

I Pharm		Genotyp	ing Report	np _{ana}	
Strain ID	T026095	Strain Type	CKO(Cas9)	Genetic Background	C57BL/6JGpt
Designer	Ya'nan Xu	Gene Name		Wwox	°C
Strategy of (F1	RI	F2 @2	R2	
_//	→ ①5'arm		-	'arm <u>↓</u>	
	Loxp 🔶 Prime	er (1) (2) PC	R Reaction Number	- Sequencing Region	

Wild type: ①PCR reaction obtains a single WT band; ②PCR reaction obtains a single WT band. Heterozygote: ①PCR reaction obtains a WT band and a Targeted band; ②PCR reaction obtains a WT band and a Targeted band. Homozygote: ①PCR reaction obtains a single Targeted band; ②PCR reaction obtains a single Targeted band.

Note: The sizes of WT and Targeted band are shown below.

2. Primer Information

PCR No.	Primer No.	Sequence	Band Size	
(1)(5'arm)	T026095-F1	ACTGTCTCGCTACACTCTTAACATCCT	WT:377 bp	
	T026095-R1 GAACTCAGAAATCTGCCTGCTTCTG		Targeted:482bp	
@(3'arm)	m) T026095-F2 CAGCAGTAGATTGTGACTTTGGATCC T026095-R2 GACGCTAATGTCAGATGTCAATGGAC		WT:282 bp	
			- Targeted:388bp	

3. Gel Image & Conclusion







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.
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	The star		
Seg.	read	Volume (µl)		
1 6	2 × Rapid Taq Master Mix(2 × Rapid Taq Master Mix (Vazyme P222)		
2	ddH2O	Co ,	9.5	
3	Primer A(10pmol/µl)	no. Ko		
4	Primer B(10pmol/µl)	12		
5	Template(≈100ng/µl)			
PCR program	① priority selection		Chr. A	
Seg.	Temp.	Time	Cycle	
1	95℃	5min		
2	98°C	30s	20×	
3	65℃*(-0.5℃/cycle)	30s	°C í C	
4 ⁽)	72°C	45s*		
5	98°C	30s	20×	
6	55℃*	30s	CLUS -	
7 6	72°C	45s*	Ga at a	
8 7	72°C	5min	No. C	
9	10 ℃	hold		



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PCR progran	n ${}^{\textcircled{0}}$ the second choice	20	·s/x	1	× 'C
Seg.	Temp.		Time		Cycle
1 ()	95°C	CC/	5min	C	nate in
2	98°C	6	30s		35×
3	58℃*	Ma,	30s	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~	
4	72°C	nar.	45s*		m the
5	72 °C	3	5min	6	
6 2	10°C	C.	hold	γ_{S}	$^{\sim}$ C

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