



Atmin Cas9-KO Strategy

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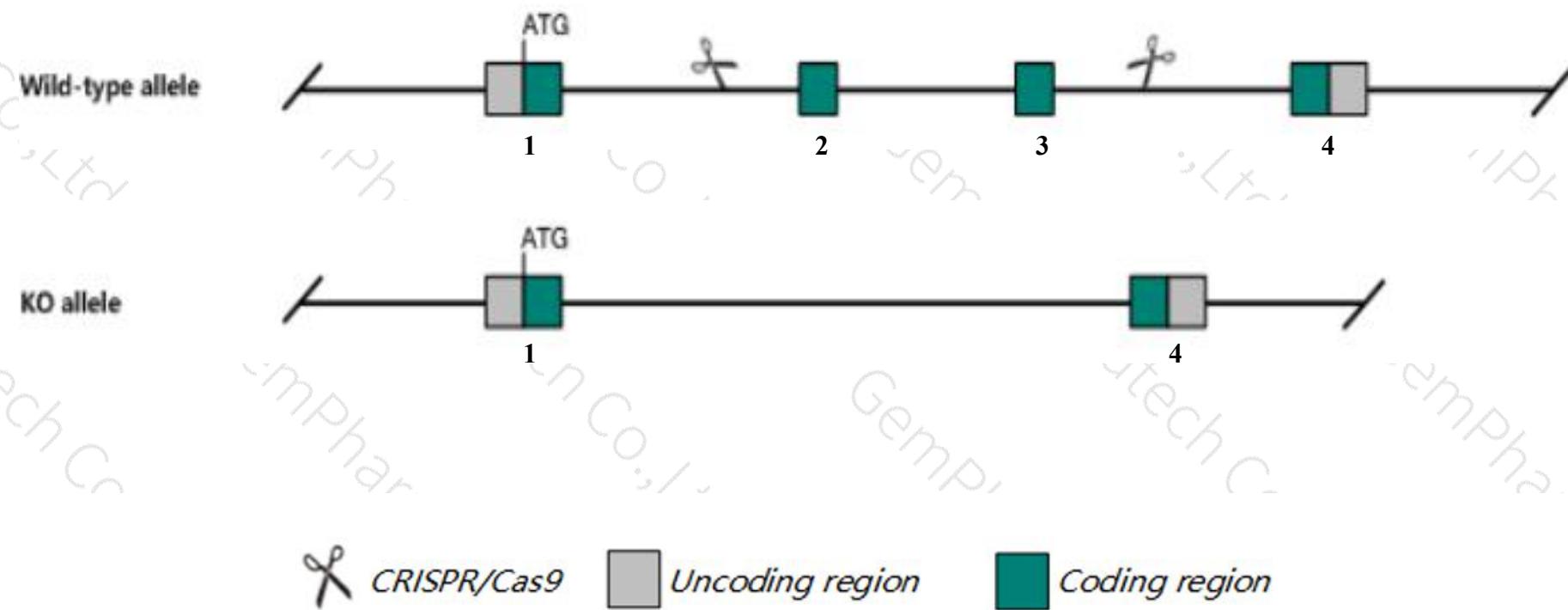
Design Date: 2020-7-22

Project Overview

Project Name	<i>Atmin</i>
Project type	Cas9-KO
Strain background	C57BL/6JGpt

Knockout strategy

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



Technical routes

- The *Atmin* gene has 1 transcript. According to the structure of *Atmin* gene, exon2-exon3 of *Atmin-201*(ENSMUST00000109099.3) transcript is recommended as the knockout region. The region contains 326bp coding sequence. Knock out the region will result in disruption of protein function.
- In this project we use CRISPR/Cas9 technology to modify *Atmin* 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.



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Notice

- According to the existing MGI data, mice homozygous for a knock-out allele exhibit fetal lethality, craniofacial defects, midbrain exencephaly, and premature senescence of mouse embryonic fibroblasts. Homozygotes for an ENU-induced mutation exhibit left-right patterning defects.
- The *Atmin* 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.



