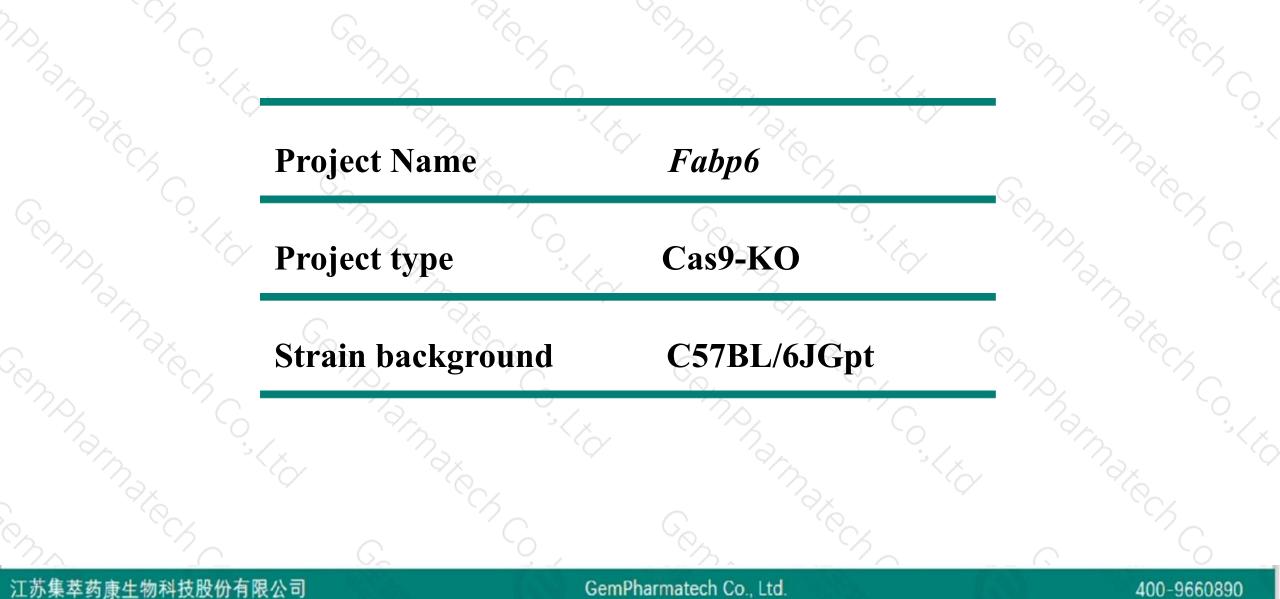


# Fabp6 Cas9-KO Strategy

Designer: Reviewer: Design Date: Yang Zeng Jing Jin 2019-11-1

### **Project Overview**

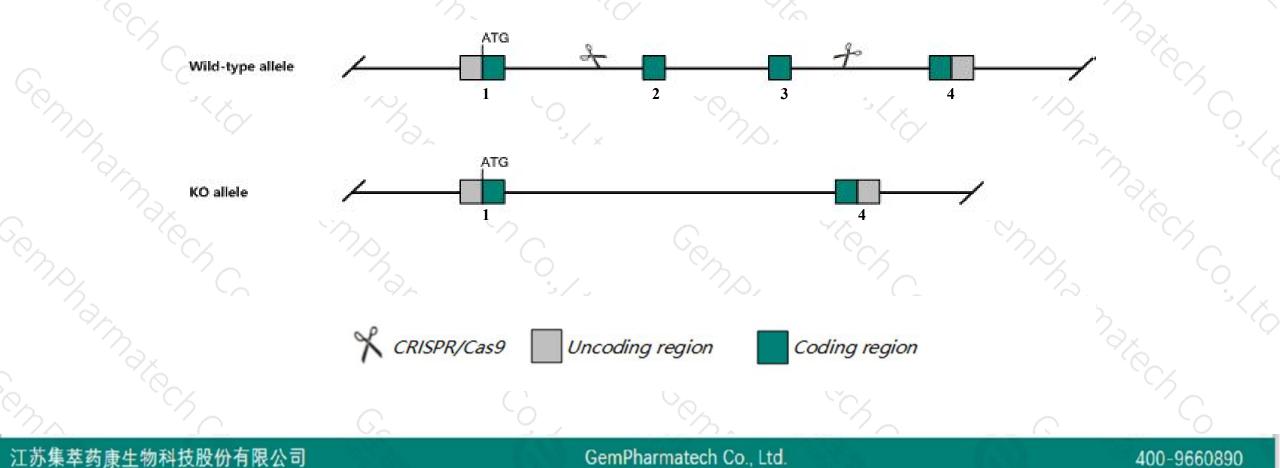




### **Knockout** strategy



This model will use CRISPR/Cas9 technology to edit the *Fabp6* gene. The schematic diagram is as follows:





