



Zbtb16 Cas9-CKO Strategy

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Design Date: 2019-9-19
Reviewer: JiaYu

Project Overview

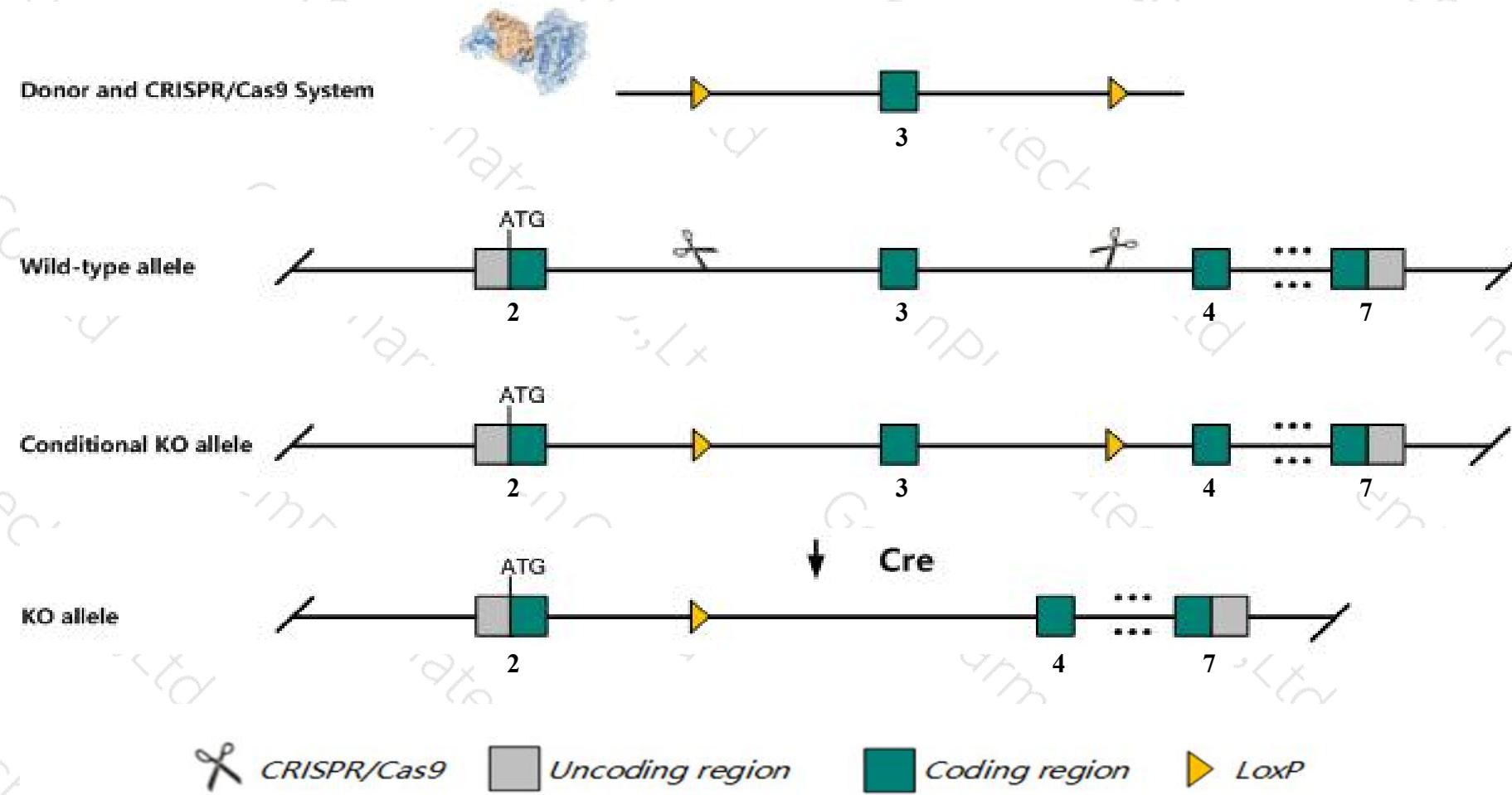
Project Name**Zbtb16**

Project type**Cas9-CKO**

Strain background**C57BL/6JGpt**

Conditional Knockout strategy

This model will use CRISPR/Cas9 technology to edit the *Zbtb16* gene. The schematic diagram is as follows:



Technical routes

- The *Zbtb16* gene has 2 transcripts. According to the structure of *Zbtb16* gene, exon3 of *Zbtb16-201* (ENSMUST00000093852.4) transcript is recommended as the knockout region. The region contains 98bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Zbtb16* 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.



