

**ASSAY NAME: HSV-2**

**Quantity: 100 x 20µL PCR reactions**

**2-plex assay: Herpes Simplex Virus 2 and human RPP30 DNA**

**SKU #: PNP-HSV2-D-QS-100**

**(RUO). Research Use Only. Not for use in Diagnostic Procedures.**

**SCOPE OF THIS PRODUCT INFORMATION SHEET (PIS):**

The oligonucleotide recipes are optimized for each instrument (QuantStudio, BioRad). The verification data presented in this PIS were performed using PNP-HSV2-D-QS-100 on a QuantStudio 7 Pro Real-Time System. The performance of the other SKUs on their corresponding instrument should be similar. Contact PCRassays.com if you are planning to use a different instrument.

**CONTENTS**

The primers and probes in the HSV-2 kit are provided in Tube 1 as a 20X concentrated working solution that detects 1 pathogen and a human control. The probes are designed as TaqMan<sup>5</sup> cleavage mechanism and thus the reaction requires a DNA polymerase with 5'-exonuclease activity (we recommend InhibiTaq Standard qPCR Master Mix).

**Table of Dyes used in this assay:**

Pathogen/Target	Dyes	Quencher	Refs.
HSV-2	<b>FAM</b>	BHQ-1	1,2
RPP30-DNA control	<b>TAMRA</b>	BHQ-2	3, 4

**ASSAY HANDLING**

The HSV-2 assay is shipped at ambient temperature, and should be stored at -20 °C. The assay should be kept on ice once thawed. Do not subject the enzyme to multiple freeze-thaw cycles.

Contamination should be avoided by using appropriate personal protective equipment (PPE), powder free gloves, aerosol barrier pipette tips, and a clean hood.

**ASSAY CONTENTS:**

**Tube 1:** 20X Primer/Probe mix for HSV-2 and hRPP30DNA.

**Tube 2:** Positive control (**Do NOT add to specimen unknowns**)

Positive control: 5000 copies/µl of synthetic 500 bp DNA fragments for HSV-2 and hRPP30DNA.

**Tube 3:** InhibiTaq Standard qPCR enzyme Mastermix (enough for 100 rxns. with 20 µL total volume). This is a custom formulation from Fortis Life Sciences to the specifications of PCRassays.com.



**EXPERIMENTAL**

Perform nucleic acid extraction/purification (recommended).

Set up your PCR reaction (20 µL) as follows on ice:

Component	Volume (µL)
InhibiTaq enzyme mastermix (2X)	10
Primer/Probe mix (20X)	1
Sample or positive control	2
Water (Molecular Biology Grade)	7

**Notes: To improve assay sensitivity, up to 9 µL of sample can be added (water volume adjusted accordingly) for a total reaction volume of 20 µL. For positive control rxns., add 2 µL of the solution from Tube 2.**

A PCR protocol was used for verification on a QuantStudio™ 7 Pro Real-Time System, with the following program:

Step	Thermocycling Protocol:
1	Incubate @ 95 °C for 2 minutes
2	Incubate @ 95 °C for 3 seconds
3	Incubate @ 55 °C for 22 seconds
4	Plate Read
5	Go to Step 2, repeat 44x more

## RESULT INTERPRETATION

After running the qPCR reaction, use the instrument software to determine the quantification cycle,  $C_q$  (or use  $C_T$  if your instrument does not have the capability to compute a  $C_q$ ). Fluorescence channels with a  $C_q < 38$  cycles, and final RFU  $>$ Threshold is considered “positive” or “+” in the Table below. The “Threshold” value for calling a PCR positive is dependent on the instrument model, well size, and sample volume; thus the user must determine the threshold that is appropriate for their method. For our QuantStudio 7 Pro with 96 well plate with 200  $\mu$ L wells and 20  $\mu$ L reaction volume, the average RFU was approximately 200,000 and we used a threshold of 20,000 for calling positives or “+” in the Table below.

