

Wdr41 Cas9-KO Strategy

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Project Overview



Project Name

Wdr41

Project type

Cas9-KO

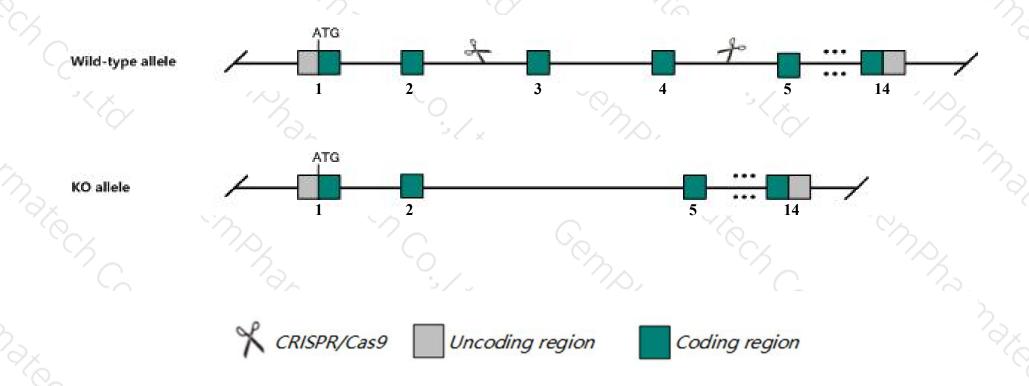
Strain background

C57BL/6JGpt

Knockout strategy



This model will use CRISPR/Cas9 technology to edit the *Wdr41* gene. The schematic diagram is as follows:



Technical routes



- ➤ The *Wdr41* gene has 9 transcripts. According to the structure of *Wdr41* gene, exon3-exon4 of *Wdr41-205*(ENSMUST00000160801.7) transcript is recommended as the knockout region. The region contains 179bp coding sequence. Knock out the region will result in disruption of protein function.
- ➤ In this project we use CRISPR/Cas9 technology to modify *Wdr41* gene. The brief process is as follows: CRISPR/Cas9 system

Notice



- > According to the existing MGI data, Mice homozygous for an ENU-generated allele exhibit increased susceptibility to DDS-induced colitis with increased activated single positive T cells.
- The *Wdr41* gene is located on the Chr13. 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)



Wdr41 WD repeat domain 41 [Mus musculus (house mouse)]

Gene ID: 218460, updated on 31-Jan-2019

Summary

Official Symbol Wdr41 provided by MGI

Official Full Name WD repeat domain 41 provided by MGI

Primary source MGI:MGI:2445123

Ensembl:ENSMUSG00000042015 See related

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 B830029I03Rik, MSTP048

Summary This gene encodes a protein of unknown function, but which contains a WD40 domain consisting of six WD40 repeats. The WD40 domain is

one of the most abundant protein domains in eukaryotes, and is found in proteins with widely varying cellular functions. However, proteins with

this domain often provide a rigid scaffold for protein-protein interactions. [provided by RefSeq, Aug 2012]

Expression Ubiquitous expression in CNS E18 (RPKM 6.7), cerebellum adult (RPKM 6.6) and 28 other tissuesSee more

Orthologs human all

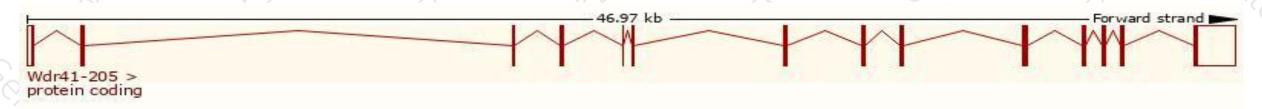
Transcript information (Ensembl)



The gene has 9 transcripts, all transcripts are shown below:

