

GENOTYPING PROTOCOL

MUTANT MOUSE RESOURCE & RESEARCH CENTER: UC DAVIS

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530-754-MMRRC

Protocol Name: CR1626 Trappc1 GenDel

Protocol: GoTaq® G2 Colorless Master Mix(Promega)

Reagent/Constituent	Volume (μL)
Water	3.9
GoTaq® G2 Colorless Master Mix,2X	7.5
Primer 1. (stock concentration is 20μM) comF	0.6
Primer 2. (stock concentration is 20μM) wtR	1.2
Primer 3. (stock concentration is 20μM) mutR	0.3
DNA (example) extracted w/ "Qiagen DNeasy columns or other similar silica based kits"	1.5
TOTAL VOLUME OF REACTION:	15.00 μL

Comments on protocol:

- Protocol may work with other DNA extraction methods.

Strategy:

Steps	Temp (°C)	Time (m:ss)	# of Cycles
1. Initiation/Melting HOT START? ☐	94	2:00	1x
2. Denaturation	94	0:10	
3. Annealing steps 2-3-4 cycle in sequence	65 (↓1°C/cycle)	0:30	10x
4. Elongation	68	2:00	
5. Denaturation	94	0:15	
6. Annealing steps 5-6-7 cycle in sequence	55	0:30	25x
7. Elongation	68	2:00 (↑20sec/cycle)	
8. Finish	4	∞	n/a

Primers:

Name	Nucleotide Sequence (5' - 3')	Agarose: 1.5%	V: 90
1. CR_Trappc1-comF	GCACATCACTGGGCTTCAGAGG	Estimated Running Time: 90 min.	
2. CR_Trappc1-wtR	GAGGACTGGACTGGGACTATGTGC	Primer Combination	Band (bp)
3. CR_Trappc1-mutR	AACTCACACTCAGGCAGCAGAGG	1 & 2, 1 & 3	445,1946*
		1 & 3	307
			wildtype
			mutant

Electrophoresis Protocol:

Allele Description: Exon 1-5 ([ENSMUSE00000662059](#), [ENSMUSE00000384026](#), [ENSMUSE00000374380](#), [ENSMUSE00000366224](#), [ENSMUSE00000662058](#)) were constitutively deleted between the 5'UTR through the 3' UTR from the Trappc1 gene [ENSMUST00000102602.7](#) using CRISPR Cas9 gene editing technology in mouse zygotes. Subsequent founders were backcrossed to C57BL6/N to produce sequence confirmed heterozygous animals.

*May not see larger wildtype band