HSV-2 FAM <sup>TM</sup>	hRPP30 TAMRA <sup>TM</sup>	Recommended Interpretation
—	—	The PCR reaction failed. Please repeat the experiment.
—	+	The sample contains human RPP30 DNA. The sample doesn't contain pathogen DNA.
+	—	The sample contains HSV-2 DNA. The sample may not contain human RPP30 DNA.
+	+	The sample contains HSV-2 DNA and human RPP30 DNA.

## VERIFICATION EXPERIMENTS

The HSV-2 assay verification was carried out as a 2-plex assay, which simultaneously detects DNA from Herpes Simplex Virus 2 and human RPP30 DNA, which serves as a positive extraction-control assay.

Experiments were performed in triplicate using the experimental procedure given above, but with different samples added to each reaction. The samples used for the verification experiments contained  $1 \times 10^4$  copies/reaction of 500 bp synthetic DNA constructs (from Twist Biosciences) harboring the regions of interest from the pathogen genomes, human RPP30 DNA gene, and human genomic DNA. The results of these experiments are shown in **Figure 1** and indicate that the 2-plex specifically detects the pathogen gene in the human genomic DNA matrix.

The limit of detection (LOD) was estimated by performing serial dilution experiments in triplicate (**Figure 2**). For dilution series only target construct was added. The results show a limit of detection (LOD)  $< 10$  copies/reaction.

**Conclusion:** The data in **Figure 1** indicate that the HSV-2 primers and probe are compatible with RPP30 DNA positive control primers and probe.

## NOTES

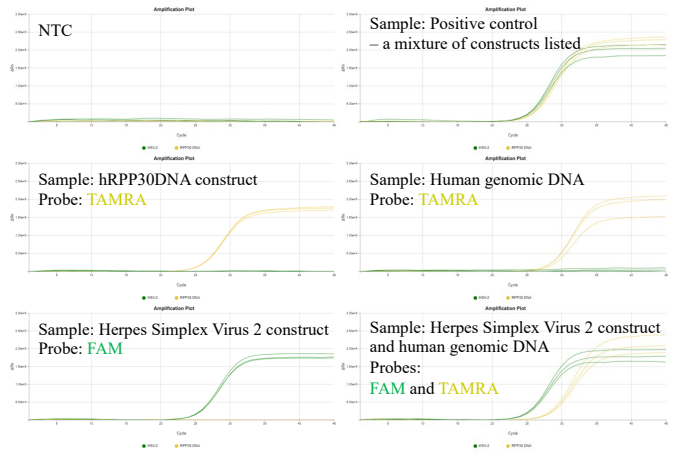
<sup>1</sup> FAM<sup>TM</sup> (Carboxyfluorescein) is a trademark of Life Technologies, Inc

<sup>2</sup> BHQ-1<sup>TM</sup> (Black Hole Quencher) is a trademark of Biosearch Technologies, Inc.

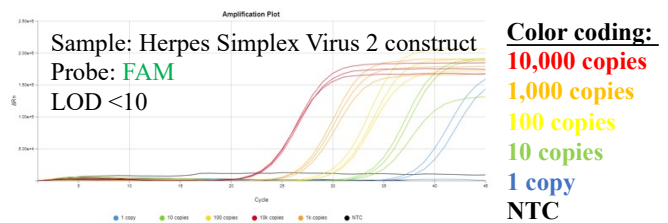
<sup>3</sup> TAMRA (Carboxyltetramethylrhodamine) is a trademark of Applera Corp.

<sup>4</sup> BHQ-2<sup>TM</sup> (Black Hole Quencher) is a trademark of Biosearch Technologies, Inc.

<sup>5</sup> “TaqMan” is a trademark of Roche Molecular Systems, Inc.



**Figure 1:** Verification experiments with single targets (given in text boxes for each panel). All sets of probes and primers are present in every reaction, but positive signal is only observed when the target(s) is present, indicating that the amplification is specific. Signal from single molecule events show  $C_q > 38$  and are considered negative.



**Figure 2:** Serial dilution experiments show LOD  $< 10$  molecules for the synthetic DNA construct of each target.

## CONTACT US

For assistance, please contact DNA Software using the link:  
<https://www.pcrassays.com/contact/>

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