

# Cbfa2t3 Cas9-CKO Strategy

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Reviewer: Huimin Su

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## **Project Overview**



**Project Name** 

Cbfa2t3

**Project type** 

Cas9-CKO

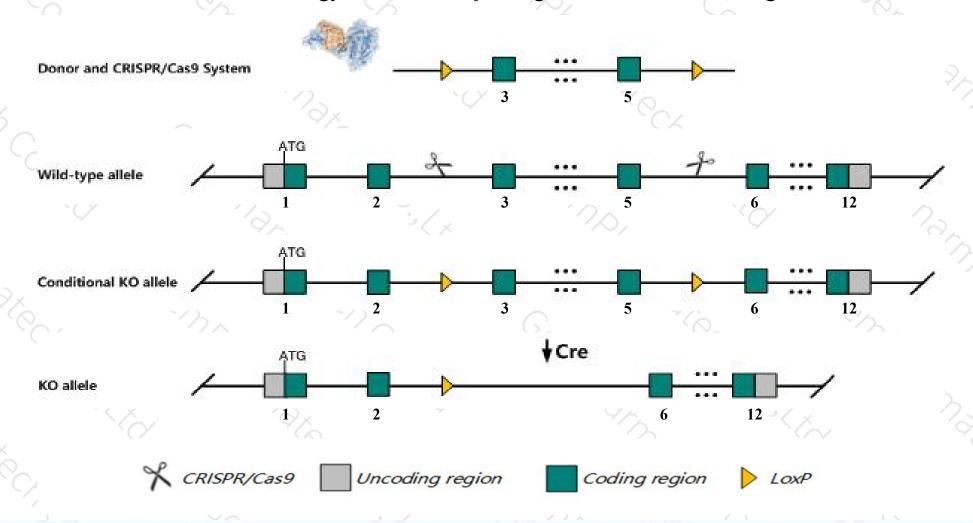
Strain background

C57BL/6JGpt

## Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Cbfa2t3 gene. The schematic diagram is as follows:



### Technical routes



- The *Cbfa2t3* gene has 7 transcripts. According to the structure of *Cbfa2t3* gene, exon3-exon5 of *Cbfa2t3-203* (ENSMUST00000127984.8) transcript is recommended as the knockout region. The region contains 410bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Cbfa2t3* 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, Mice that are homozygote null for this gene display skewing of the early myeloid progenitor cells toward the granulocytic/macrophage lineage while reducing the numbers of megakaryocyte-erythroid progenitor cells.
- > The *Cbfa2t3* 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 loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at existing technological level.

## Gene information (NCBI)



#### Cbfa2t3 CBFA2/RUNX1 translocation partner 3 [Mus musculus (house mouse)]

Gene ID: 12398, updated on 31-Jan-2019

#### Summary

↑ ?

Official Symbol Cbfa2t3 provided by MGI

Official Full Name CBFA2/RUNX1 translocation partner 3 provided by MGI

Primary source MGI:MGI:1338013

See related Ensembl: ENSMUSG00000006362

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 A630044F12Rik, Al465270, AW229127, Cbfa2t3h, ETO-2, Eto2, MTGR2

Expression Broad expression in lung adult (RPKM 8.3), thymus adult (RPKM 7.0) and 24 other tissuesSee more

Orthologs <u>human</u> all

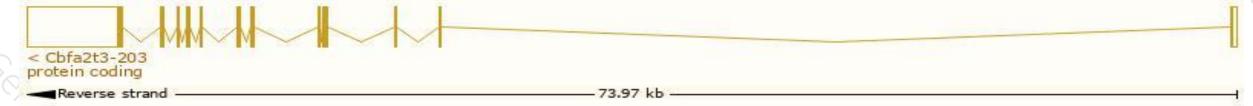
## Transcript information (Ensembl)



The gene has 7 transcripts, all transcripts are shown below:

