MAYO CLINIC LABORATORIES

Test Definition: FHV7P

Herpesvirus 7 (HHV-7) DNA, Quantitative Real-Time PCR

Overview

Method Name
Real-Time PCR

NY State Available
Yes

Specimen

Specimen Type
Varies

Specimen Required
Submit only 1 of the following specimens:

Source is required

Whole Blood

Specimen Type: Whole Blood
Container/Tube: Lavender-top (EDTA)
Specimen Volume: 1 mL
Collection Information: Draw blood in a lavender-top (EDTA) tube(s) and send 1 mL whole blood refrigerated (DO NOT FREEZE).
Stability: Ambient 48 hours; Refrigerated 7 days

Serum

Specimen Type: Serum
Container/Tube: Red-top
Submission Container/Tube: 12x75 mm screw-capped vial
Specimen Volume: 1 mL
Collection Instructions: Draw blood in a plain red-top tube(s). Spin down and send 1 mL serum in a plastic, screw-capped vial. Send specimen refrigerated.
Stability: Ambient 48 hours; Refrigerated 7 days; Frozen 30 days

Plasma

Collection Container/Tube: Lavender-top (EDTA)

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Submission Container/Tube: 12x75 mm screw-capped vial
Specimen Volume: 1 mL
Collection Instructions: Draw blood in a lavender-top (EDTA) tube(s). Spin down and transfer 1 mL EDTA plasma into a plastic screw-capped vial. Send specimen refrigerated.
Stability: Ambient 48 hours; Refrigerated 7 days; Frozen 30 days

Specimen Minimum Volume
0.3 mL

Reject Due To

Thawing:	Warm reject; Cold OK
Other:	Samples collected in Lithium, Sodium Heparin, ACD or PPT

Specimen Stability Information

Specimen Type	Temperature	Time	Special Container
Varies	Refrigerated (preferred)	7 days	
	Ambient	48 hours	

Clinical & Interpretive

Reference Values
Not Detected

Performance

PDF Report
No

Day(s) Performed
Monday through Sunday

Report Available
3 to 5 days

Performing Laboratory Location
Quest Diagnostics

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Fees & Codes

Fees

- Authorized users can sign in to Test Prices for detailed fee information.
- Clients without access to Test Prices can contact Customer Service 24 hours a day, seven days a week.
- Prospective clients should contact their account representative. For assistance, contact Customer Service.

Test Classification

This test was developed and its analytical performance characteristics have been determined by Quest Diagnostics. It has not been cleared or approved by FDA. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.

CPT Code Information

87799

LOINC® Information

Test ID	Test Order Name	Order LOINC® Value
FHV7P	Herpes Virus 7 DNA, Quant RT-PCR	49397-3

Result ID	Test Result Name	Result LOINC® Value
Z5814	Source	31208-2
Z5815	Herpes Virus 7 DNA, QN PCR	49397-3
Z5816	Herpes Virus 7 DNA, QN PCR	Not Provided