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Gene information (NCBI)

Atmin ATM interactor [Mus musculus (house mouse)]

Gene ID: 234776, updated on 13-Mar-2020

Summary ✖ ?

Official Symbol Atmin provided by [MGI](#)

Official Full Name ATM interactor provided by [MGI](#)

Primary source [MGI:MGI:2682328](#)

See related [Ensembl:ENSMUSG00000047388](#)

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 Asciz, gpg6, mKIAA0431

Expression Ubiquitous expression in testis adult (RPKM 16.7), cerebellum adult (RPKM 10.6) and 28 other tissues [See more](#)

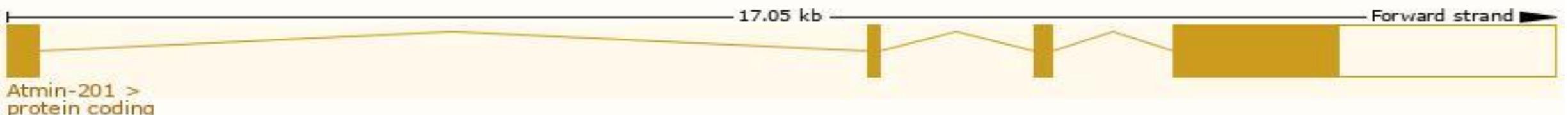
Orthologs [human](#) [all](#)

Transcript information (Ensembl)

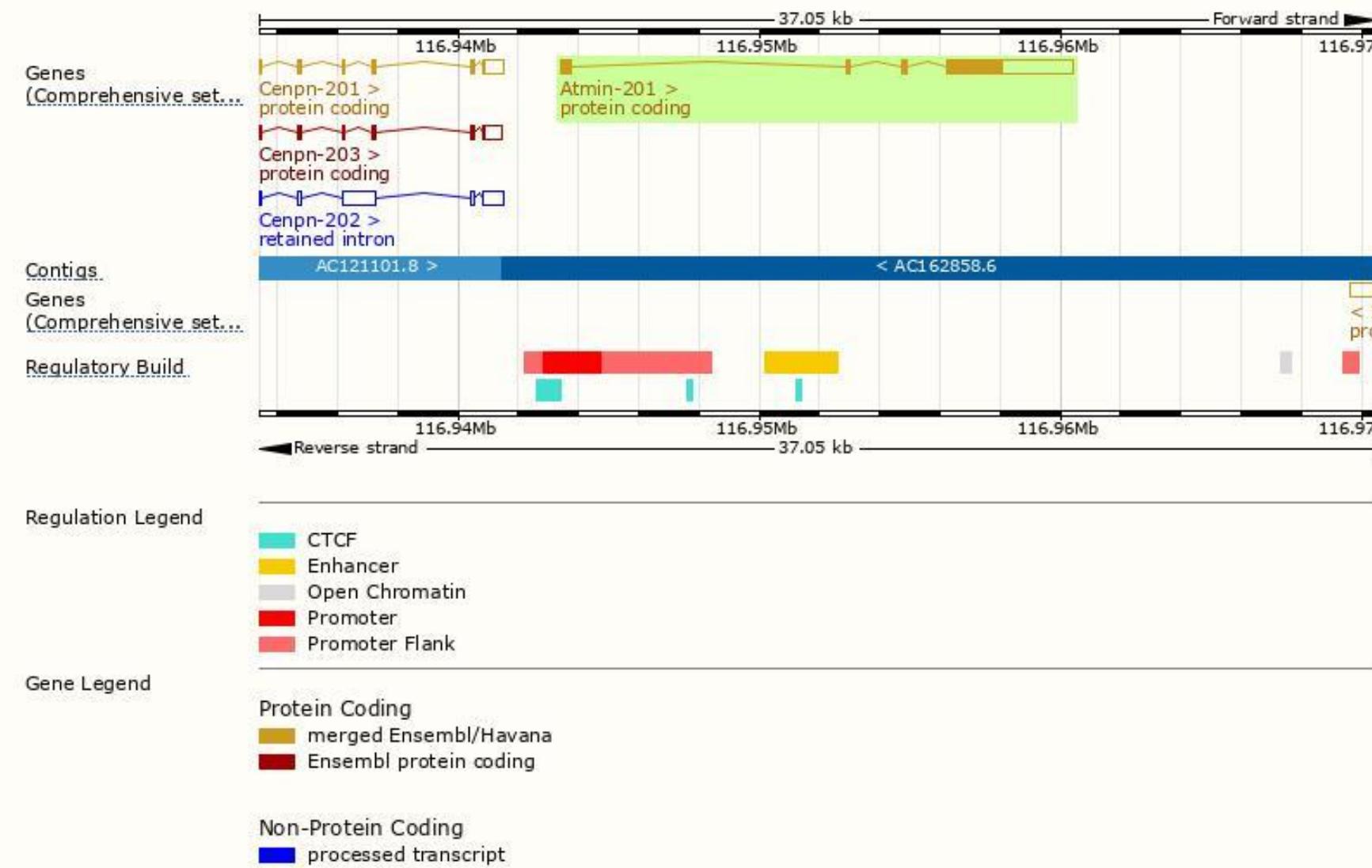
The gene has 1 transcript, and the transcript is shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Atmin-201	ENSMUST00000109099.3	4877	818aa	Protein coding	CCDS22694	Q6P9S1	TSL:1 GENCODE basic APPRIS P1