- The Fabp6 gene has 1 transcript. According to the structure of Fabp6 gene, exon2-exon3 of Fabp6-201 (ENSMUST0000020672.4) transcript is recommended as the knockout region. The region contains 266bp coding sequence. Knock out the region will result in disruption of protein function.
- > In this project we use CRISPR/Cas9 technology to modify Fabp6 gene. The brief process is as follows: CRISPR/Cas9 system

- According to the existing MGI data, Mice homozygous for a knock-out allele exhibit sex-specific altered bile acid absorption and transport.
- The *Fabp6* gene is located on the Chr11. 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.

Notice

### Gene information (NCBI)



☆ ?

Fabp6 fatty acid binding protein 6 [ Mus musculus (house mouse) ]

43474306

C1qtnf2

Cenil

Gene ID: 16204, updated on 12-Aug-2019

Summary

Official Symbol Fabp6 provided by MGI Official Full Name fatty acid binding protein 6 provided by MGI Primary source MGI:MGI:96565 See related Ensembl:ENSMUSG0000020405 Gene type protein coding RefSeq status REVIEWED Organism Mus musculus Lineage Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha; Muroidea: Muridae: Murinae: Mus: Mus GT; ILBP; I-15P; ILBP3; IIIbp; I-BABP Also known as The protein encoded by this gene is part of the fatty acid binding protein family (FABP). FABPs are a family of small, highly conserved, Summary cytoplasmic proteins that bind long-chain fatty acids and other hydrophobic ligands and participate in fatty acid uptake, transport, and metabolism. This protein functions within the ileum, the distal 25-30% of the small intestine, and plays a role in enterohepatic circulation of bile acids and cholesterol homeostasis. In humans, it has been reported that polymorphisms in FABP6 confer a protective effect in obese individuals from developing type 2 diabetes. In mice deficiency of this gene affects bile acid metabolism in a gender-specific manner and was reported to be required for efficient apical to basolateral transport of conjugated bile acids. [provided by RefSeq, Jan 2013] Restricted expression toward large intestine adult (RPKM 1460.5) See more Expression Orthologs human all

