

GENOTYPING PROTOCOL

MUTANT MOUSE RESOURCE & RESEARCH CENTER: UC DAVIS

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530-754-MMRRC

Protocol Name: CR1430 1700034E13Rik EXDEL

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:

Electrophoresis Protocol:

Name	Nucleotide Sequence (5' - 3')	Agarose: 1.5%	V: 90
1. CR_1700034E13Rik-comF	AGAGTGTAGACAGACCTGTTTCCTGC	Estimated Running Time: 90 min.	
2. CR_1700034E13Rik-wtR	ACGCCTTATCATTGTTGGCTGG	Primer Combination	Band (bp)
3. CR_1700034E13Rik-mutR	GTAGCTGGTTGGGGGCATCTCC	1 & 2, 1 & 3	695,1337*
		1 & 3	397
			wildtype
			mutant

Allele Description: Exon 3 [ENSMUSE00000143102](#) and flanking splicing regions were constitutively deleted from the 1700034E13Rik gene [ENSMUST00000118724.7](#) 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

