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## Mouse Genotyping with KAPA Kit in 2 hours (#KK7302) V.1

DOI

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**Protocol status:** Working

**We use this protocol and it's working**

**Created:** January 29, 2020

**Last Modified:** January 29, 2020

**Protocol Integer ID:** 32461




## DNA Extraction

30m

1 Collect mouse tissues (ear or tail clips) into PCR strips.

2 Make  100  $\mu$ L extraction buffer mix for each sample.

10x Extraction buffer	10 $\mu$ L
H <sub>2</sub> O	88 $\mu$ L
Extraction Enzyme	2 $\mu$ L
Total	100 $\mu$ L

3 Add  100  $\mu$ L buffer into each tissue sample and run the reaction as follows:

 75 °C  00:15:00

 95 °C  00:05:00

 4 °C  00:00:00

### Note

Tissues are visible or intact after extraction (this is normal).  
Vortex DNA extract and spin down before PCR.

## PCR

30m

4 Make  20  $\mu$ L PCR master mix for each sample.

2x PCR buffer	10 $\mu$ L
H <sub>2</sub> O	8 $\mu$ L
10 $\mu$ M Primers mix	1 $\mu$ L
DNA	1 $\mu$ L
Total	20 $\mu$ L

5 Run the reaction as follows:

 95 °C  00:03:00



35 cycles



95 °C



00:00:15



57 °C adjustable



00:00:15



72 °C



00:00:10 per kb

35 cycles



4 °C



00:00:00

## Run DNA Gel

30m

6 Take  10 µL PCR reaction and run DNA gel.