Chromosome 11 - NC\_000077.6

Fabp6 + Gm12149

江苏集萃药康生物科技股份有限公司

#### GemPharmatech Co., Ltd.

Gm34963

43722633

400-9660890

## **Transcript information (Ensembl)**



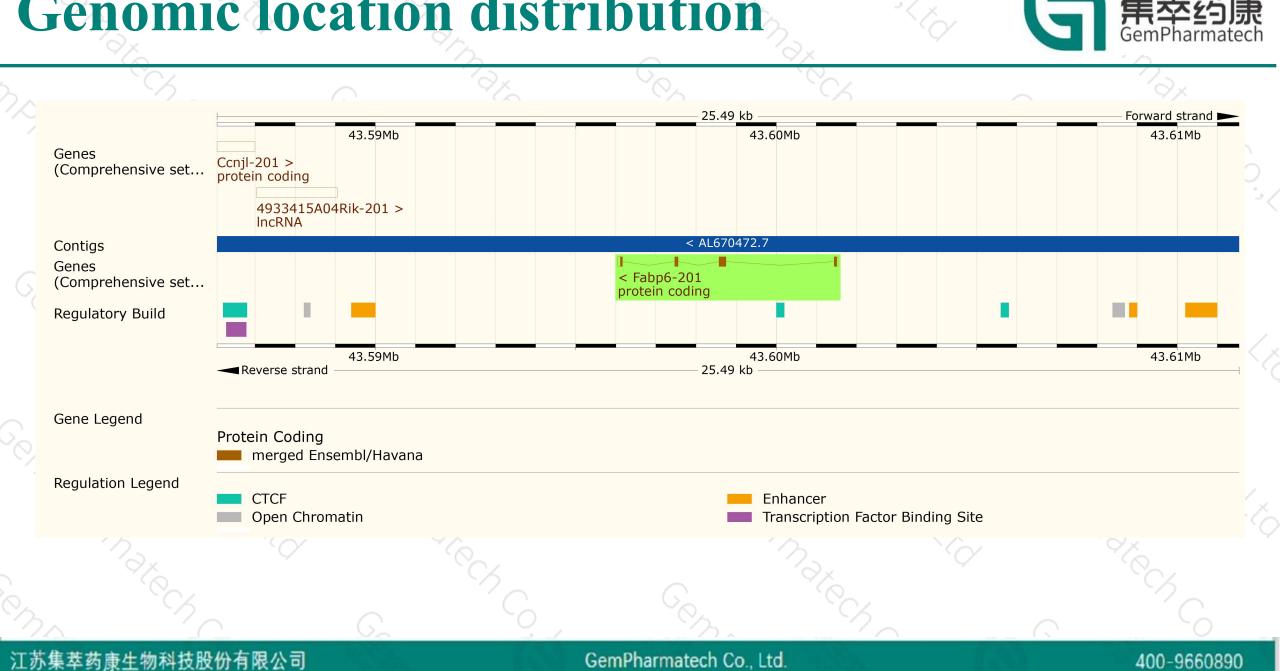
The gene has 1 transcript, and the transcript is shown below:

Name 💧	Transcript ID 💧	bp 🍦	Protein 🖕	Translation ID	Biotype 💧	CCDS 🖕	UniProt 🖕		Flags	4
Fabp6-201	ENSMUST0000020672.4	479	<u>128aa</u>	ENSMUSP00000020672.4	Protein coding	<u>CCDS24560</u> &	<u>P51162</u> &	TSL:1	GENCODE basic	APPRIS P1

The strategy is based on the design of *Fabp6-201* transcript, The transcription is shown below

< Fabp6-201 protein coding Reverse strand		5.4	19 kb			
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### **Genomic location distribution**



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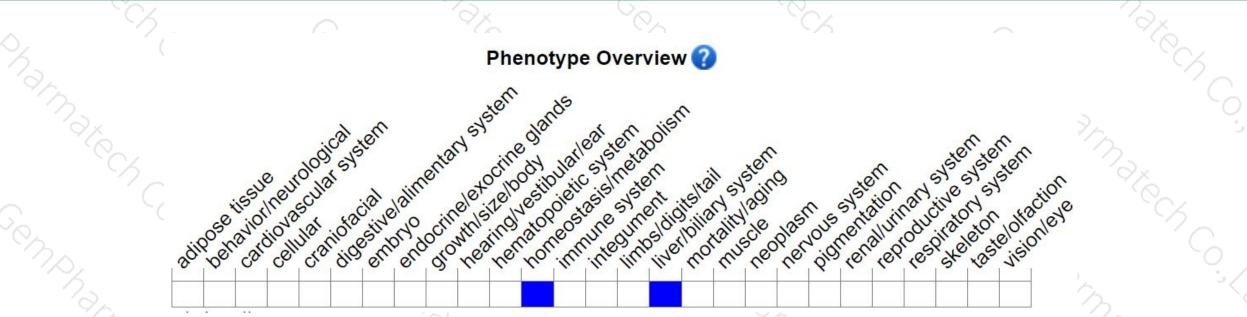
### **Protein domain**



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ENSMUSP00000020 Superfamily	Caluain							
Prints	Calycin Cytosolic fatty-acid b	inding					\	
Pfam	PF14651	inding					<u> </u>	
PROSITE patterns	Cytosolic fatty-ad	id binding						
PANTHER	PTHR11955:SF69	-					245°	
	Intracellular lipid bindi	ng protein						
Gene3D	Calycin							
All sequence SNPs/i	Sequence variants (db	SNP and all other sources)					R R	
Variant Legend	stop lost		splice re	gion variant	syno	synonymous variant		
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### 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 homozygous for a knock-out allele exhibit sex-specific altered bile acid absorption and transport.



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



