



INTERPATH
LABORATORY

Interpath Laboratory, Inc.
Test File Update

As an Interpath customer who receives electronic results or sends electronic orders you may need to be notified when we update our Service Manual. Although we try to keep these changes to a minimum, laboratory medicine is an evolving industry requiring changes to our technology from time to time. Depending on the requirements of your EMR or Hospital Information System you may be required to make similar changes to your system in order to correctly process inbound electronic results and create outbound electronic orders.

If you are uncertain that you are required to update your system we recommend that you contact your vendor for more information. As your laboratory service provider we are available to participate in the discussion with your vendor so that you clearly understand the impact of these changes.

Included in this email:

- This cover letter with a summary of the changes
- Microsoft Word® Document with the detail of these changes to our Service Manual
- Interpath Master Order/Result Compendium

Additional information including our most recent Service Manual and additional contact information can be found at www.interpathlab.com

Effective Date: March 23, 2026



Order Code	Test Name	NC Name Change	CC Component Change	CPT CPT Change	SRC Specimen Requirements Change	RRC Reference Range Change	NT New Test	DT Discontinued Test	AOE Ask on Order Entry Questions
90014	Epstein-Barr Virus by Quantitative NAAT, Plasma								◆
5290	Primary High-Risk HPV (Self-Collected Vaginal)						◆		
1749	Unexplained Fatigue Panel				◆				



90014 Epstein-Barr Virus by Quantitative NAAT, Plasma
Please take note of changes to Ask-on-Order-Entry questions.

AOE

5290 Primary High-Risk HPV (Self-Collected Vaginal)

NT

Specimen:	
Collect:	Self-Collect Vaginal Swab in ThinPrep PreservCyt Media Also Acceptable Self-Collect Vaginal Swab in Vaginal Self-Collect Kit
Submit:	20 mL (Min:10 mL) Self-Collect Vaginal Swab in ThinPrep PreservCyt Media. Submit Ambient.
Special Handling:	State source. Follow manufacturer's FDA labeling for self-collection. FDA-cleared for self-collected vaginal specimens obtained in a healthcare setting for primary HPV screening in adults 25 years or older with a cervix. Not validated for medico-legal use, including suspected sexual abuse. Only self-collection using FDA-approved/cleared devices for Cobas® HPV is permitted.
Rejection Criteria:	Any specimen not using FDA-approved vaginal self-collect device; any specimen not submitted in ThinPrep PreserveCyt Media; improper labeling; specimen >30 days old; frozen specimens; leakage; insufficient quantity; name discrepancies; patient <25 years; patient without a cervix; non-vaginal source; excessively bloody specimen
Stability:	Ambient: 1 Month(s); Refrigerated: 1 Month(s); Frozen: Unacceptable; Incubated: Unacceptable
Methodology:	Real-Time Polymerase Chain Reaction
Performed:	Mon-Fri
Reported:	1-4 Day(s)
CPT Codes:	87626
Interpretive Data:	Please see report for interpretive data.
Components:	5291 - HPV16 5293 - HPV, HIGH-RISK, OTHER 5292 - HPV18 5294 - ANATOMICAL SOURCE

Please note new test.



1749 Unexplained Fatigue Panel

SRC

Specimen:	
Collect:	Two SST Also Acceptable One Standard Transport Tube
Submit:	6 mL (Min:4 mL) Serum in Standard Transport Tube. Submit Refrigerated. Submit in a Standard Transport Tube.
Special Handling:	Allow specimen to clot completely at room temperature Avoid Repeated Freeze/Thaw Cycles Separate from cells ASAP If positive ANA Titer, the following EIA tests will be performed: Centromere Antibody, Jo-1 Antibody, U1-RNP Antibody, dsDNA Antibody, Scleroderma 70-S Antibody, Smith DP-S Antibody, Ribosomal-P Antibody, Sjogren Antibodies A/RO & B/LA. Additional charges will apply. If the dsDNA Antibody is Positive, then the dsDNA AB Titer (Test# 2866) will be performed. Additional charges will apply.
Rejection Criteria:	Grossly Hemolyzed Samples Heat inactivated Lipemic Samples Microbially Contaminated Particulate matter
Stability:	See Individual Components
Methodology:	See Individual Components
Performed:	Varies
Reported:	1-4 Day(s)
CPT Codes:	86038 86200 86235 86376 86431 86800 86258x2 86364x2
Interpretive Data:	Please see report for interpretive data.
Components:	2859 - CCP ANTIBODY IGG 2752 - TISSUE TRANSG.IgA 2726 - GLIADIN (DGP)-IgG 2755 - ANTI-THYROGLOBIN 2337 - RHEUMATOID FACTOR 2727 - TISSUE TRANSG.IgG 2725 - GLIADIN (DGP)-IgA 2868 - dsDNA ANTIBODY 2754 - ANTI-T PEROXIDASE 2867 - ANA HEP2 TITER 2869 - ANA HEP2 PATTERN 2510 - ANA REFLEX

Please note specimen requirement changes.