Technical Bulletin

## M-Fast PCR Genotyping Kit (Cat. #: VGT-203/VGT-203p)

Cat: GT-203 Size: 1000 Extractions (Reagent-A: 250mL; Reagent-B: 50mL)

Storage: Room Temperature

Cat: GT-203p Size: 1000 Extractions

(Reagent-A: 250mL; Reagent-B: 50mL; 2x Green PCR mixture: 12.5mL)

\* 2x Green PCR mixture stored at -20°C

## \* Protocol for gDNA Extractions

1. Prepare 0.1-0.2cm mouse tail/ear biopsy sample in a 1.5/0.5mLmicrocentrifuge tube.

- 2. Add <u>250ul reagent-A</u> into the sample tube.
- 3. Place the tube in a PCR machine (or dry heat bath or heat block) and incubate at 95°C for 30 minutes. (Cover the sample tubes with a heavy book or others to prevent the cap opening during incubation)
- 4. Add <u>50ul reagent-B</u> into the sample tube and mix well by simple vortexing.
- 5. Pipette **2ul lysate supernatant** into an 23ul PCR mastermix (total: 25ul reaction)
- Run PCR reactions at thermal cyclers.

Note: These DNA samples are stable at room temperature for 1-3 weeks, or 1-3 months at 2-8°C and more than 3 years at -20°C.

## **Suggested PCR Protocol:**

I. Preparation of PCR Master Mix for a single reaction (total volume: 25uL) in a 0.2mL tube.

Component	Volume (μL)	Final Concentration
2x PCR Mastermix	12.5	1x
Forward primer (10μM)	1	250nM
Reverse primer (10μM)	1	250nM
DNA Template	2	Determined by user
PCR grade water	8.5 μL	Determined by user

## II. Setup typical thermal cycling parameters

After thermal cycling, the PCR products can be loaded directly onto an agarose gel and run gels as usual.

Precautions and Disclaimer: This product and procedure described are intended for R&D use only. Purchase of this product does not convey a license to perform any patented process.