Lab Dept: Microbiology/Virology

Test Name: ADENOVIRUS DNA PCR, PLASMA

General Information

Lab Order Codes: ADVPP

Synonyms: ADV PCR; Adenovirus Molecular Detection

CPT Codes: 87798 – Infectious agent detection by nucleic acid, not otherwise

specified; amplified probe technique, each organism

Test Includes: Adenovirus detection in plasma specimens by PCR. This test **does not**

include adenovirus culture. Refer to Viral Culture order.

Logistics

Test Indications: To detect Adenovirus DNA in clinical specimens.

Human adenoviruses cause a variety of diseases including pneumonia, cystitis, conjunctivitis, diarrhea, hepatitis, myocarditis, and encephalitis. In humans, adenoviruses have been recovered from almost every organ system. Infections can occur at any time of the year and in all age groups. Currently, there are 51 adenovirus serotypes that have been

grouped into 6 separate subgenera.

Culture is the gold standard for the diagnosis of adenovirus infection;

however, it can take up to 3 weeks to achieve culture results.

Serological tests have faster turnaround times, but can be less sensitive compared to culture. PCR offers a rapid, specific, and sensitive means

of diagnosis by detecting adenovirus DNA.

Lab Testing Sections: Microbiology/Virology - Sendouts

Referred to: Mayo Clinic Laboratories (Mayo Test: LCADP)

Phone Numbers: MIN Lab: 612-813-6280

STP Lab: 651-220-6550

Test Availability: Daily, 24 hours

Turnaround Time: 2 – 5 days, set up Monday, Wednesday and Friday

Special Instructions: Viral culture is the standard method and is recommended along with

PCR.

Specimen

Specimen Type: Blood

Container: Lavender top (EDTA) tube

Draw Volume: 3 mL (Minimum: 0.9 mL) blood

Processed Volume: 1 mL (Minimum: 0.3 mL) plasma

Collection: Routine blood collection

Special Processing: Lab staff: Centrifuge specimen, remove plasma aliquot into a screw-

capped, plastic vial. Ship and store specimen at refrigerated

temperatures.

Specimen stable refrigerated (preferred) or frozen for 7 days.

Sample Rejection: Mislabeled or unlabeled sample; QNS; gross contamination; gross

hemolysis

Interpretive

Reference Range: None detected

A positive report indicates the presence of adenovirus. A negative report does not rule out the presence of adenoviruses because organisms may be present at levels below the detection limits of this

assay.

Limitations: • Asymptomatic shedding may occur up to 18 months after infection.

Test results should be interpreted in conjunction with other clinical

information.

• A negative result **does not** rule out the presence of PCR inhibitors in the patient specimen or adenoviral nucleic acid in concentrations below

the level of detection of the assay.

• Test results should be used as an aid in diagnosis and should not be

considered diagnostic in themselves.

Methodology: Real-Time Polymerase Chain Reaction (PCR)/DNA probe hybridization

References: Mayo Clinic Laboratories January 2018

Updates: 11/15/2023: Biennial review, removed frozen as reason for rejection.