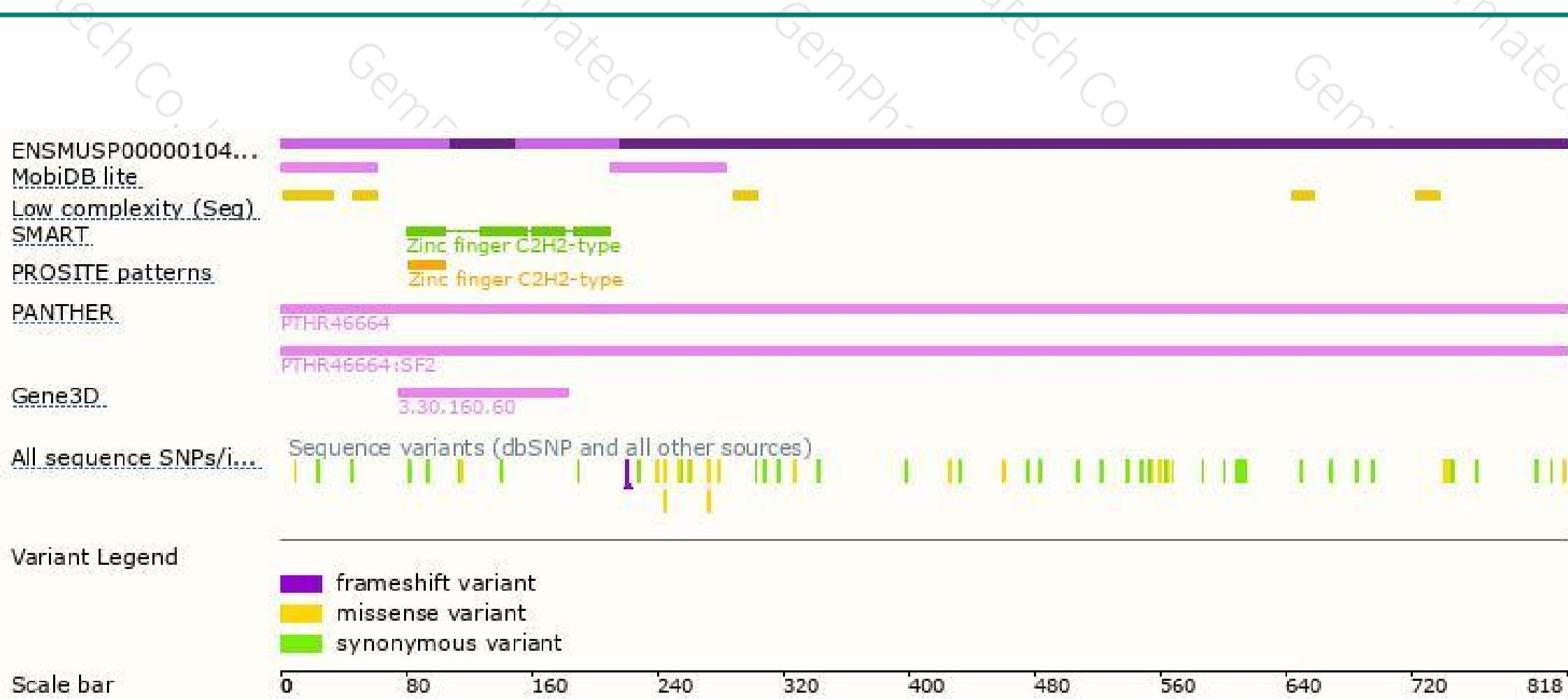
The strategy is based on the design of *Atmin-201* transcript, the transcription is shown below:



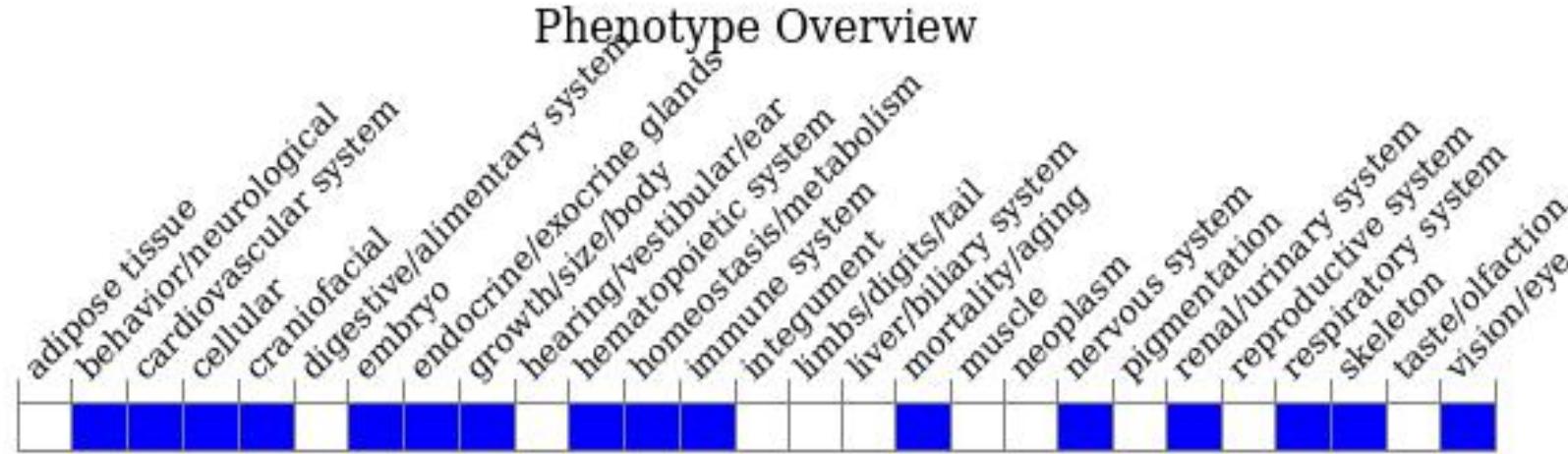
Genomic location distribution



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, mice homozygous for a knock-out allele exhibit fetal lethality, craniofacial defects, midbrain exencephaly, and premature senescence of mouse embryonic fibroblasts. Homozygotes for an ENU-induced mutation exhibit left-right patterning defects.



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

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