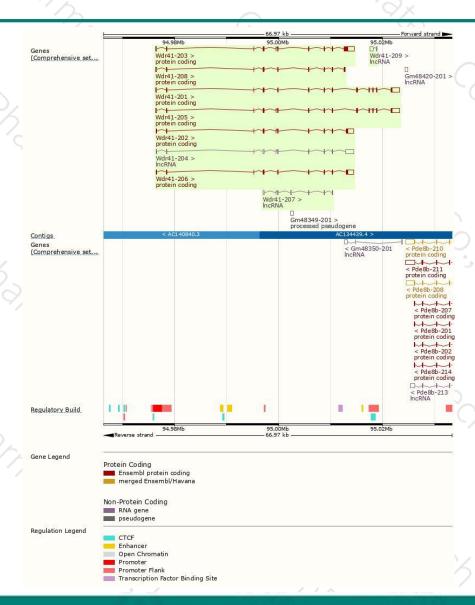
25' 170'	1997			2 2000			A Resident
Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Wdr41-205	ENSMUST00000160801.7	3062	<u>460aa</u>	Protein coding	CCDS56896	Q3UDP0	TSL:5 GENCODE basic APPRIS P1
Wdr41-201	ENSMUST00000056512.13	3063	<u>460aa</u>	Protein coding	-	Q3UDP0	TSL:1 GENCODE basic APPRIS P1
Wdr41-203	ENSMUST00000160115.7	2851	<u>467aa</u>	Protein coding	828	S4R290	TSL:5 GENCODE basic
Wdr41-202	ENSMUST00000159647.7	2289	329aa	Protein coding	1028	B2RQV3	TSL:5 GENCODE basic
Wdr41-206	ENSMUST00000167155.1	2289	329aa	Protein coding	181	B2RQV3	TSL:1 GENCODE basic
Wdr41-208	ENSMUST00000222995.1	1112	240aa	Protein coding	-	A0A1Y7VMD8	TSL:5 GENCODE basic
Wdr41-204	ENSMUST00000160409.1	2339	No protein	IncRNA	828	v	TSL:5
Wdr41-209	ENSMUST00000223422.1	756	No protein	IncRNA	727	-	TSL:5
Wdr41-207	ENSMUST00000222948.1	644	No protein	IncRNA	-		TSL:5

The strategy is based on the design of Wdr41-205 transcript, The transcription is shown below



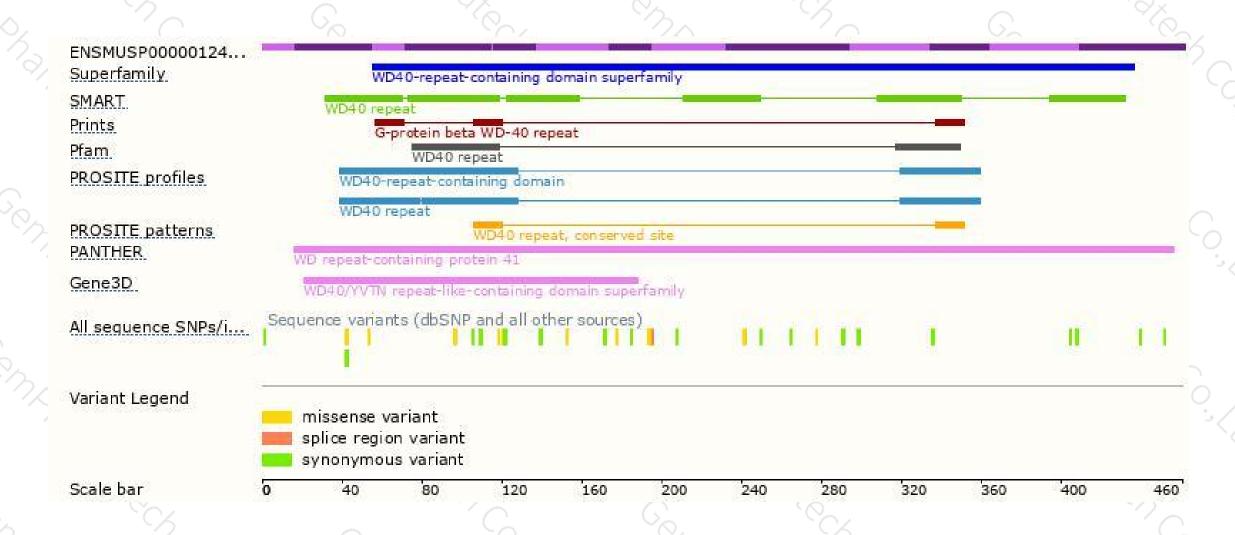
Genomic location distribution





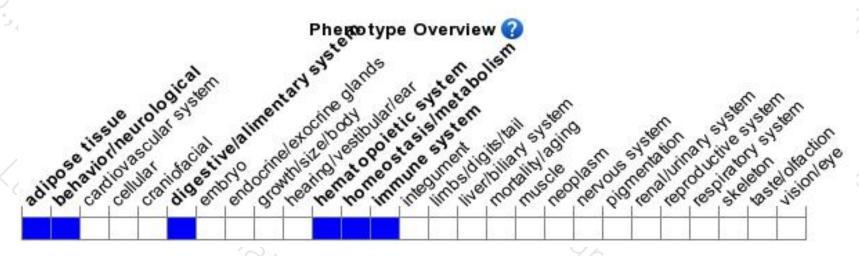
Protein domain





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 homozygous for an ENU-generated allele exhibit increased susceptibility to DDS-induced colitis with increased activated single positive T cells.



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





