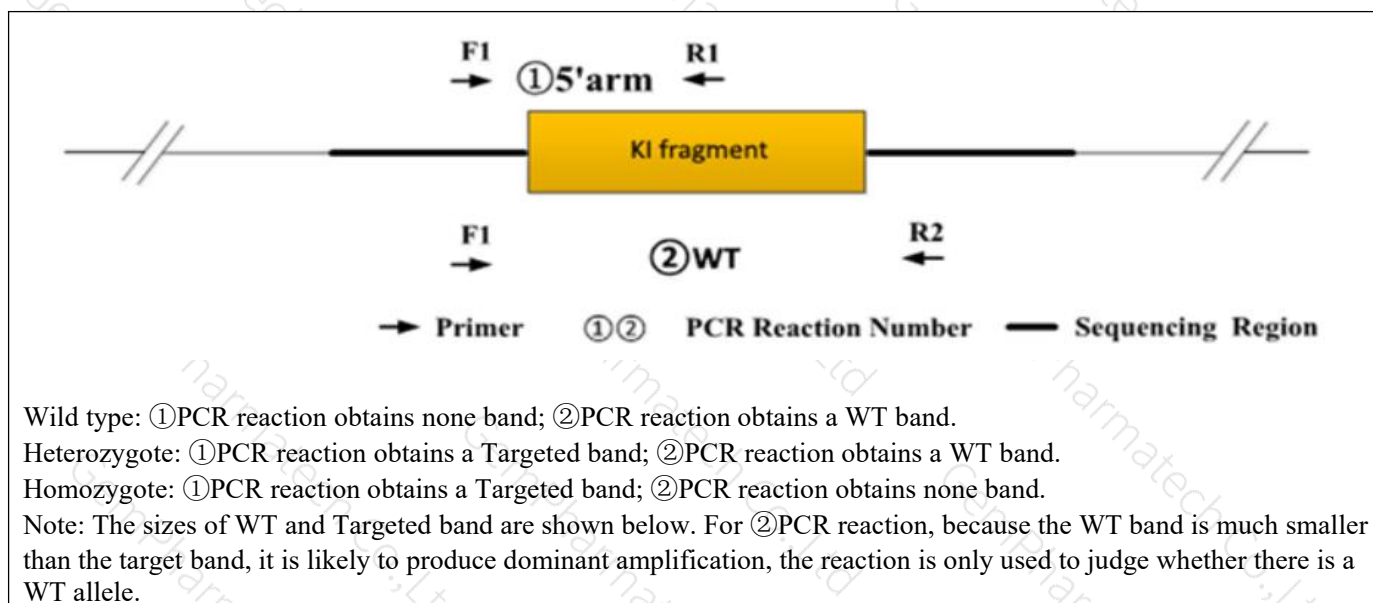


## Genotyping Report

Strain ID	T059381	Strain Type	KI(Cas9)	Genetic Background	C57BL/6JGpt
Designer	Tiantian Sun	Gene Name	<i>Cd81-NLS-tdtomato-P2A-DreERT2</i>		

### 1. Strategy of Genotyping



### 2. Primer Information

PCR No.	Primer No.	Primer Name	Sequence	Band Size
① 5'arm GC:62.6%	F1	PO-GJS0220221010507-01-Cd81-wt-tF1	CGGGACCTAGAGGTCTATAAAGAGTG	WT:0bp Targeted:326bp
	R1	tdTomato-tR1	GCGCATGAACTCTTTGATGACC	
② WT GC:62.6%	F1	PO-GJS0220221010507-01-Cd81-wt-tF1	CGGGACCTAGAGGTCTATAAAGAGTG	WT:334bp Targeted:3922bp
	R2	PO-GJS0220221010507-01-Cd81-wt-tR1	CTGCTCACCCAGAAGACGAAATTG	

### 3. Gel Image & Conclusion

--



Note: P:Heterozygous samples; WT: Wildtype control; B: Blank control (ddH<sub>2</sub>O); M: DNA Ladder

① Control (WT) : It is an important reference mark for whether the PCR reaction is successful and whether the product band position and size meet the theoretical requirements.

② Control (B) : PCR amplification was performed without template in the PCR reagent to monitor whether the reagent was contaminated.

#### 4. PCR Condition

PCR Reaction Component			
Seg.	reaction component	Volume (μl)	
1	2 × Rapid Taq Master Mix (Vazyme P222)	12.5	
2	ddH <sub>2</sub> O	9.5	
3	Primer A(10pmol/μl)	1	
4	Primer B(10pmol/μl)	1	
5	Template(20~80ng/μl)	1	
PCR program I priority selection			
Seg.	Temp.	Time	Cycle
1	95℃	5min	
2	98℃	30s	20×
3	65℃* (-0.5℃/cycle)	30s	
4	72℃	45s*	
5	98℃	30s	15×
6	55℃*	30s	
7	72℃	45s*	
8	72℃	5min	
9	10℃	hold	
PCR program II the second choice			
Seg.	Temp.	Time	Cycle
1	95℃	5min	
2	98℃	30s	35×
3	58℃*	30s	
4	72℃	45s*	

5	72℃	5min	
6	10℃	hold	

Note\*: Annealing temperature and extension time can be determined according to the actual amplification situation and amplification enzyme efficiency.