ARUP LABORATORIES
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Jonathan R. Genzen, MD, PhD, Chief Medical Officer

Client: Example Client ABC123
123 Test Drive
Salt Lake City, UT 84108
UNITED STATES

Physician: Doctor, Example

Patient: Patient, Example

DOB	10/22/2024
Gender:	Unknown
Patient Identifiers:	01234567890ABCD, 012345
Visit Number (FIN):	01234567890ABCD
Collection Date:	00/00/0000 00:00

Human Immunodeficiency Virus Types 1 and 2 (HIV-1/2) Antibody Differentiation, Supplemental, With Reflex to HIV-1/HIV-2 by Qualitative NAAT

ARUP test code 2012669

HIV-1/2 Ab Differentiation Immunoassay	See Below INTERPRETIVE INFORMATION: HIV-1/2 Ab Diff, Supplemental This assay should not be used for blood donor screening, associated re-entry protocols, or for screening Human Cell, Tissues, and Cellular and Tissue-Based Products (HCT/P).	
HIV-1 Antibody	Negative	(Ref Interval: Negative)
HIV-2 Antibody	Negative	(Ref Interval: Negative)
HIV Serologic Interpretation	HIV Abs Neg HIV Serologic Interpretation - Indeterminate HIV nucleic acid testing is recommended. The specimen is negative for both HIV-1 and HIV-2 antibodies by the HIV-1/HIV-2 antibody differentiation assay. Qual HIV-1/HIV-2 nucleic acid testing will be performed on this specimen.	

Human Immunodeficiency Virus 1 and 2 (HIV-1/HIV-2) by Qualitative NAAT

ARUP test code 3017779

HIV-1 Qualitative by NAAT	Not Detected	(Ref Interval: Not Detected)
HIV-2 Qualitative by NAAT	Not Detected	(Ref Interval: Not Detected)

H=High, L=Low, *=Abnormal, C=Critical

INTERPRETIVE INFORMATION: HIV 1/2 Qualitative

This test detects human immunodeficiency virus type 1 (HIV-1) RNA from Group M, N and O subtypes and human immunodeficiency virus type 2 (HIV-2) RNA from Group A and B subtypes; it does not detect HIV proviral DNA. A result of "Not Detected" does not rule out HIV-1/HIV-2 RNA concentrations below the limit of detection of the assay or the presence of inhibitors in the patient specimen. The diagnosis of HIV infection should not be made based solely on a single HIV test result. Diagnosis requires repeat and confirmatory testing as recommended by U.S. Health and Human Services guidelines. Improper specimen handling can cause false negatives or contamination.

This assay should not be used for blood donor screening, associated reentry protocols, or for screening human cells, tissues, and cellular- and tissue-based products (HCT/P).

VERIFIED/REPORTED DATES

Procedure	Accession	Collected	Received	Verified/Reported
HIV-1/2 Ab Differentiation Immunoassay	24-297-103817	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
HIV-1 Antibody	24-297-103817	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
HIV-2 Antibody	24-297-103817	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
HIV Serologic Interpretation	24-297-103817	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
HIV-1 Qualitative by NAAT	24-297-103817	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
HIV-2 Qualitative by NAAT	24-297-103817	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00

END OF CHART