# **Package insert**



## GeneProof Herpes Simplex Virus (HSV-1/2) PCR Kit

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In vitro diagnostic medical device

The kit has been manufactured according to EC Directive 98/79/EC as an *in vitro* diagnostic medical device and it has been designed for professional use in specialized clinical and research laboratories.

### **KIT** CONTENT

	ISIN Version IS included in the MasterMix			ISEX Version IS supplied in a separate tube Nucleic acid isolation and PCR inhibition control		
REF	HSV/ISIN/025	HSV/ISIN/050	HSV/ISIN/100	HSV/ISEX/025	HSV/ISEX/050	HSV/ISEX/100
	25 rxn	50 rxn	100 rxn	25 rxn	50 rxn	100 rxn
MasterMix HSV	1x750 μl	2x750 μl	4x750 μl	1x750 μl	2x750 μl	4x750 μl
<b>CALIBRATOR</b> HSV 10 <sup>4</sup> copy/μl	1x200 μl	1x200 μl	1x200 μl	1x200 μl	1x200 μl	1x200 μl
<b>CALIBRATOR</b> HSV 10 <sup>3</sup> copy/μl	1x200 μl	1x200 μl	1x200 μl	1x200 μl	1x200 μl	1x200 μl
<b>CALIBRATOR</b> HSV 10 <sup>2</sup> copy/μl	1x200 μl	1x200 μl	1x200 μl	1x200 μl	1x200 μl	1x200 μl
<b>CALIBRATOR</b> HSV 10 <sup>1</sup> copy/μl	1x200 μl	1x200 μl	1x200 μl	1x200 μl	1x200 μl	1x200 μl
Internal Standard HSV	-	-	-	1x1000 μl	1x1000 μl	2x1000 μl

## STORAGE AND TRANSPORTATION CONDITIONS

The kits should be transported and stored at temperatures between -85 °C and -10 °C. The kit will remain stable at least until the expiry date printed on the package, if the storage temperature is kept. Repeated freezing and thawing of the kit components may result in lower detection quality.

## TECHNICAL SPECIFICATION

Target sequence	DNA conservative section of a single-copy gene encoding the B glycoprotein B (gB)
Specificity	Herpes simplex virus type 1 (HSV-1)
	Herpes simplex virus type 2 (HSV-2)
Sensitivity (LOD)	for HSV-1 reaches 0.764 copies/µl with the probability of 95 %
	for HSV-2 reaches 1.309 copies/µl with the probability of 95 %
Accuracy of measurement	for HSV-1 within the range of 10 <sup>4</sup> -0.764 copies/µl the detection accuracy is 0.5 log
	for HSV-2 within the range of 10 <sup>4</sup> -1.309 copies/µl the detection accuracy is 0.5 log
Linear range of measurement	for HSV-1 within the range of 10 <sup>7</sup> -0.764 copies/µl
	for HSV-2 within the range of 10 <sup>7</sup> -1.309 copies/µl
Sample types	whole blood in EDTA, plasma, cerebrospinal fluid, swabs, urine
Reporting units	copies/ml
Quality Control	regularly tested by QCMD and Instand e.V. External Quality Assessment Panels



### METHOD PRINCIPLES

The PCR kit is designed for detection and differentiation of both types of Herpes simplex viruses (HSV-1 and HSV-2) by the real-time Polymerase Chain Reaction (multiplex PCR) method. The HSV detection is based on the amplification of a specific conservative DNA sequence of a single-copy gene encoding glycoprotein B (gB) and measuring the amplification product concentration using the PCR process and fluorescence labelled probes. HSV-1 presence is indicated by FAM fluorophore fluorescence growth and the presence of HSV-2 is indicated by Cy5 fluorophore fluorescence growth. An Internal Standard (IS) is included in the reaction mix, controlling the possible inhibition of the PCR reaction (ISIN version) and possibly also the DNA extraction process quality (ISEX version). IS positive amplification is detected in the HEX fluorophore fluorescence channel. The detection kit takes advantage of the "hot start" technology, minimizing non-specific reactions and assuring maximum sensitivity. Ready to Use MasterMix contains uracil-DNAglycosylase (UDG), eliminating possible contamination of the PCR reaction by amplification products. The kit performs very sensitive HSV-1 and HSV-2 detection in clinical material (EDTA whole blood, plasma, cerebrospinal fluid, swab, urine). The kit is designed for *in vitro* diagnostics and provides qualitative and quantitative detection.

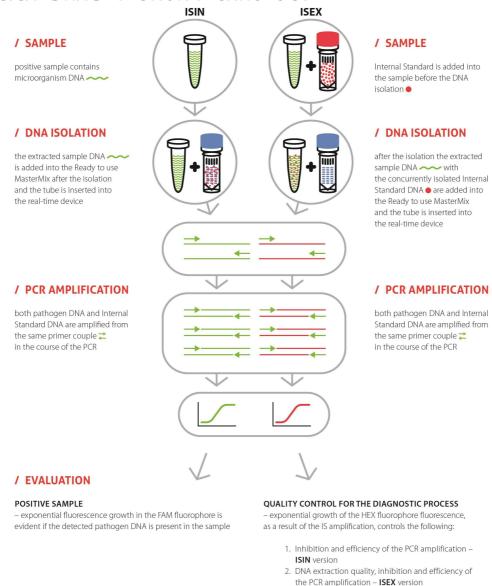
#### ISIN version

Internal Standard is included in the MasterMix tube. This PCR kit version enables PCR inhibition control.

