

GENOTYPING PROTOCOL

MUTANT MOUSE RESOURCE & RESEARCH CENTER: UC DAVIS

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530-754-MMRRC

Protocol Name: CR10314 Rab11fip2 EXDEL

Protocol: GoTaq® G2 Colorless Master Mix(Promega)

Reagent/Constituent	Volume (μL)
Water	4.5
GoTaq® G2 Colorless Master Mix,2X	7.5
Primer 1. (stock concentration is 20μM)	0.5
Primer 2. (stock concentration is 20μM)	0.5
Primer 3. (stock concentration is 20μM)	0.5
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_Rab11fip2-ivqc5F	ATAGCTTTCTGTCAGGAATCACTTGAGC	Estimated Running Time: 90 min.
2. CR_Rab11fip2-ivqc3F	AGTCGCCAGATAGATTCTTTGGAG	Primer Combination Band (bp) Genotype
3. CR_Rab11fip2-ivqc3R	TGTGCATTTGGGTGTCTAAGGAAGCTC	2 & 3, 1 & 3 375,1479 wildtype
		1 & 3 716 mutant

Electrophoresis Protocol:

Allele Description: Exon 2 [ENSMUSE00001256751](#) and flanking splicing regions were constitutively deleted from the Rab11fip2 gene [ENSMUSG00000040022](#) using CRISPR Cas9 gene editing technology in mouse zygotes. Subsequent founders were backcrossed to C57BL6/N to produce sequence confirmed heterozygous animals.