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Notice

- According to the existing MGI data, Homozygous mutants exhibit abnormal anterior-posterior patterning, with skeletal abnormalities of the limb, especially the hindlimb, and homeotic transformations of anterior skeletal elements into posterior structures. Males develop infertility due to loss of germline cells with age.
- The *Zbtb16* gene is located on the Chr9. 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)

Zbtb16 zinc finger and BTB domain containing 16 [Mus musculus (house mouse)]

Gene ID: 235320, updated on 19-Mar-2019

Summary



Official Symbol Zbtb16 provided by [MGI](#)

Official Full Name zinc finger and BTB domain containing 16 provided by [MGI](#)

Primary source [MGI:MGI:103222](#)

See related [Ensembl:ENSMUSG00000066687](#)

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 AI467657, PLZF, Zfp145, lu

Expression Broad expression in lung adult (RPKM 8.1), subcutaneous fat pad adult (RPKM 7.3) and 24 other tissues [See more](#)

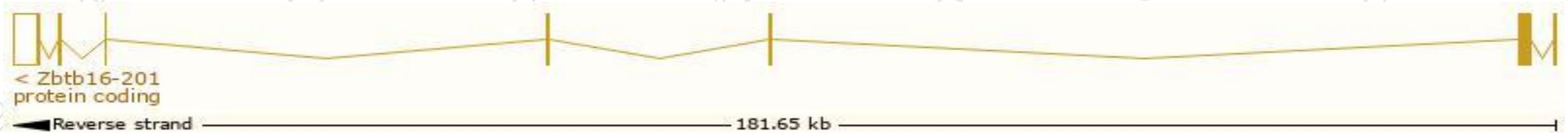
Orthologs [human](#) [all](#)

Transcript information (Ensembl)

The gene has 2 transcripts, all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Zbtb16-201	ENSMUST00000093852.4	5114	673aa	Protein coding	CCDS23158	Q3UQ17	TSL:1 GENCODE basic APPRIS P1
Zbtb16-202	ENSMUST00000216150.1	2505	673aa	Protein coding	CCDS23158	Q3UQ17	TSL:1 GENCODE basic APPRIS P1

The strategy is based on the design of *Zbtb16-201* transcript, The transcription is shown below



