

Gdf1 Cas9-KO Strategy

Designer: Daohua Xu

Reviewer: Xueting Zhang

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Project Overview



Project Name Gdf1

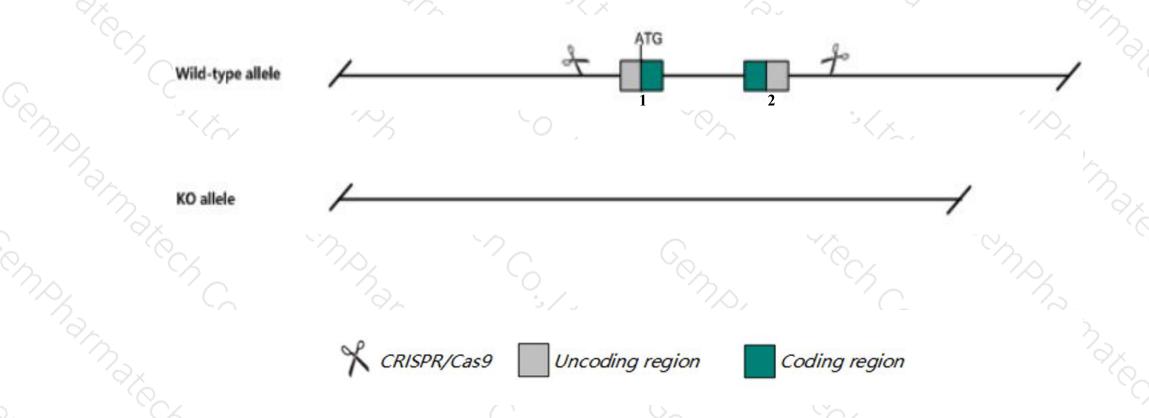
Project type Cas9-KO

Strain background C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Gdf1* gene. The schematic diagram is as follows:



Technical routes



- The *Gdf1* gene has 1 transcript. According to the structure of *Gdf1* gene, exon1-exon2 of *Gdf1-201*(ENSMUST00000207684.1) 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 *Gdf1* gene. The brief process is as follows: CRISPR/Cas9 system 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.

Notice



- > According to the existing MGI data, homozygous null mice display partially penetrant late embryonic lethality, neonatal lethality, situs inversus, right pulmonary isomerism, and other left-right patterning defects, cardiac septal defects, annular pancreas, and abnormal spleen morphology.
- The KO region contains functional region of the *Upf1,Cers1* gene.Knockout the region may affect the function of *Upf1,Cers1* gene.
- The *Gdf1* 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 the gene knockout on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.

Gene information (NCBI)



Gdf1 growth differentiation factor 1 [Mus musculus (house mouse)]

Gene ID: 14559, updated on 13-Mar-2020

Summary

☆ ?

Official Symbol Gdf1 provided by MGI

Official Full Name growth differentiation factor 1 provided by MGI

Primary source MGI:MGI:95683

See related Ensembl:ENSMUSG00000087408 Ensembl:ENSMUSG00000109523

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

Also known as Al385651, Gdf-1

Summary This gene encodes a secreted ligand of the TGF-beta (transforming growth factor-beta) superfamily of proteins. Ligands of this family bind

various TGF-beta receptors leading to recruitment and activation of SMAD family transcription factors that regulate gene expression. The encoded preproprotein is proteolytically processed to generate each subunit of the disulfide-linked homodimer. This protein is involved in the establishment of left-right asymmetry in early embryogenesis and in neural development in later embryogenesis. This protein is translated from a monocistronic mRNA early in development, and from a bicistronic mRNA in later stages that also encodes ceramide synthase 1. [provided by

RefSeq, Jul 2016]

Expression Biased expression in frontal lobe adult (RPKM 27.2), cortex adult (RPKM 24.4) and 11 other tissuesSee more

Orthologs human all

Transcript information (Ensembl)



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

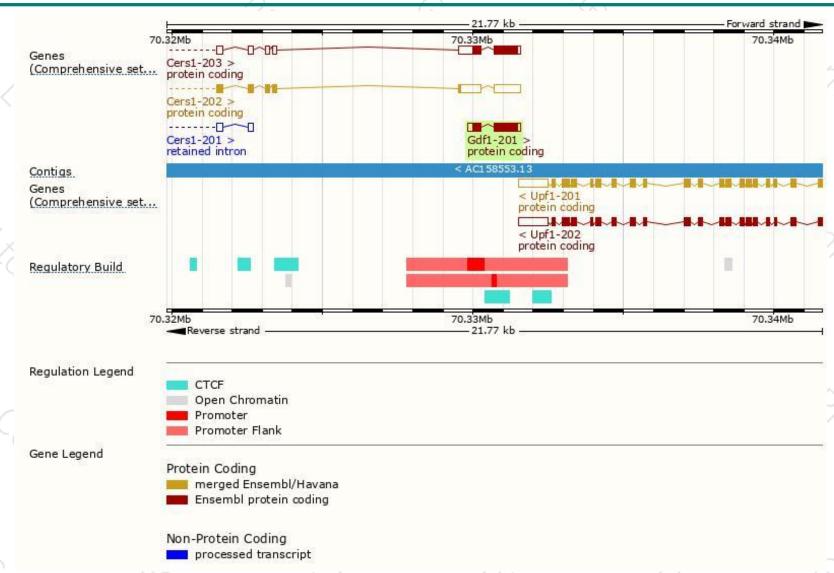
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Gdf1-201					0000000000		TSL:1 GENCODE basic APPRIS is a system to annotate alternatively spliced transcripts based on a range of computational methods to identify the most functionally important transcript(s) of a gene. APPRIS P1

The strategy is based on the design of *Gdf1-201* transcript, the transcription is shown below:



Genomic location distribution





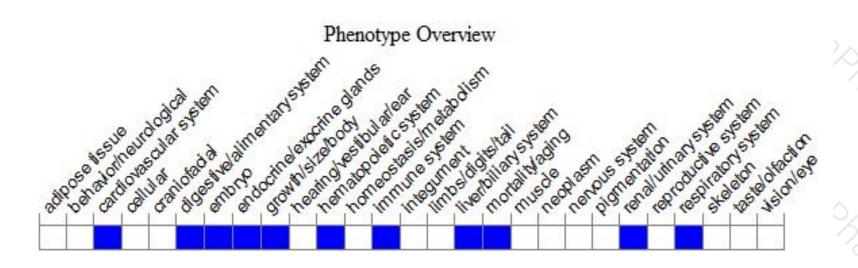
Protein domain





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, homozygous null mice display partially penetrant late embryonic lethality, neonatal lethality, situs inversus, right pulmonary isomerism, and other left-right patterning defects, cardiac septal defects, annular pancreas, and abnormal spleen morphology.



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





