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Test NameTest CodeEffective Date4th Generation HIV Test ResultsB688October 23, 2023

Please be advised that 4th Generation HIV Test Results have been updated, and will appear differently on reports.

Previously the Elecyses Duo HIV Ag/Ab test result was reported as Reactive or Non-Reactive. The Elecyses Duo HIV Ag/Ab now includes HIV Ag/Ab test result Reactive or Non/ Reactive, as well as the individual test components for HIV-1 p24 (the antigen) and HIV 1/2 (the antibodies) as Reactive or Non- Reactive. HIV-1 p24 antigen and HIV 1/2 antibodies results will appear under the final HIV Ag/Ab result.

This format of reporting is mandated by the NYS Department of Health. Providing the individual results as well as the final interpretation will allow healthcare providers to know if the sample represents an early HIV infection. (HIV-1 p24 reactive and confirmed on the Qualitative HIV-1 Aptima TMA).

Please see sample report below, and contact your dedicated Account Executive or call Customer Service if you have any questions.

MISCELLANEOUS					
Test	Result	Abnormal	Reference	Units	Rpt Date
HIV Ag/Ab	Non-Reactive	!	Non-Reactive		10/24/23
Assay Information:	Assay for the detecti	on of HIV-1 p	24 antigen and		
	antibodies to Human Immunodeficiency Virus, HIV-1				
	(groups M and O) and HIV-2				
	Method: Electrochemiluminescence Elecsys Duo				
HIV-1 p24 Ag	Non-Reactive	:	Non-Reactive		10/24/23
HIV 1 +2 Ab	Non-Reactive	!	Non-Reactive		10/24/23
	Final Report				

Activated Protein C Resistance

5704

October 17, 2023

Please be advised that APC Resistance (Test code 5704) was retired. APC Resistance-modified (Test code 5705) can be ordered as an alternative test for APC resistance, which is more indicative of Factor V Leiden mutation when positive.

ACTH 0507 October 2023

Due to a change in instrumentation, test information for ACTH (Test Code 0507) has been updated. Please refer to the table below for details.

	Previous Test Information	New Test Information
Methodology	Chemiluminescence	Electrochemiluminescence immunoassay
Reference Range	<46pg/mL	7.2-63.3 pg/mL

Antinuclear Ab (ANA) W/ Reflex

0038, A233

November 1, 2023

Turn Around Time* for Antinuclear Ab (ANA) W/ Reflex has been updated (Test Codes 0038 and A233). Please refer to the table below for details.

	Previous Test Information	New Test Information
Turn Around Time*	2 days	2-4 days

^{*} TAT is based upon receipt of the specimen at the laboratory.

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Test NameTest CodeEffective DateAnti-CCP3747November 2023

Due to a change in instrumentation, test information for **Anti-CCP** (**Test Code 3747**) has been updated. Please refer to the table below for details.

	Previous Test Information	New Test Information
Methodology	Chemiluminescence	Electrochemiluminescence immunoassay
Reference Range	<4.0U/mL	< 17 U/mL

Cortisol Multiple, see below October 2023

Due to a change in instrumentation, test information for Cortisol (Test Codes 0066, 0067, 0374, 0900, 1495, 1496, 1851, 3314, and 3315) has been updated. Please refer to the table below for details.

	Previous Test Inforn	mation	New Test Information	
Stability	7 days, 90 days from	zen	4 days refrigerated, 12 months	frozen
Methodology	Chemiluminescence	е	Electrochemiluminescence imm	nunoassay
Reference Range	AM:	5-25ug/dL	Morning Hours 6 am - 10am	4.9 – 19.5 ug/dL
	PM:	2.5-12.5 ug/dL	Afternoon Hours 4pm – 8pm	2.5 – 11.9 ug/dL
	Random:	2.5-25.0ug/dL	Random	2.5 – 19.5 ug/dL
			Cortisol I0 min	Not Established
			Cortisol 30min	Not Established
			Cortisol 60min	Not Established
			#2	Not Established
			#3	Not Established

ER/ PR/ DNA/ KI-67/ HER2 - Reflex to HER2 by FISH

5408

Immediately

Please be advised that ER/ PR/ DNA/ KI-67/ HER2 - Reflex to HER2 by FISH (Test code 5408) will be retired. ER/ PR/ KI-67/ HER2 - Reflex to HER2 by FISH. (Test Code 5407) can be ordered as an alternative test.

