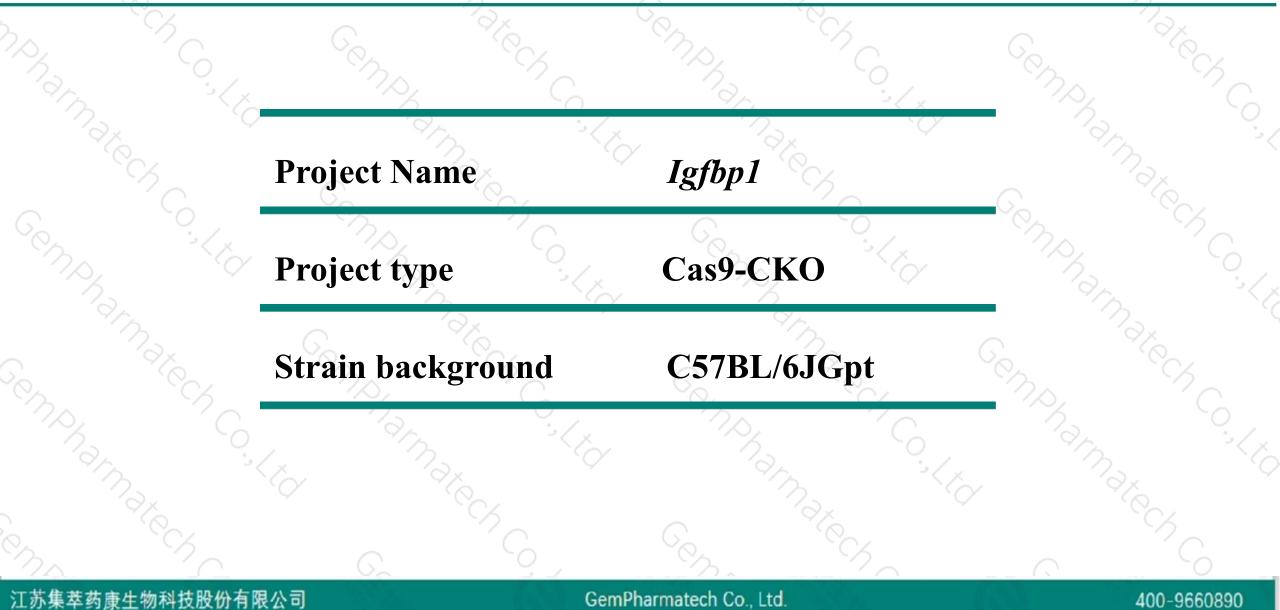


Igfbp1 Cas9-CKO Strategy

Designer: Reviewer: Design Date: Yang Zeng Jing Jin 2019-10-31

Project Overview



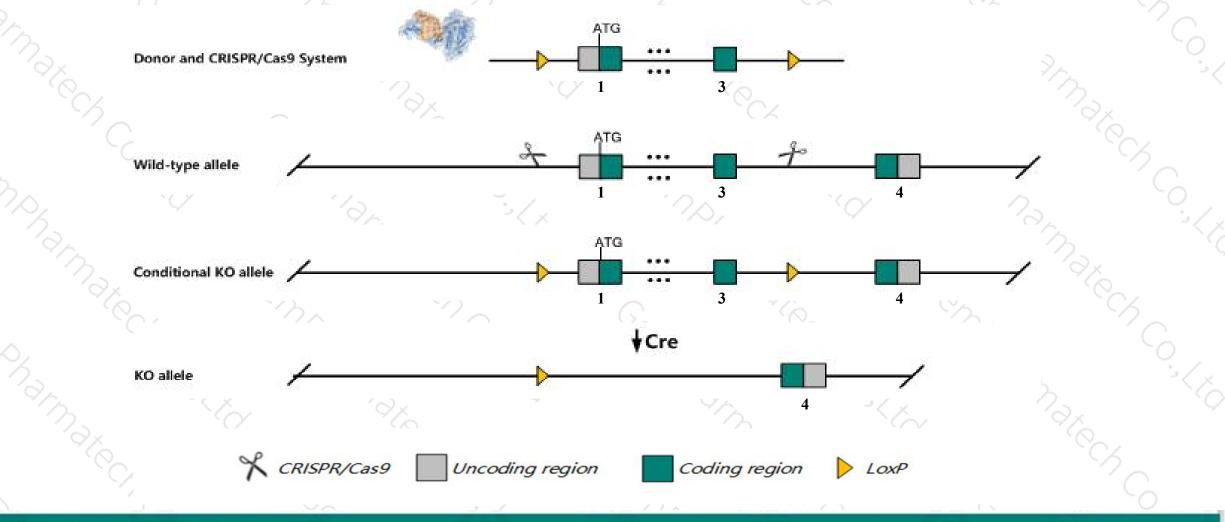


Conditional Knockout strategy



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This model will use CRISPR/Cas9 technology to edit the *Igfbp1* gene. The schematic diagram is as follows:



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The *Igfbp1* gene has 1 transcript. According to the structure of *Igfbp1* gene, exon1-exon3 of *Igfbp1-201* (ENSMUST0000020704.7) transcript is recommended as the knockout region. The region contains start codon ATG. Knock out the region will result in disruption of protein function.

