Lab Dept:

Microbiology/Virology

Test Name: HHV7 PCR, QUANTITATIVE

General Information

Lab Order Codes:	HHV7P
Synonyms:	Human Herpes Virus 7 PCR; HHV-7; Herpesvirus 7 (HHV-7) DNA, Quantitative Real-Time PCR
CPT Codes:	87799 – Infectious agent detection by nucleic acid, not otherwise specified, quantification, each organism
Test Includes:	HHV7 detection by PCR reported in copies/mL and Log copies/mL
Logistics	
Test Indications:	Useful as an adjunct in the rapid diagnosis of HHV-7 infection. HHV-7 is closely related to HHV-6 and CMV, and can cause reactivation disease in transplant patients or other immune-compromised individuals.
Lab Testing Sections:	Microbiology - Sendouts
Referred to:	Mayo Clinic Laboratories forwarded to Quest Diagnostics (MML Test: FHV7P)
Phone Numbers:	MIN Lab: 612-813-6280
	STP Lab: 651-220-6550
Test Availability:	Daily, 24 hours
Turnaround Time:	3 – 5 days
Special Instructions:	N/A
Specimen	
Specimen Type:	Blood
Container:	Blood: Lavender top (EDTA) tube Alternate tube: Red NO gel
Draw Volume:	3 mL blood (minimum 0.3 mL whole blood or 1.2 mL to obtain plasma or serum)

Processed Volume:	1 mL (Minimum: 0.3 mL) whole blood or plasma/serum
Collection:	Routine blood collection
Special Processing:	Lab Staff: Centrifuge specimen, remove plasma aliquot within 2 hours of collection into a screw-cap plastic vial. Store and ship at refrigerated temperatures. Forward promptly.
	Specimen stable 7 days refrigerated (preferred), 30 days frozen or 48 hours ambient, with the exception of whole blood cannot be frozen.
Patient Preparation:	None
Specimen Rejection:	Specimens collected in Lithium/Sodium heparin; mislabeled or unlabeled specimens
Interpretive	
Reference Range:	Not Detected
Limitations:	The sensitivity of the assay is very dependent upon the quality of the specimen submitted. Results less than 500 copies/mL does not indicate that the patient is not infected with HHV7.
	The results obtained should be used in conjunction with clinical findings to make accurate diagnosis.
Methodology:	Real-Time Polymerase Chain Reaction (PCR)
References:	Mayo Clinic Laboratories (December 2023)
Updates:	 4/21/2004: Test moved from Eastern Virginia Medical School Department of Pathology to UTSW. 12/5/2006: Test moved to Focus as the UTSW test was not quantitative. 4/25/2011: Whole blood removed as a specimen type option. 3/22/2022: Added Log copies/mL, change forward to Quest. 12/27/2023: Removed outdated reference to Focus Technologies test limitations and interpretations, serum gel tube; added specimen stability; removed ACD as an acceptable blood container.