Genomic location distribution

Contigs

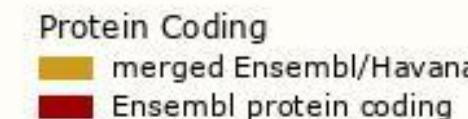
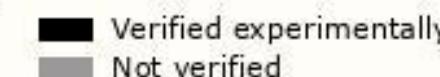
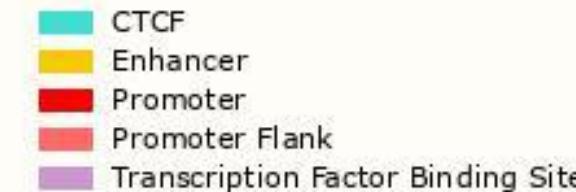
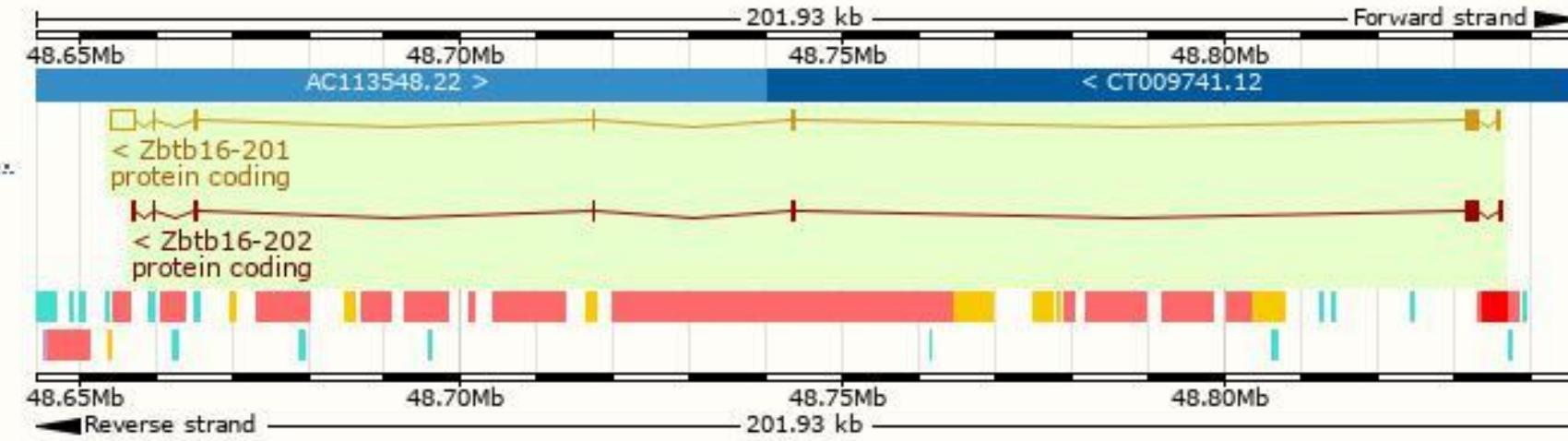
Genes
(Comprehensive set...)