In this project we use CRISPR/Cas9 technology to modify *Igfbp1* 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.

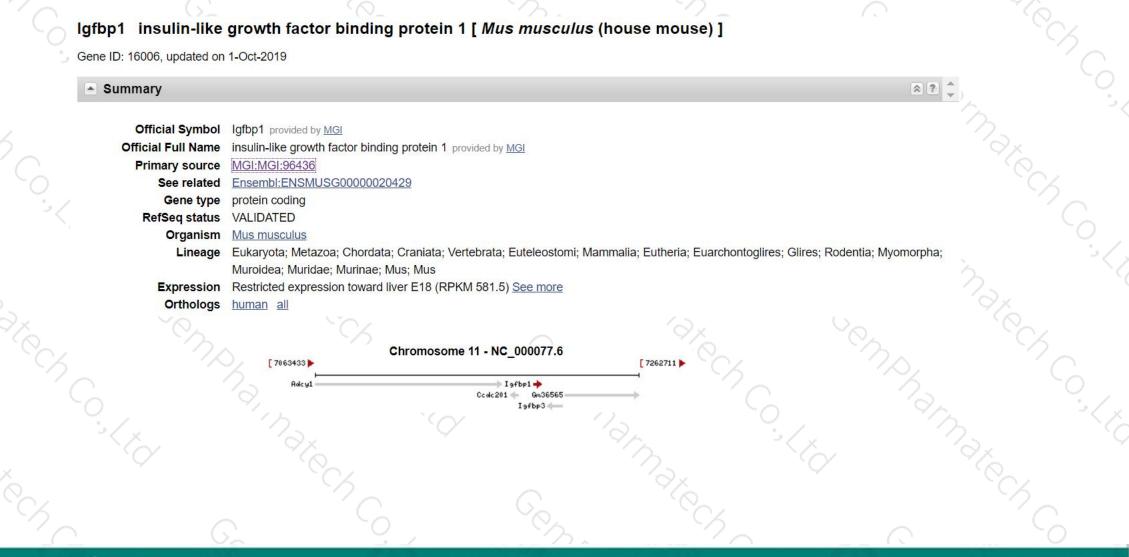
The flox mice will be knocked out after mating with mice expressing Cre recombinase, resulting in the loss of function of the target gene in specific tissues and cell types.



- According to the existing MGI data, Mice homozygous for disruptions in this gene desplay a grossly normal phenotype but are more susceptible to liver injury.
- The *Igfbp1* 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 loxp insertion on gene transcription, RNA splicing and protein translation cannot be predicted at existing technological level.

Gene information (NCBI)