ICD-10 Reminder All October 1, 2023

As we continue through the Fall season, please remember to review the ICD-10 diagnosis code updates effective October 1. This update included additions, deletions, and revisions. Most recently, we have seen removal of "unspecified" diagnosis codes with addition of more specific diagnosis codes which require 4th or 5th digit specificity. Since incorrect or outdated diagnosis coding can affect payment as much as incorrect procedure coding, it's important to update your code files and references for diagnosis codes your practice uses most often.

Please download the ICD-10 guide from the CDC here: https://www.cdc.gov/nchs/icd/Comprehensive-Listing-of-ICD-10-CM-Eiles.htm, or refer to the on-line browser tool to search diagnosis codes here:

https://www.cdc.gov/nchs/icd/icd10cm_browsertool.htm

Interferon-Alpha, EIA 3997 November 2023

Please be advised that Interferon-Alpha, EIA (Test code 3997) was retired by our referral lab, with no alternate test recommended.

Interferon-Beta, IgG 3541 November 2023

Please be advised that Interferon-Beta, IgG (Test code 3541) was retired by our referral lab, with no alternate test recommended.

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Test NameTest CodeEffective DateMI-2 Antibodies6328November 2023

Please be advised that MI-2 Antibodies (Test code 6328) was retired by our referral lab, with no alternate test recommended.

MYOSITIS ASSESSR + J0-1

2676

November 2023

Please be advised that MYOSITIS ASSESSR + J0-1 (Test code 2676) will be retired. Extended Myositis Panel (Test code TQ33) can be ordered as an alternative test. Please see alternate test information on next page.

	Previous Test Information	Alternate Test Information
Primary Container	SST - SST Tube	SST - SST Tube
Minimum Volume	2 ml	3 aliquots of 1mL
Turn Around Time*	15 Days	9-20 Days
Transportation Temp	Refrigerate	Refrigerated
Stability	7 Days	Ambient: 48 hours; Refrigerated: 2 weeks; Frozen: 1 month
Methodology	Immunoassay	ELISA
Reference Range	PL 7 AUTOANTIBODIES – Detected/Not Detected PL 12 AUTOANTIBODIES - Detected/Not Detected MI 2 AUTOANTIBODIES - Detected/Not Detected KU AUTOANTIBODIES - Detected/Not Detected EJ AUTOANTIBODIES - Detected/Not Detected OJ AUTOANTIBODIES - Detected/Not Detected SRP AUTOANTIBODIES - Detected/Not Detected SRP AUTOANTIBODIES - Detected/Not Detected JO-1 ANTIBODY - <1.0 NEGATIVE AI	Smith/RNP (ENA) Ab, IgG - 19 Units or less SSA-52 (Ro52) (ENA) Antibody, IgG - 40 AU/mL or less Jo-1 (Histidyl-tRNA Synthetase) Ab, IgG - 40 AU/mL or less PM/ScI 100 Antibody, IgG - Negative Mi-2 (nuclear helicase protein) Antibody - Negative PL-7 (threonyl-tRNA synthetase) Antibody - Negative PL-12 (alanyl-tRNA synthetase) Antibody - Negative P1-55/140 Antibody - Negative EJ (glycyl-tRNA synthetase) Antibody - Negative Ku Antibody - Negative SRP (Signal Recognition Particle) Ab - Negative OJ (isoleucyl-tRNA synthetase) Antibody - Negative SSA-60 (Ro60) (ENA) Antibody, IgG - 40 AU/mL or less Fibrillarin (U3 RNP) Ab, IgG - Negative SAE1 (SUMO activating enzyme) Ab - Negative MDA5 (CADM-140) Ab - Negative NXP2 (Nuclear matrix protein-2) Ab - Negative TIF-1 gamma (155 kDa) Ab - Negative
Collection Instructions	SST: Fill tube, invert gently 5 times, label with patient name, let stand for minimum of 30 minutes, maximum of 1 hr, spin for 10-15 minutes	Separate from cells ASAP or within 2 hours of collection. Transfer three 1 mL serum aliquots to Standard Transport Tubes.
Profile Components	PL 7 AUTOANTIBODIES PL 12 AUTOANTIBODIES MI 2 AUTOANTIBODIES KU AUTOANTIBODIES EJ AUTOANTIBODIES OJ AUTOANTIBODIES SRP AUTOANTIBODIES JO-1 ANTIBODY	SSA-52 (Ro52) (ENA) Antibody, IgG SSA-60 (Ro60) (ENA) Antibody, IgG Smith/RNP (ENA) Antibody, IgG Jo-1 Antibody, IgG
CPT Code(s)**	83516x5, 86235x3	83516x8, 86235x6, 84182x4
List Price	\$630.00	\$520.00
Clinical Utility (If applicable)		May be useful for differential evaluation of polymyositis, dermatomyositis, necrotizing autoimmune myopathy, or overlap syndromes associated with connective tissue disease

