

**GENOTYPING BY PCR PROTOCOL  
MUTANT MOUSE REGIONAL RESOURCE CENTER**

sacoord@mrrrc.org  
800-910-2291 North America, +1-530-757-5710 International

Please provide the following information required for genetic analysis of your mutant mice.

*Note to MAC users: to ensure your graphic can be viewed on a PC please follow the steps below when inserting the graphic into this document. DO NOT drag and drop or copy/paste the graphic into this document.*

- Open the original graphic in the program that created it
- Choose File, Save As
- Select No Compression in the save options.
- Save as JPG or PNG or similar format that's compatible with both PC and Mac Word versions.
- Switch to Word, choose Insert, Picture, From File and choose the newly saved picture.

*These instructions are very generic. The menu options for your graphics program may be different.*

Donating Investigator/PI			Frank Furnari			Email			ffurnari@ucsd.edu			Institution			Ludwig Institute for Cancer Research			Address			9500 Gilman Dr CMMME 3020			City			La Jolla			Lab Contact			Rachel Reed			Email			ratakara@ucsd.edu			Telephone			858 534 7809			FAX			858 534 7830			Strain Name			K125R-BL6			MMRRC Stock Number			43570		
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MMRRC: 0-CTR

NAME OF PCR: K125

Protocol: (PCR protocol provided by Donating Investigator)

Reagent/Constituent	Volume (µL)
Water	42.1
10x Buffer	5
MgCl <sub>2</sub> (stock concentration is mM)	
Betaine (stock concentration is 5M) <i>Optional</i>	
DNTPs (stock concentration is 10mM)	1
DMSO <i>Optional</i>	
Primer 1. (stock concentration is 20µM)	0.2
Primer 2. (stock concentration is 20µM)	0.2
Primer 3. (stock concentration is 20µM)	
Primer 4. (stock concentration is 20µM)	
Tag Polymerase 5units/µL	0.5
DNA (50-200ng/ µL) extracted w/ "Qiagen DNeasy columns or other similar silica based kits"	1
<i>The total volume is auto-calculated based on volumes entered, right click the total and update field to show/recalculate the total volume.</i>	
<b>TOTAL VOLUME OF REACTION:</b>	<b>50.000 µL</b>

*Comments on protocol:*

Strategy:

Steps	Temp (°C)	Time (m:ss)	# of Cycles
1. Initiation/Melting <input type="checkbox"/> HOT START?	94	3.0	1
2. Denaturation	94	0.30	
3. Annealing steps 2-3-4 cycle in sequence	50	0.45	30x
4. Elongation	72	1.0	
5. Amplification	72	5.0	1
6. Finish	4	∞	n/a

Primers: Electrophoresis Protocol:

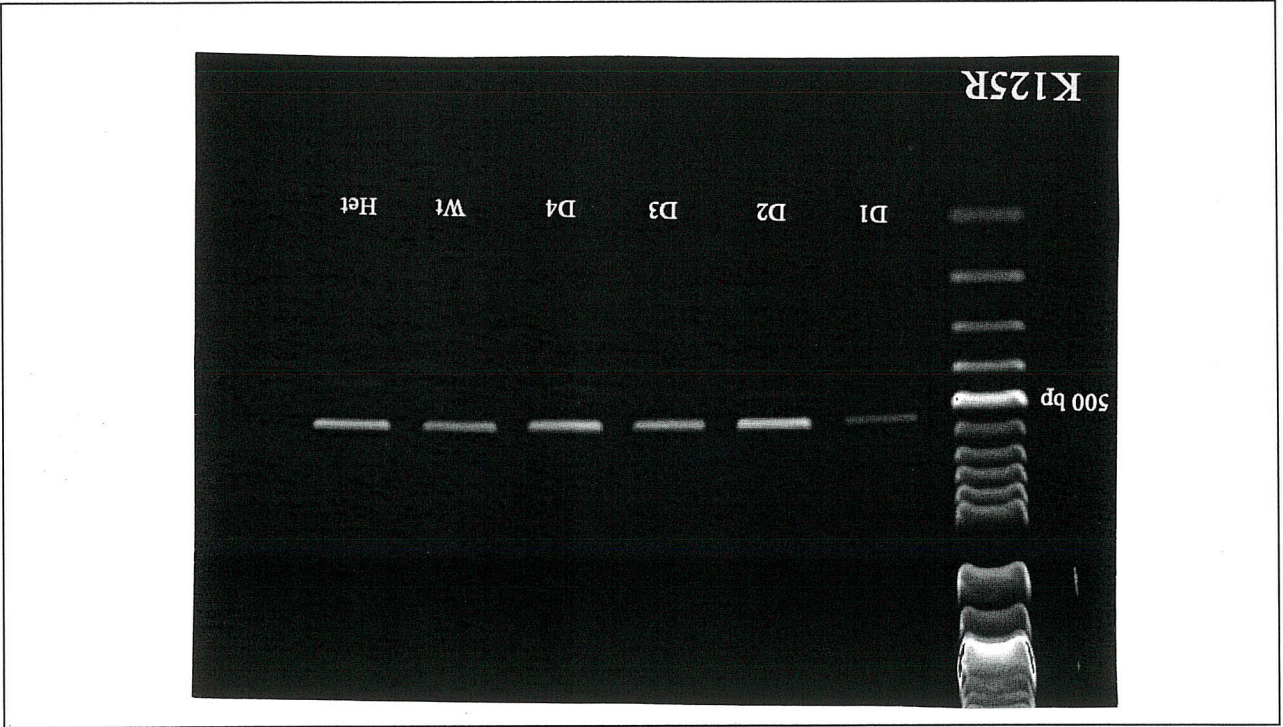
Name	Nucleotide Sequence (5' - 3')	Argarose: 1% V:110 min.	Estimated Running Time: min.
1. RTL1	CCTGAAAAGTTAG GCTTCTTTAAG		
2. RTL2	AACAGTTCTCAAAAGCATCAGACTG		
3.	RTL1 + RTL2		520 bp
4.			bp
5.			bp

