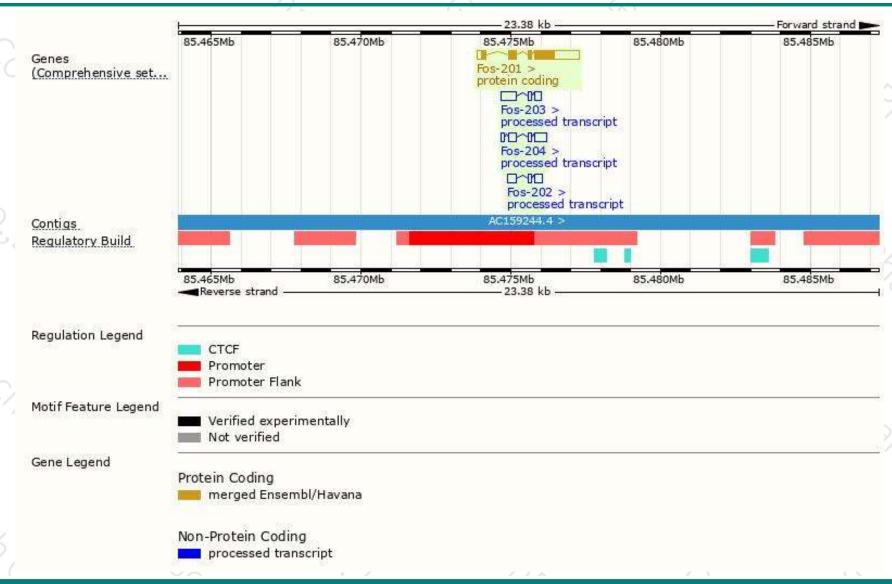
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Fos-201	ENSMUST00000021674.6	2108	<u>380aa</u>	Protein coding	CCDS26059	P01101	TSL:1 GENCODE basic APPRIS P1
Fos-204	ENSMUST00000140525.7	834	No protein	Processed transcript	-	-	TSL:5
Fos-203	ENSMUST00000136122.7	795	No protein	Processed transcript	-	-	TSL:2
Fos-202	ENSMUST00000134311.1	631	No protein	Processed transcript	-	-	TSL:2

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



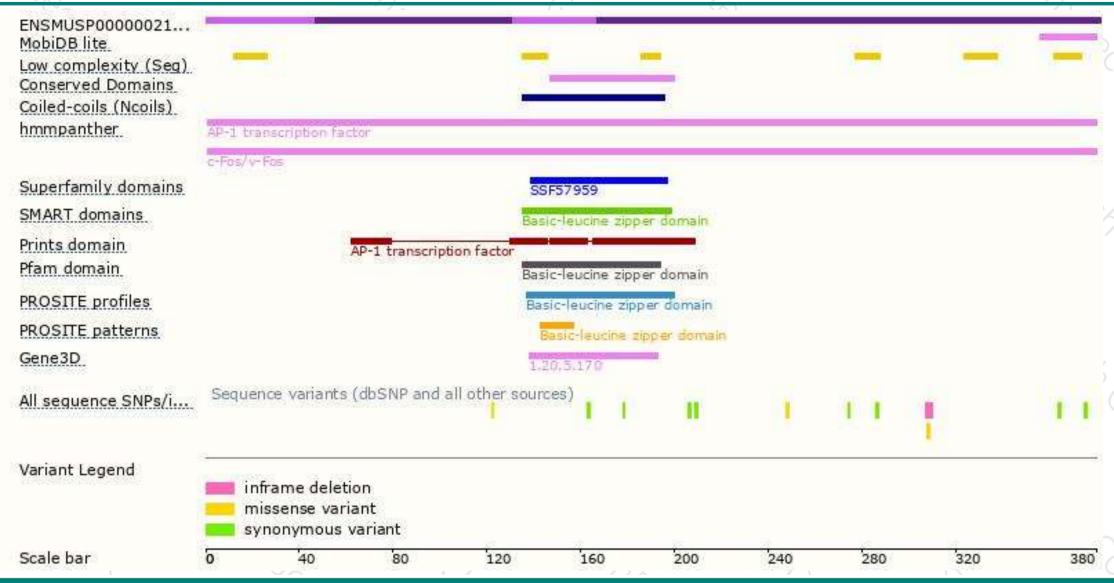
### Genomic location distribution





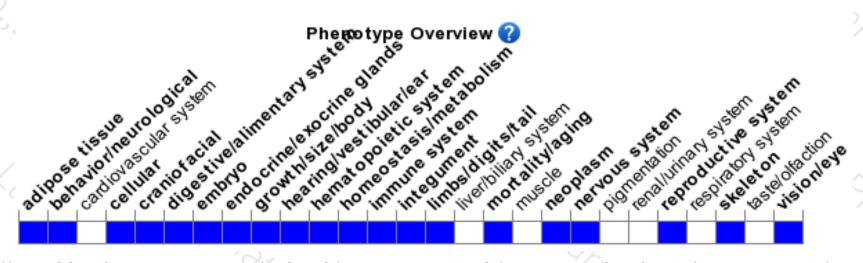
### Protein domain





# 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, Null mutants are growth-retarded, most dying perinatally. Survivors have osteopetrosis and abnormal tooth eruption, gametogenesis, hemopoiesis, behavior and photoreceptor apoptosis. Hippocampal-specific mutants have seizures and highly excitable neurons.



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

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