Parainfluenza Ab types 1-3

0946

November 2023

Please be advised that **Parainfluenza Ab types 1-3 (Test code 0946)** was retired by our referral lab, with no alternate test recommended.

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Test NameTest CodeEffective DateNatural Killer Cells, FunctionalTN20November 2023

Please be advised that **Natural Killer Cells, Functional (Test code TN20)** was retired by our referral lab, with no alternate test recommended.

Neutrophil Cytoplasmic AB/ ANCA

0636 November 1, 2023

Turn Around Time* for Neutrophil Cytoplasmic AB/ ANCA (Test Code 0636) has been updated. Please refer to the table below for details.

	Previous Test Information	New Test Information
Turn Around Time*	3 days	4 days

Phosphatidylcholine (G,A,M), Non-NY

3114

November 2023

Please be advised that **Phosphatidylcholine** (G,A,M), Non-NY (Test code 3114) was retired by our referral lab, with no alternate test recommended.

Thyroglobulin, Serum

0577

November 2023

Due to a change in test instrumentation, test information for **Thyroglobulin, Serum (Test Code 0577)** has been updated. Please refer to the table below for details.

	Previous Test Information	New Test Information
Turn Around Time*	2 days (MON,TUE,WED,THUR,FRI,SAT,SUN)	3 days (MON,TUE,WED,THUR,FRI)
Stability	3 days, 2 months frozen	2 days, 7 days frozen
Reference Range	<55.0 ng/mL	1.59 – 50.03 ng/mL

Unconjugated Estriol

3116

November 2023

Due to a change in test instrumentation, test information for **Unconjugated Estriol (Test Code 3116)** has been updated. Please refer to the table below for details.

	Previous Test Information		New Test Information	
Turn Around Time*	2 days (MON,TUE,WED,THUF	R,FRI,SAT,SUN)	3 days (MON,TUE,WED,TH	UR,FRI)
Reference Range	Gestational Week	Range (ng/mL)	Gestation Week	Expected Range (ng/mL)
	27	2.3-6.4	12	0.30-1.00
	28	2.3-7.0	14	0.40-1.60
	29	2.3-7.7	16	1.40-6.50
	30	2.4-8.6	18	1.60-8.50
	31	2.6-9.9	20	2.10-13.00
	32	2.8-11.4	22	2.70-16.00
	33	3.0->12.0	26	3.00-18.00
	34	3.3->12.0	32	4.60-23.00
	35	3.9->12.0	36	7.20-29.00
	36	4.7->12.0	40	8.00-39.00
	37	5.6->12.0	Male/Non-Pregnant Female	<2.0
	38	6.6->12.0		
	39	7.3->12.0		
	40	7.6->12.0		
	Male/Non-Pregnant Female	<2.0		

^{*} TAT is based upon receipt of the specimen at the laboratory.