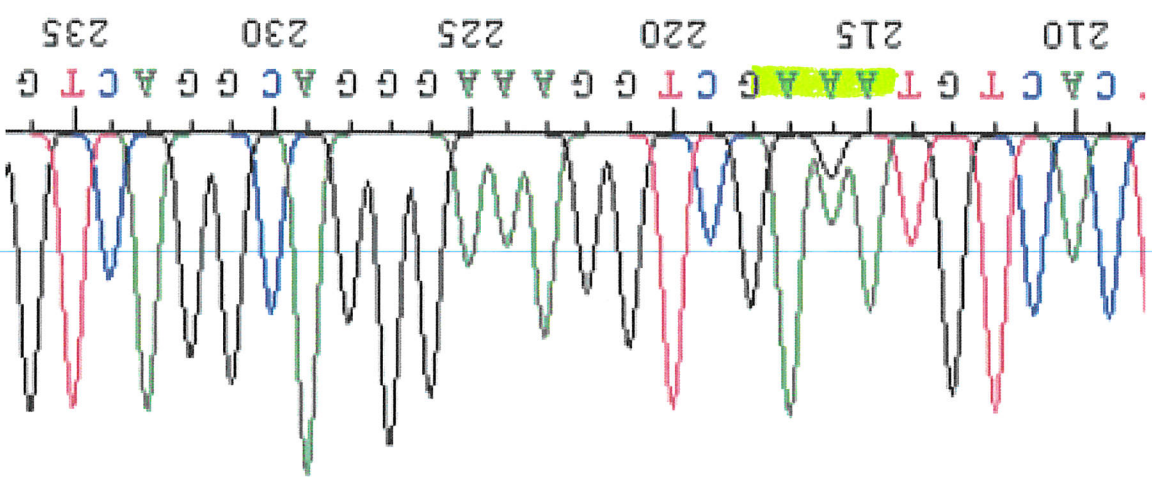
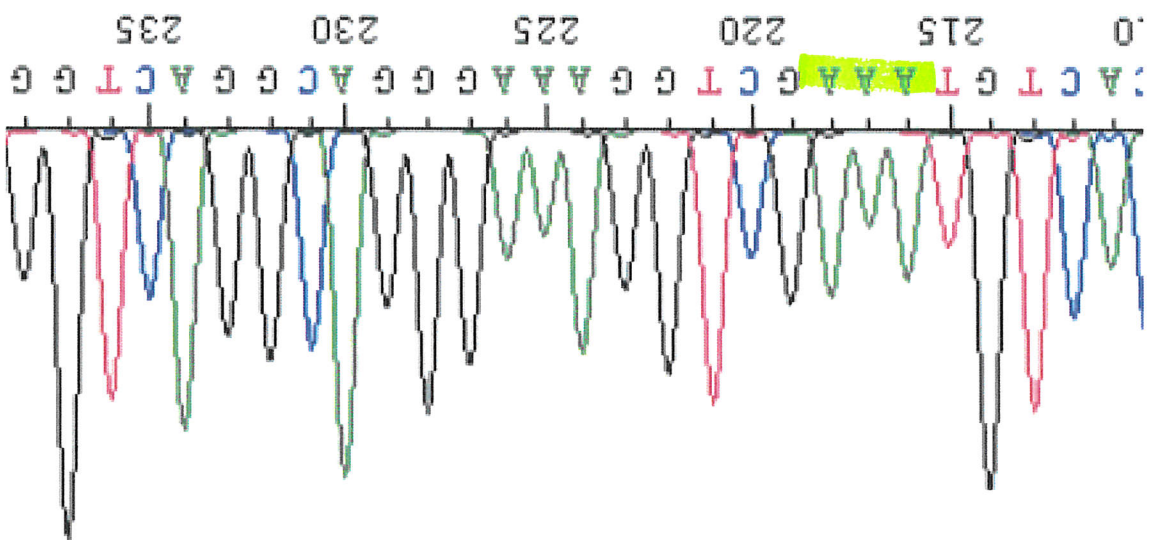
Please size gel images to fit in this space

**Protocol / Gel Comments:**

After running the PCR, purify, and send to sequencing with RLT1 and see if wildtype, homo, or het.

**Gel pictures:**





Trace: D1\_Rtl.ab1