

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

Physician: Doctor, Example

**Patient: Patient, Example**

**DOB:** 4/26/1971  
**Gender:** Male  
**Patient Identifiers:** 01234567890ABCD, 012345  
**Visit Number (FIN):** 01234567890ABCD  
**Collection Date:** 00/00/0000 00:00

**Human Immunodeficiency Virus 1 (HIV-1) by Quantitative NAAT with Reflex to HIV-1 Drug Resistance by Next Generation Sequencing**

ARUP test code 3000870

HIV-1 Qnt by NAAT (copies/mL) 7,130 cpy/mL

HIV-1 Qnt by NAAT (log copies/mL) 3.85 log cpy/mL

HIV-1 Drug Resistance by Next Generation Sequencing will be added.

HIV-1 Qnt by NAAT Interp

**Detected \* (Ref Interval: Not Detected)**

INTERPRETIVE INFORMATION: HIV-1 by Quantitative NAAT, Plasma

Normal range for this assay is "Not Detected".  
The quantitative range of this assay is 1.47-7.00 log copies/mL (30-10,000,000 copies/mL).

An interpretation of "Not Detected" does not rule out the presence of inhibitors or HIV-1 RNA concentration below the level of detection of the assay. Care should be taken in the interpretation of any single viral load determination. The clinical significance of changes in HIV-1 RNA concentration has not been fully established; however, a change of 0.5 log copies/mL may be significant.

This assay should not be used for blood donor screening, associated re-entry protocols, or for screening Human Cell, Tissues and Cellular Tissue-Based Products (HCT/P).

**Human Immunodeficiency Virus 1 Drug Resistance by Next Generation Sequencing**

ARUP test code 3003853

HIV-1 Drug Resistance by NGS See Note

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

**Integrase Strand Transfer Inhibitor Drug Class**  
Bictegravir, BIC Susceptible  
Dolutegravir, DTG Susceptible  
Elvitegravir, EVG Susceptible  
Raltegravir, RAL Susceptible

IN drug resistance mutations identified: None

IN accessory resistance mutations identified: None

IN additional mutations identified: E10D, E11D, V31I, V32I, M50T, I72V, I84M, L101I, K111R, I191D, G192W, V201I, K211N, L234I, R269K

**Protease Inhibitor Drug Class**  
Atazanavir, ATV Susceptible  
Darunavir, DRV Susceptible  
Fosamprenavir, FPV Susceptible  
Indinavir, IDV Susceptible  
Lopinavir, LPV Susceptible  
Nelfinavir, NFV Susceptible  
Saquinavir, SQV Susceptible  
Tipranavir, TPV Susceptible

PR drug resistance mutations identified: None

PR accessory resistance mutations identified: None

PR additional mutations identified: G16E, R41K, R57K, I62V, L63T, T74S, V77I

**Nucleoside Reverse Transcriptase Inhibitor Drug Class**  
Abacavir, ABC Susceptible  
Zidovudine, AZT Susceptible  
Stavudine, D4T Susceptible  
Didanosine, DDI Susceptible  
Emtricitabine, FTC Susceptible  
Lamivudine, LMV Susceptible  
Tenofovir, TDF Susceptible

NRTI drug resistance mutations identified: None

**Non-nucleoside Reverse Transcriptase Inhibitor Drug Class**  
Doravirine, DOR Susceptible  
Efavirenz, EFV Susceptible  
Etravirine, ETR Susceptible  
Nevirapine, NVP Susceptible  
Ralpivirine, RPV Susceptible

NNRTI drug resistance mutations identified: None

RT accessory resistance mutations identified: None

RT additional mutations identified: V35T, K122P, D123A, I135R, I178L, R211K, V245D, V245A, E248T, A272P, T286A, Q334N, A360T, A376T, K390R, E399D, A400T, V435I

HIVGenotyper software version: 1.0.0.0

Stanford HIV Drug Resistance Database Version: HIVDB\_8.9-1

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

**INTERPRETIVE INFORMATION: HIV-1 Drug Resistance by NGS**

This assay predicts HIV-1 resistance to protease inhibitors, nucleoside reverse transcriptase inhibitors, non-nucleoside reverse transcriptase inhibitors and integrase inhibitors. The protease gene, integrase gene and the reverse transcriptase gene of the viral genome are sequenced using Next Generation Sequencing. Drug resistance is assigned using the Stanford hivdb database.

This test should be used in conjunction with clinical presentation and other laboratory markers. A patient's response to therapy depends on multiple factors, including patient adherence, percentage of resistant virus population, dosing, and drug pharmacology issues.

This test detects populations down to 10 percent of the total population which may account for resistance interpretation differences between methods. Some insertions or deletions may be difficult to detect using this software.

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.

**EER HIV-1 Drug Resistance by NGS**

See Note

Authorized individuals can access the ARUP Enhanced Report using the following link:

**VERIFIED/REPORTED DATES**

Procedure	Accession	Collected	Received	Verified/Reported
HIV-1 Qnt by NAAT (copies/mL)	23-088-119256	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
HIV-1 Qnt by NAAT (log copies/mL)	23-088-119256	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
HIV-1 Qnt by NAAT Interp	23-088-119256	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
HIV-1 Drug Resistance by NGS	23-088-119256	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
EER HIV-1 Drug Resistance by NGS	23-088-119256	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00

**END OF CHART**

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

Unless otherwise indicated, testing performed at: