

# **TEST DIRECTORY UPDATE**

#### **JULY 2022 IMMEDIATE ACTION**

Update Notes	

<b>Update Summary</b>			
	<b>New Test Activation</b>	7/21/2022	MOPOX - "Monkeypox Virus DNA, QL PCR"
	Update Existing Test	7/19/2022	DHEA - "DHEA (DEHYDROEPIANDROSTERONE), UNCONJUGATED, LC/MS/MS"

<b>New Test Activation</b>			
Effective Date	7/21/2022		
Name	Monkeypox Virus DNA, QL PCR		
Code	MOPOX		
CPT Code(s)	87798 x 2		
Notes			
Specimen Requirements			
Specimen Required	Collect: Lesion swab  Specimen Preparation: Swab the pustule/lesion vigorously and place the swab in 3.0 mL Viral transport medium (VTM) tube. Each individual specimen submitted for Monkeypox virus testing should be accompanied by its own separate requisition and transported in its own sealed bag. Multiple specimens collected on a single patient should be submitted separately.  Minimum Volume: 0.5 mL  Transport Temperature: Frozen  Additional Collection Information: https://www.cdc.gov/poxvirus/monkeypox/clinicians/prep-collection-specimens.html		
Alternate Specimen	Viral culture media (VCM), Universal Transport Media (UTM)		
Rejection Criteria	Calcium Alginate swabs, Wooden swabs		
Stability	Room temperature: Unacceptable Refrigerated: 7 days Frozen: 30 days		

LAST EDITED: 2022-07-21 PAGE 1 OF 3



# **TEST DIRECTORY UPDATE**

## **JULY 2022 IMMEDIATE ACTION**

Performing Information					
Methodology	Real-Time Polymerase Chain Reaction (PCR)				
Reference Range	Orthopoxvirus DNA, QL PCR Not detected  Monkeypox Virus DNA, QL PCR Not detected				
Performed Days	Sunday - Saturday				
Turnaround Time	3 - 5 days				
Performing Laboratory	Quest SJC				
Interface Information					
Legacy Code <sup>1</sup>	MOPOX				
Interface Order Code	3400644				
Result Code	Name	LOINC Code	AOE/Prompt <sup>2</sup>		
3400645	Patient Race:	Not available	Yes		
3400646	Ethnicity:	Not available	Yes		
3400647	Orthopoxvirus DNA, QL PCR	Not available	No		
3400648	Monkeypox Virus DNA, QL PCR	Not available	No		

LAST EDITED: 2022-07-21 PAGE 2 OF 3



#### LABORATORY REPORT

Example Client, XYZ123 1234 Warde Road Ann Arbor MI 48108 **EXAMPLE, REPORT** 

WX0000003039 M 12/05/1988 33 Y

Referral Testing

Collected: 07/20/2022 11:01 Received: 07/21/2022 12:38

Test Name Result Flag Ref-Ranges Units Site

Monkeypox Virus DNA, QL PCR

Patient Race: Unknown QCRL
Ethnicity: Unknown QCRL
Orthopoxvirus DNA, QL PCR NOT DETECTED QCRL
Monkeypox Virus DNA, QL PCR NOT DETECTED

This specimen is negative for Monkeypox virus DNA. If clinically indicated, consider collecting another specimen.

REFERENCE RANGE: NOT DETECTED

The Monkeypox Virus DNA Real-time PCR is intended for the qualitative detection of non-variola Orthopoxviruses and Monkeypox virus (West African clade) DNA using swabs from human pustular or vesicular rash specimens. These results must be used in conjunction with clinical observations and epidemiological risk factors.

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

Test Performed at: Quest Diagnostics Nichols Institute 33608 Ortega Highway San Juan Capistrano, CA 92675-2042

Performing Site:

QCRL: QUEST DIAGNOSTICS REFERENCE LAB CAPISTRANO 33608 Ortega Highway San Juan Capistrano CA 92675

I Maramica MD, PhD



# **TEST DIRECTORY UPDATE**

## **JULY 2022 IMMEDIATE ACTION**

Update Existing Test			
Effective Date	7/19/2022		
Name	DHEA (Dehydroepiandrosterone), Unconjugated, LC/MS/MS		
Code	DHEA		
Interface Order Code	3700510		
Legacy Code	3700510		
Notes	Updates to rejection criteria, transport temperature and stability.		
Required Testing Change	9S		
Specimen Required	Patient Preparation: Overnight fasting is preferred.  Collect: Red top  Specimen Preparation: Centrifuge, separate serum from cells and send 0.5 mL serum in a screw capped plastic vial.  Minimum Volume: 0.3 mL  Transport Temperature: Frozen		
Alternate Specimen	Plasma: EDTA (lavender or Dark Blue), sodium heparin or lithium heparin (green top)		
Rejection Criteria	Serum Separator Tube (SST), gross hemolysis		
Stability	Room temperature: 6 hours Refrigerated: 72 hours Frozen: 90 days		

LAST EDITED: 2022-07-21 PAGE 3 OF 3