

TEST CHANGE

Cytomegalovirus by Qualitative PCR

0060040, CMVPCR

Specimen Requirements:	
Patient Preparation:	
Collect:	Bone marrow aspirate in lavender Lavender (EDTA) or pink; Pink (K2EDTA), amniotic or Serum Separator Tube (SST). Also acceptable: Amniotic fluid, bronchoalveolar lavage (BAL), CSF, ocular fluid, tissue, urine, or dried blood spot (DBS).
Specimen Preparation:	Separate serum or plasma from cells. Transfer 1 mL plasma, serum, whole blood, bone marrow, amniotic fluid, BAL, CSF, ocular fluid, or urine to a sterile container— (Min: 0.5 mL). Dried Blood Spot: Whole blood collected on newborn screening card (3/16 inch punch). Transport punch in an ARUP standard transport tube. Standard Transport Tube. Tissue: Transfer to a sterile container and freeze immediately.
Transport Temperature:	Frozen. Whole Blood or Bone Marrow: Refrigerated. Dried Blood Spot: Room temperature. All others: Frozen
Unacceptable Conditions:	Heparinized specimens, tissues in optimal cutting temperature compound. Saliva (Refer to ARUP test code 2008555, CMVPCR SAL.)
Remarks:	Specimen source is required.
Stability:	Ambient: 8 hours; Refrigerated: 72 hours; Frozen: 3 months Whole Blood or Bone Marrow: Ambient: 1 week; Refrigerated: 1 week; Frozen: 1 week Dried Blood Spot: Ambient: 90 28 days; Refrigerated: 8 days; Frozen: 8 days Tissue: Ambient: Unacceptable; Refrigerated: Unacceptable; Frozen: 3 months All others: Ambient: 8 hours; Refrigerated: 72 hours; Frozen: 3 months
Methodology:	Qualitative Polymerase Chain Reaction (PCR)
Performed:	Sun-Sat
Reported:	1-3 days
Note:	
CPT Codes:	87496
New York DOH Approval Status:	This test is New York DOH approved.



A nonprofit enterprise of the University of Utah
and its Department of Pathology

Effective Date: **November 13, 2023**

Interpretive Data:

~~This test was developed and its performance characteristics determined by ARUP Laboratories. It has not been cleared or approved by the US Food and Drug Administration. This test was performed in a CLIA certified laboratory and is intended for clinical purposes.~~

Reference Interval:

<u>Test Number</u>	<u>Components</u>	<u>Reference Interval</u>
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