

# Foxd2 Cas9-CKO Strategy

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Reviewer: Xiaojing Li

**Design Date:** 2020-3-19

# **Project Overview**



**Project Name** 

Foxd2

**Project type** 

Cas9-CKO

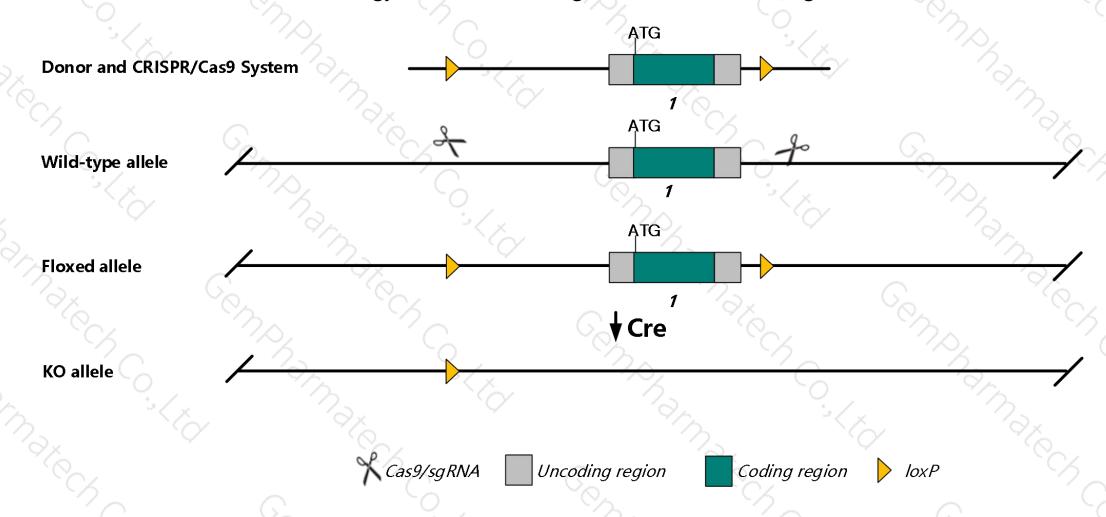
Strain background

C57BL/6JGpt

## Conditional Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Foxd2 gene. The schematic diagram is as follows:



### **Technical routes**



- The Foxd2 gene has 1 transcript. According to the structure of Foxd2 gene, exon1 of Foxd2-201 (ENSMUST00000068654.4) transcript is recommended as the knockout region. The region contains all 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 *Foxd2* 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, Homozygotes for a targeted null mutation exhibit renal abnormalities including kidney hypoplasia and hydroureter. Penetrance is reduced, and dependent upon the genetic background.
- ➤ The flox region overlap with part of the Foxd2os gene, which may affect the regulation of this gene.
- ➤ The *Foxd2* gene is located on the Chr4. 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)



#### Foxd2 forkhead box D2 [ Mus musculus (house mouse) ]

Gene ID: 17301, updated on 13-Mar-2020

Summary

2

Official Symbol Foxd2 provided by MGI

Official Full Name forkhead box D2 provided by MGI

Primary source MGI:MGI:1347471

See related Ensembl: ENSMUSG00000055210

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 Mf2; Al426778 Orthologs <u>human all</u>

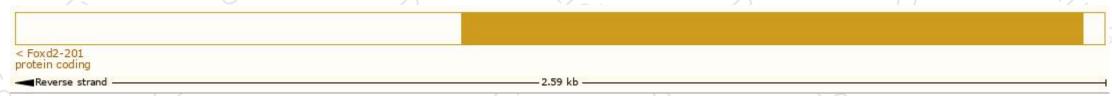
# Transcript information (Ensembl)



The gene has 1 transcript, all transcripts are shown below:

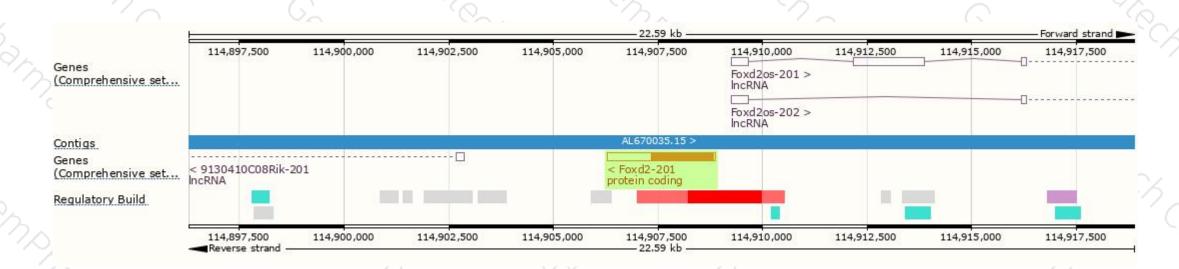
Name 🍦	Transcript ID 🍦	bp 🌲	Protein 🍦	Biotype 🍦	CCDS 🍦	UniProt 🍦	Flags		
Foxd2-201	ENSMUST00000068654.4	2592	492aa	Protein coding	CCDS18482₽	035392@	TSL:NA	GENCODE basic	APPRIS P1

The strategy is based on the design of Foxd2-201 transcript, The transcription is shown below



### Genomic location distribution





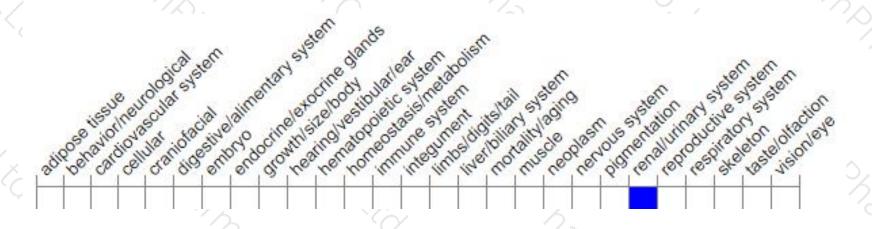
### Protein domain





# 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, Homozygotes for a targeted null mutation exhibit renal abnormalities including kidney hypoplasia and hydroureter. Penetrance is reduced, and dependent upon the genetic background.



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





