

# **Cytomegalovirus PCR Detection Kit**

# For Research Use Only

Cat. No.: PK3001 Quantity: 50 Reactions Shipment: Wet Ice Storage: −20 °C

### Description

This kit designed for qualitative detection of human sample by the method of polymerase chain reaction.

#### **Kit Contents:**

The kit for 50 amplification reactions consists of:

1. 1x PCR MIX 1000 $\mu$ l 4. Mineral Oil 2ml 2. Taq DNA polymerase 15 $\mu$ l (5 $u/\mu$ l) 5. Positive Control 50 $\mu$ l 3. DNase Free, Deionized Sterile Water 5ml

The **Kit** should be stored at - 20°C.

#### **DNA Extraction:**

DNA can be extracted using DNP<sup>™</sup> (# EX6071) or using other standard methods.

### **PCR Protocol:**

- 1. Take out the kit and unfreeze the tubes, then put all the tubes on ice. Before opening tubes vortex and spin them. The final volume of each reaction will be 25µl.
- 2. Label new 0.5ml tubes for amplification reaction(s) for test(s), positive and negative control.
- 3. Add the following reagents for each tube on ice:

 1x PCR MIX
 19.7μl

 Taq DNA polymerase
 0.3μl

- Note: To avoid contamination all reagents must be taken with separate clean tips!
  - **4.** Mix the mixture thoroughly by shaking and spin.
  - 5. To each tube add one-drop (20-25 $\mu$ l) mineral oil.
  - 6. Add 5µl DNA\*(Use specified pipette for sampling of DNA).
  - $\textbf{7.} \ \textbf{Close tubes; spin the mixtures on microfuge for 3-5sec.}$
  - **8.** Transfer the tubes to preheated thermocycler and start the program:

## Cycling parameters:

95°C - 180sec 62°C - 40sec 72°C - 40sec **1 cycle** 93°C - 40sec 61°C - 40sec 72°C - 40sec

35 cycles

Analyze  $6\mu l$  of amplified samples directly in a 2% agarose gel without adding loading buffer. The presence of 222bp fragments indicates positive test.

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