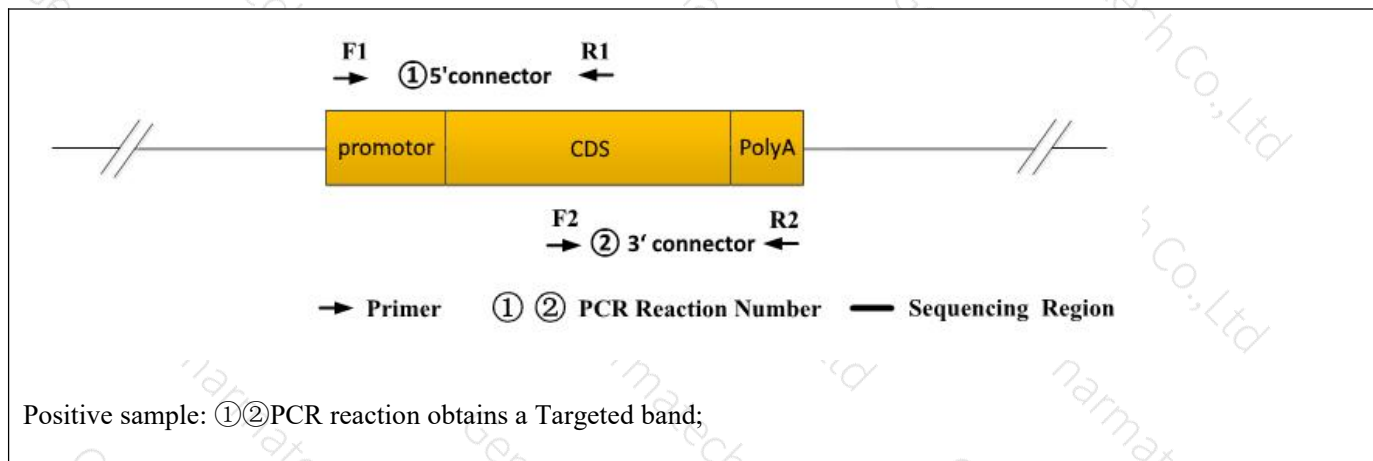


Genotyping Report

Strain ID	T052681	Strain Type	TG	Genetic Background	C57BL/6JGpt
Designer	Tianjiao Wang	Gene Name	<i>Sp7-iCre</i>		

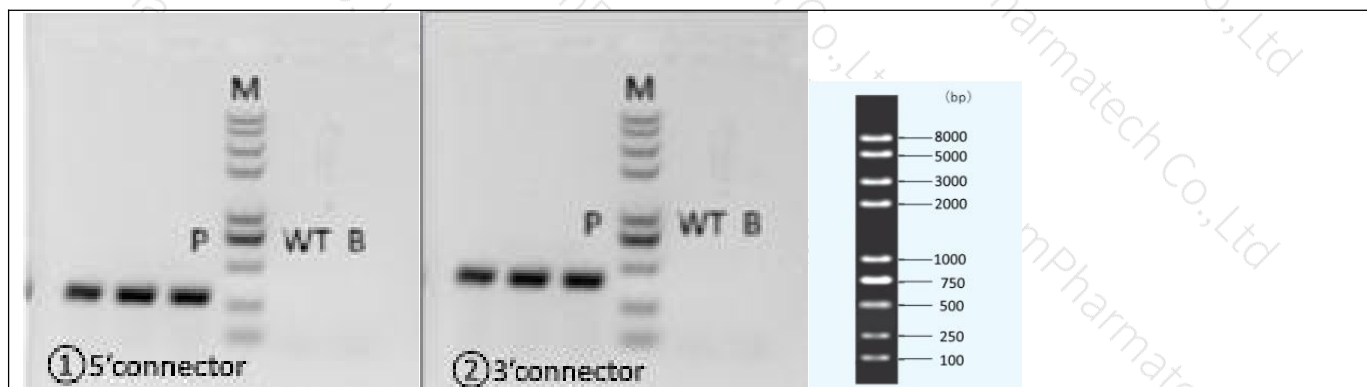
1. Strategy of Genotyping



2. Primer Information

PCR No.	Primer No.	Sequence	Band Size
①5'connector	T052681-F1	CCACCTCTGATGAAGTCAGGAAG	WT:0bp Targeted:376bp
	T052681-R1	TCAAAGTCAGTGC GTTCAAAGG	
②3'connector	T052681-F2	TTACCAAGCTGGTGGAGAGATG	WT:0bp Targeted:397bp
	T052681-R2	TGCACCTGAGGAGTTCAGTCC	

3. Gel Image & Conclusion



Note: P:Positive samples; WT: Wildtype control; B: Blank control (ddH₂O); 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	ddH2O		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	20×
2	98℃	30s	
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	35×
2	98℃	30s	
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.