

# Herpes Simplex virus I&II Detection Kit

For Research Use Only

Cat.No. : PK3051

**Quantity: 20 Reactions** 

Shipment: Wet/Dry Ice Storage: -20°C

## Description:

This kit is designed for qualitative detection of Herpes simplex Virus I&II (HSV I&II) DNA in the Human sample by the method of Polymerase Chain

Reaction.

## **Kit Contents:**

The kit for 20 amplification reactions consists of:

| 1. 1X PCR MIX                 | 400 µl        |
|-------------------------------|---------------|
| 2. Taq DNA polymerase         | 4 μl (5U/μl)  |
| 3. Positive Control           | 25μl (1pg/μl) |
| 4. Mineral Oil                | 1 ml          |
| 5. DNase Free Deionized Water | 5ml           |

## Sample preparation:

Performed in Pre-amplification, specimen, and control preparation area.

### DNA Extraction of Cerebrospinal fluid (CSF):

Reagents for DNA extraction are not included in this kit and should be ordered separately.

• DNP™ kit (Cat. No.: EX6071), SinaPure™ DNA Extraction (Cat No.:EX6011) is recommended for DNA extraction from CSF samples.

DNA can also be extracted by other standard method like Phenol/Chloroform.

Label tubes for patient, negative & positive controls.

## PCR Protocol:

Performed in Pre-amplification, Reagent preparation area.

- 1. Defreeze reagents in room temperature and then put all the tubes on ice. Vortex and spin tubes before opening. The final volume of each PCR reaction will be 25  $\mu$ l.
- 2. Label new 0.5 ml tubes for test(s), positive and negative control.
- 3. Add the following reagents for each tube on ice:

| 1x PCR MIX   | 20 µl        |  |
|--|--------------|--|
| Taq DNA polymerase   | 0.2 μl       |  |
| • 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).  |              |  |
| 7. Close tubes; spin the mixtures on microfuge for 3-5 sec.  |              |  |
| 8. Transfer the tubes to preheated thermocycler and start the program:   |              |  |
| Cycling parameters:  |              |  |
| 94°C 2mi   | n            |  |
| 94°C 30 s  | ec           |  |
| 62°C 15 s  | ec 33 cycles |  |
| 72°C 40 s  | ec           |  |
| 72°C 5 mi  | n            |  |
| Result analysis:   |              |  |
| Performed in Pre-amplification area  |              |  |
| Analyze 10 $\mu l$ of amplified samples directly in a 2% agarose gel without adding loading buffer. The presence of <b>256 bp</b> fragment indicate positive test. |              |  |
|  |              |  |

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