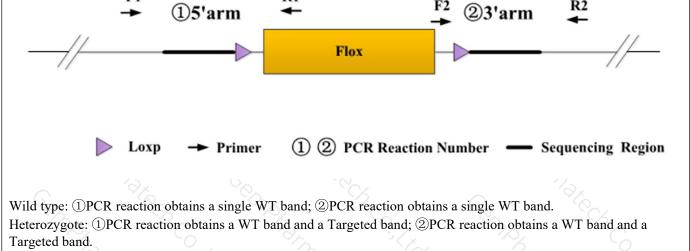


Strain ID	T019036	Strain Type	CKO(Cas9)	Genetic Background	C57BL/6JGp
Designer	Tiantian Sun	Gene Name	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~	Cav3	26
. Strategy of	Genotyping			a armax	



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
	T019036-F1	TGATGCAAGAAGAGTGGGCTAG	WT:295bp
(1)(5'arm)	T019036-R1	CATTGCAGGTTAGACATGGCAG	Targeted:400bp
	T019036-F2	GCATCGCATTGTCTGAGTAGGTG	WT:0bp
2)(3'arm)	T019036-R2	GCATCACCAGGTGAGAATACCTC	Targeted:347bp

## 3. Gel Image & Conclusion

6	<u>`</u>	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~	_%	0	G
M	м Р Ш WT В	(bp) 	- <ree< th=""><th></th><th></th></ree<>		
P WT B			× Co.		ate ch
(1)(5'arm)	(2)(3'arm)	250 100		- Adres	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~



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Note: P: Positive control; WT: Wildtype control; B: Blank control ( $ddH_2O$ ); 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.

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

## 4. PCR Condition

PCR Reaction Comp	ponent	X Con	10 <sup>2</sup>
Seg.	reaction compo	onent	Volume (µl)
1 90	2 × Rapid Taq Master Mix (Vazyme P2	12.5	
2 7	ddH2O	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~	9.5
3	Primer A(10pmol/µl)		1 6
4 6.	Primer B(10pmol/µl)	× ~ ~	1 6
5	Template(≈100ng/μl)		
PCR program $\textcircled{1}$ pr	iority selection		
Seg.	Temp.	Time	Cycle
1	95°C	5min	197 <sub>8</sub>
2	98°C	30s	20×
3 <sup>9</sup> 12	65℃* (-0.5℃/cycle)	30s	
4	72℃	45s*	nay '
5 %	98°C	30s	20×
6	55°C*	30s /	
7 75	72℃	45s*	
80.	72℃	5min	122
9 7	10°C	hold	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~
PCR program ② tl	ne second choice	- 10 - 17	6
Seg.	Temp.	Time	Cycle
1	95℃	5min	
2	98°C	30s	35×
3	58℃*	30s	
4 %	72°C	45s*	
5	72℃	5min	(9/2)
6 6	10°C	hold	925

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



