

Herpes Simplex virus I&II Detection Kit

Cat. No.: PR8240C Shipment:Wet/Dry Ice Quantity: 20 Reactions Storage: -20°C

Description:

This kit is designed for qualitative detection of Herpes simplex Virus I & Π (HSV I & Π) DNA in the Human sample by the method of Polymerase Chain Reaction. CinnaGen HSV PCR Detection Kit may be used in clinical medicine to detect HSV I & II DNA.

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.

■ Cinnagen DNPTM kit (Cat. No.: DN8115C),

Cinnapure DNA Extraction (Cat No.: PR881613) 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.

- 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 25µl.
- 2. Label new 0.5 ml tubes for test(s), positive and negative control.

3. Add the following reagents for each tube on ice:

 $\begin{array}{ccc} \text{1x PCR MIX} & 20 \ \mu\text{I} \\ \text{Taq DNA polymerase} & 0.2 \ \mu\text{I} \end{array}$

- 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 μ 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:

Result analysis:

Performed in Pre-amplification area Analyze 10 μ l of amplified samples directly in a 2% agarose gel without adding loading buffer. The presence of 256 bp fragment indicate positive test.