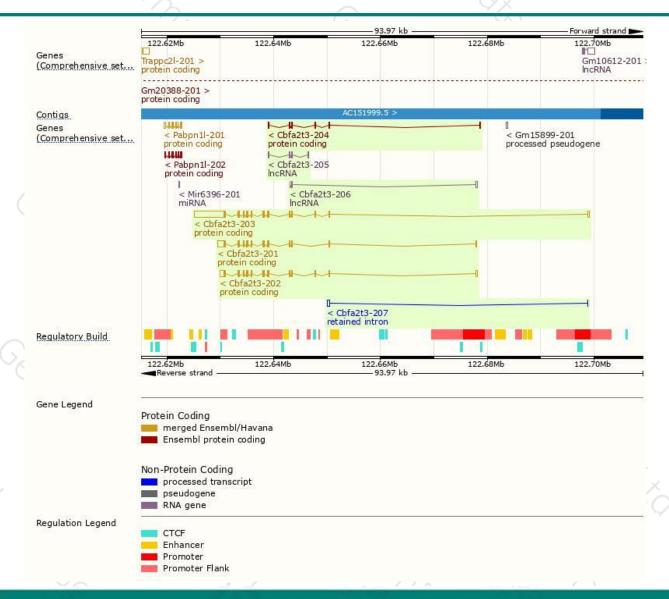
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Cbfa2t3-203	ENSMUST00000127984.8	7645	620aa	Protein coding	CCDS22744	B7ZP57	TSL:1 GENCODE basic APPRIS P3
Cbfa2t3-201	ENSMUST00000006525.13	3161	<u>559aa</u>	Protein coding	CCDS52696	<u>054972</u>	TSL:2 GENCODE basic APPRIS ALT2
Cbfa2t3-202	ENSMUST00000064674.12	2867	<u>585aa</u>	Protein coding	CCDS52697	E9QM80	TSL:1 GENCODE basic APPRIS ALT2
Cbfa2t3-204	ENSMUST00000134045.1	639	<u>185aa</u>	Protein coding	29	D3Z3P7	CDS 3' incomplete TSL:5
Cbfa2t3-207	ENSMUST00000148630.1	537	No protein	Retained intron	56		TSL:2
Cbfa2t3-206	ENSMUST00000147374.1	542	No protein	IncRNA			TSL:3
Cbfa2t3-205	ENSMUST00000142535.7	408	No protein	IncRNA	29	-	TSL:5

The strategy is based on the design of Cbfa2t3-203 transcript, The transcription is shown below



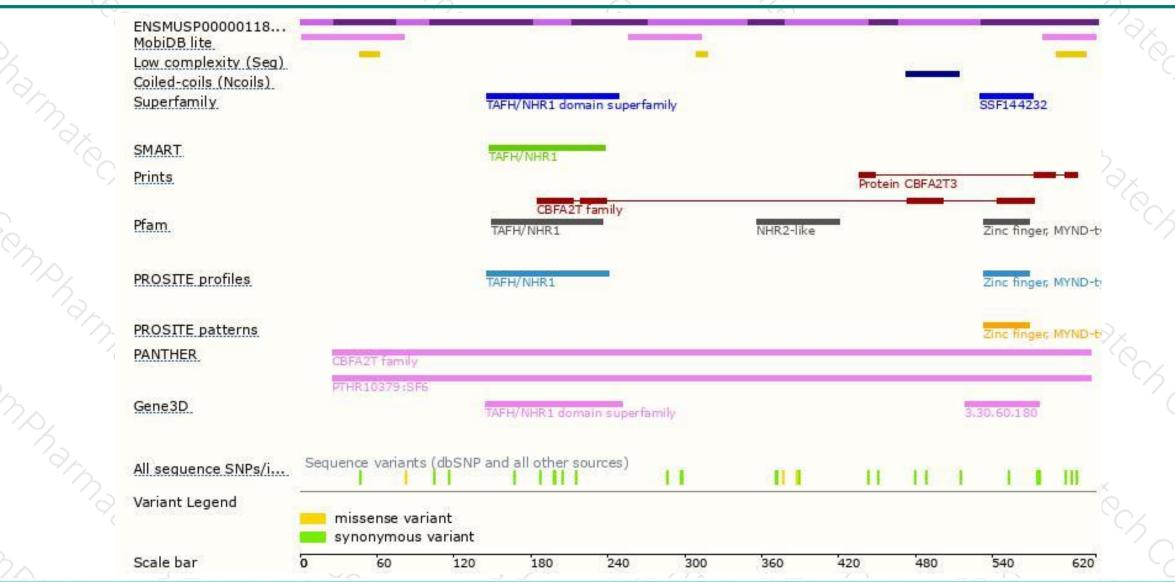
### Genomic location distribution





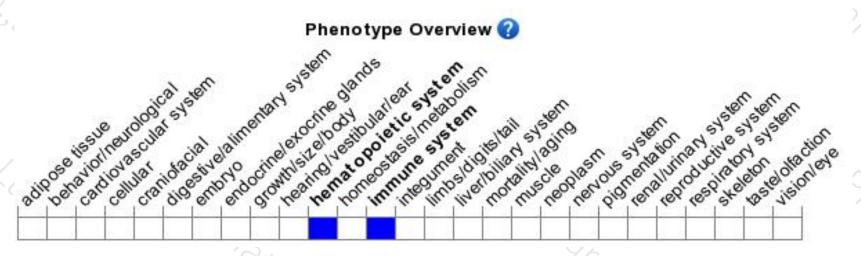
### Protein domain





## 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, Mice that are homozygote null for this gene display skewing of the early myeloid progenitor cells toward the granulocytic/macrophage lineage while reducing the numbers of megakaryocyte-erythroid progenitor cells.



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