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Test Name Test Code Effective Date

REMINDERS

Obstetric Panels Multiple November 1, 2023

BioReference® and GenPath® strive to ensure that you have the most up to date and guideline driven testing options. Coming in November, we will be making updates to our **Obstetric/Prenatal Panel** (**Test Code 0008**), **Obstetric AMA Panel** (**test code 0010**), and **Obstetric Panel** (**Test Code 7307**) to reflect current ACOG and CDC recommendations. Please see the tables below for the anticipated test names changes and test components updates.

	Test Code 0008 - Previous Test Information		Test Cod	Test Code 0008 - New Test Information	
Test Name	Obstetric/Prenatal I		Obstetric	ACOG Panel w/HIV	
Test Components	0039	ANTIBODY SCREEN	0039	ANTIBODY SCREEN	
	0053	CBC WITH DIFF	0053	CBC WITH DIFF	
	0142	RPR	0142	RPR	
	0156	ABO/Rh BLOOD TYPE	0156	ABO/Rh BLOOD TYPE	
	0973	RUBELLA, IGG	0973	RUBELLA, IGG	
			B688	HIV	
			B125	HEP C ANTIBODY W/RFX RT PCR	
			0800	URINE CULTURE	
				HEP B SURF AG W/CONF, IF POS, RFX to DNA, QUANT,	
			TP85	PRENATAL	

	Test Code 0010 - Previous Test Information		Test Code	e 0010 - New Test Information	
Test Name	Obstetric AMA Panel		Obstetric A	Obstetric ACOG Panel w/HIV/CT/GC/Trich	
Test Components	0039	ANTIBODY SCREEN	0039	ANTIBODY SCREEN	
	0053	CBC WITH DIFF	0053	CBC WITH DIFF	
	0142	RPR	0142	RPR	
	0156	ABO/Rh BLOOD TYPE	0156	ABO/Rh BLOOD TYPE	
	0973	RUBELLA, IGG	0973	RUBELLA, IGG	
	0106	HEP B SURFACE ANTIGEN		HEP B SURF AG W/CONF, IF POS, RFX to DNA, QUANT,	
			TP85 †	PRENATAL	
			B688	HIV	
			B125	HEP C ANTIBODY W/RFX RT PCR	
			0080 ††	URINE CULTURE	
			A861 ^{††}	TRICH (URINE, RNA)	
			6368 ^{††}	CT (URINE, RNA)	
			6369 ^{††}	GC (URINE RNA)	

	Test Code 7307 - Previous Test Information		Test Code	Test Code 7307- New Test Information		
Test Name	Obstetric/Prenatal I		Obstetric	ACOG Panel w/out HIV		
Test Components	0039	ANTIBODY SCREEN	0039	ANTIBODY SCREEN		
	0053	CBC WITH DIFF	0053	CBC WITH DIFF		
	0142	RPR	0142	RPR		
	0156	ABO/Rh BLOOD TYPE	0156	ABO/Rh BLOOD TYPE		
	0973	RUBELLA, IGG	0973	RUBELLA, IGG		
	0800	URINE CULTURE	0800	URINE CULTURE		
	0106	HEP B SURFACE ANTIGEN		HEP B SURF AG W/CONF, IF POS, RFX to DNA, QUANT,		
			TP85 [†]	PRENATAL		
			B125	HEP C ANTIBODY W/RFX RT PCR		

†Hepatitis B Surface Antigen (with confirmation), if positive, reflex to DNA, Quant for prenatal patients (Test code: TP85-7) will replace 0106- Hepatitis B Surface Antigen in the standard OB Panels to align with the <u>Centers for Disease Control and Prevention (CDC)</u> recommendation that all HBsAg-positive pregnant women should be tested for HBV DNA to guide the use of maternal antiviral therapy during pregnancy for the reduction of HBV transmission to the newborn.

Please refer to the test compendium or online test search for specimen requirements, collection instructions and CPT codes. If you have a favorites or profile currently created within your EMR, test components will need to be updated in November accordingly, in order to *avoid* test not performed (TNP's) due to duplicate orders. If you have any additional questions or need more information, please contact customer service or reach out to your dedicated sales representative.

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^{††}When collecting urine culture, please pour off urine into GenProbe Aptima urine tube for CT, GC, Trich

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