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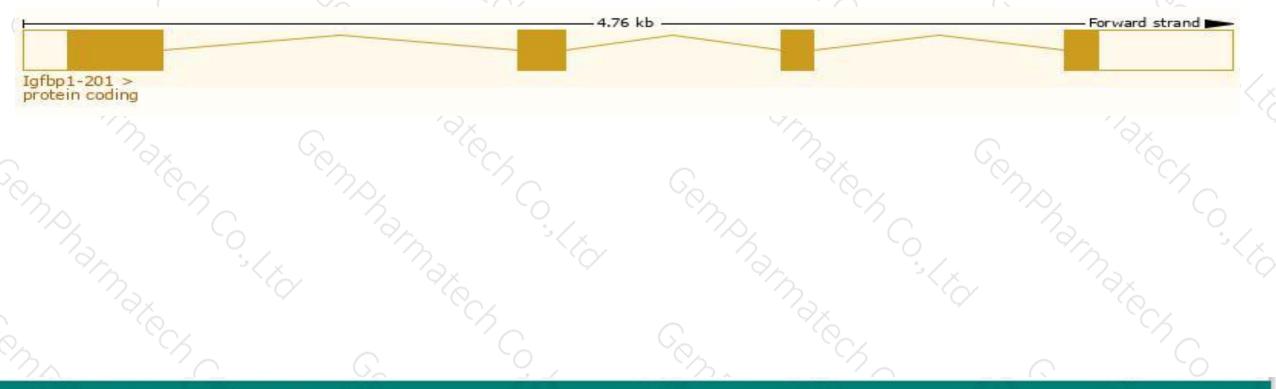
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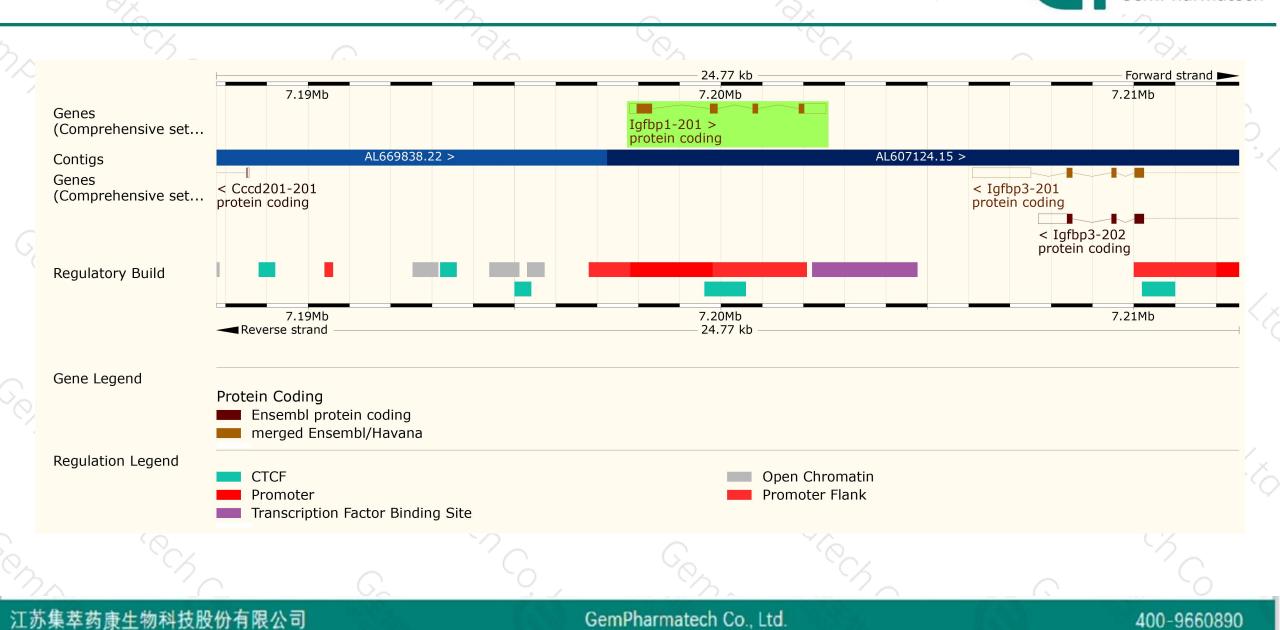
The gene has 1 transcript, and the transcript is shown below:

Name 🍦	Transcript ID 👙	bp 🖕	Protein 🛊	Translation ID	Biotype 💧	CCDS	UniProt 🖕		Flags	4
lgfbp1-201	ENSMUST0000020704.7	1525	<u>272aa</u>	ENSMUSP0000020704.7	Protein coding	<u>CCDS24427</u> &	<u>P47876</u>	TSL:1	GENCODE basic	APPRIS P1

The strategy is based on the design of *Igfbp1-201* transcript, The transcription is shown below



Genomic location distribution



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Protein domain



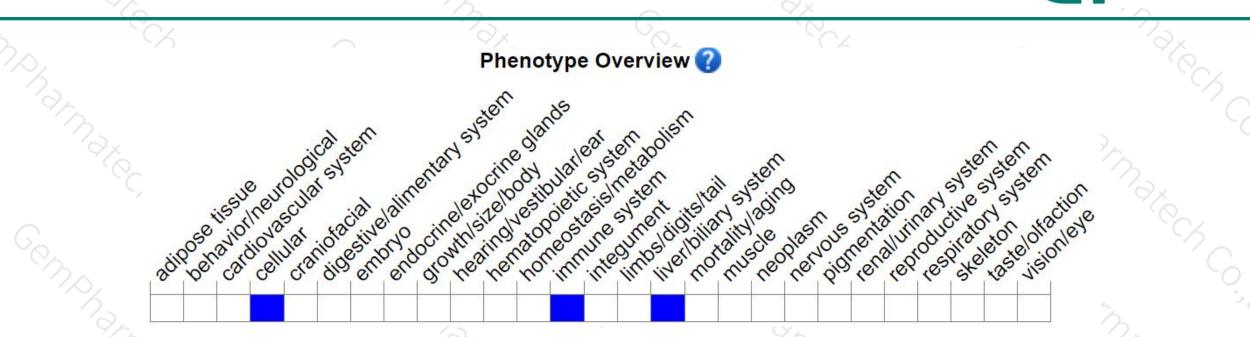
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All sequence SNPs/i Sequence variants (dbSNP and all other sources) Variant Legend missense variant Sequence variants	Gene3D	Insulin-like growth factor binding protein	4.10.40.20			Thyroglobulin type-1 superfamil	у		
Variant Legend synonymous variant synonymous variant	CDD					Thyroglobulin type-1			
missense variant synonymous variant	All sequence SNPs/i	Sequence variants (dbSNP and all other sourc	es)	1.11		с. н. с.	1.1	1.1	
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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 disruptions in this gene desplay a grossly normal phenotype but are more susceptible to liver injury.



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