Regulatory Build

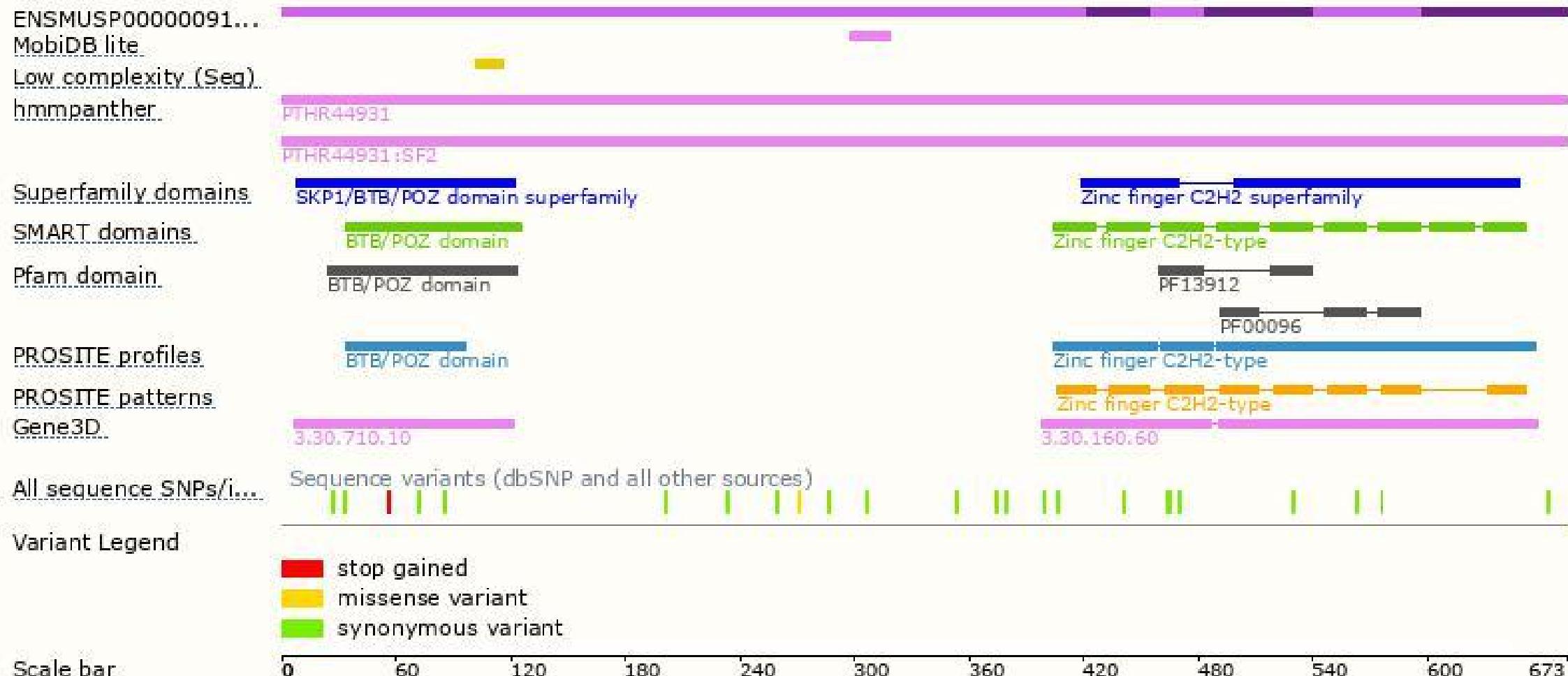
Regulation Legend

Motif Feature Legend

Gene Legend



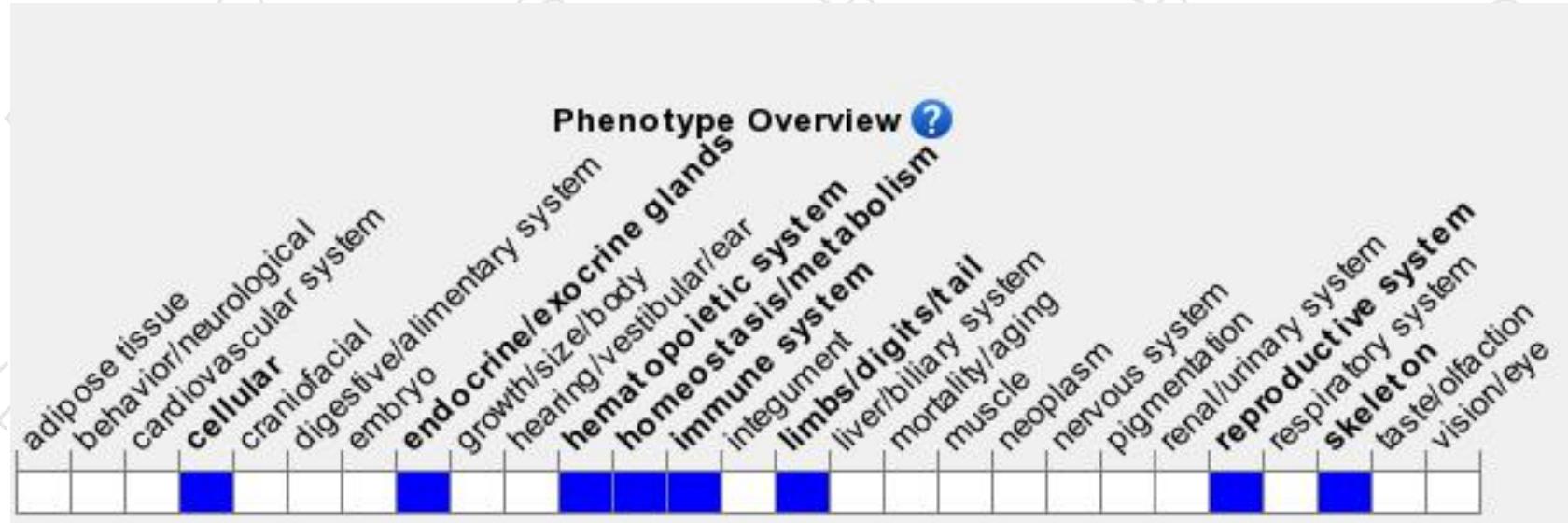
Protein domain





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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, Homozygous mutants exhibit abnormal anterior-posterior patterning, with skeletal abnormalities of the limb, especially the hindlimb, and homeotic transformations of anterior skeletal elements into posterior structures. Males develop infertility due to loss of germline cells with age.



If you have any questions, you are welcome to inquire.

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