#### ISEX version

Internal Standard is provided as independent item within the package. This PCR kit version enables both PCR inhibition control and nucleic acid purificaion process efficiency control.

#### MICROBIOLOGICAL DNA DETECTION TECHNOLOGY





### **USER MANUAL**

### SAMPLING AND SAMPLE STORAGE

Sampling of all sample types (plasma, cerebrospinal fluid, swabs, urine), except for blood, should be performed into sterile tubes without any transportation media and the samples should be transported within 12 hours at the temperature between +2 °C and +8 °C. Non-coagulating peripheral blood should be sampled into EDTA and transported to the laboratory at the temperature between +2 and +8 °C within 24 hours. In case of longer storage keep all samples frozen at the temperature between -85 and -10 °C.

#### **NUCLEIC ACID PURIFICATION**

Nucleic acid isolation should be performed by isolation kits available at the market according to protocols for the particular clinical material isolation. The manufacturer recommends the following isolation kits:

GeneProof PathogenFree DNA Isolation Kit croBEE NA16 Nucleid Acid Extraction System

When using the ISEX versions of the PCR kits the IS should be added directly into the sample at the beginning of the isolation process so that in the end 1  $\mu$ l of the resulting elution volume contains 0.1  $\mu$ l of the IS:

Elution volume	25 μl	50 μ1	100 μl	200 μl
Internal Standard	2,5 μl	5 μl	10 μl	20 μl

#### **PCR SETUP**

- 1. Add 30 µl of MasterMix into PCR tubes.
- 2. Add 10  $\mu$ l of the isolated nucleic acid sample or 10  $\mu$ l of Positive Control into the individual PCR tubes. The final reaction mix volume will be 40  $\mu$ l.

It is necessary to keep all components at +2°C to +8°C during the PCR preparation.

3. Close the tubes, centrifuge shortly, insert them into the device and let them amplify according to the following PCR profile.

Be very careful when handling the Positive Control or the clinical material, incorrect handling could result in contamination and the consequent impairment of the kit components or the MasterMix! The manufacturer is not responsible for the kit impairment due to incorrect handling.

#### **AMPLIFICATION PROFILE**

Step	Temperature	Time	Data collection	Cycles
1. Hold	37 °C	2 min		1
2. Hold	95 °C	10 min		1
	95 °C	5 s		
3. PCR	60 °C	40 s	FAM+HEX+Cy5	45
	72 °C	20 s		

## VALIDATED INSTRUMENTS

GeneProof PCR kits are designed for use with real-time devices from various manufacturers. This PCR kit has been validated with the following devices:

Applied Biosystems 7500 Real-Time PCR System

AriaMx Real-Time PCR System

Dx/CFX96™ Real-Time PCR Detection System

LightCycler ® 480

LineGene 9600\*

Mx3000P/3005P QPCR System

Rotor-Gene 3000\*, Rotor-Gene Q\*

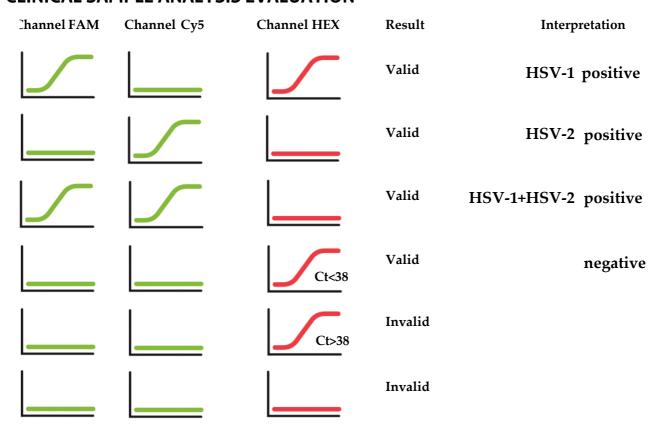
SLAN® Real-Time PCR System

GeneProof diagnostic kits are continually validated with various types of devices. Please request the current list at support@geneproof.com.



<sup>\*</sup> Validation applies to a device model providing detection in the following channels: FAM, HEX and Cy5.

### **CLINICAL SAMPLE ANALYSIS EVALUATION**



### QUANTITATIVE DETECTION EVALUATION

Use the following formula to calculate the virus concentration in copies/ml while taking into account the volume of material entering the isolation:

$$copy/ml = \frac{SC \times EV}{IV}$$

SC - Sample Concentration (copy/ $\mu$ l)

EV - Elution Volume (μl)

IV - Isolation Volume (ml)

You can use the calculator for pathogen concentration conversion at www.geneproof.com to make the calculation easier.

## WARNING

A single valid package insert for a specific kit is included in the package or to be requested for the particular lot from the manufacturer. The kit should be disposed of after use according to the current legal regulations considering the fact that the kit doesn't contain any dangerous, infectious or toxic components that would be subject to special safety regulations, and the packaging materials are made of paper and polypropylene. If you have any questions please contact our Customer Service.

Customer care and technical support

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