

June 2015 - Monthly Update, Quest Diagnostics Nichols Institute, Valencia

NEW TESTS			
Please Note: Not all test codes assigned to each assay are listed in the table of contents. Please refer to the complete listing on the page numbers indicated.			
Test Code	Test Name	Effective Date	Page #
<u>93020</u>	Cortisol, LC/MS/MS, Saliva, 2 Samples	5/18/2015	2
<u>90851</u>	Herpes Simplex Virus 1/2 Antibody (IgM), IFA with Reflex to Titer, CSF	7/20/2015	3
<u>37507</u>	Rickettsia Antibody Panel with Reflex to Titers	7/20/2015	3
<u>93022</u>	Zinc Transporter 8 (ZnT8) Antibody	7/20/2015	4
<u>17875</u>	Leptospira DNA, Qualitative Real-Time PCR	7/27/2015	5

TEST CHANGES				
Please Note: Not all test codes assigned to each assay are listed in the table of contents. Please refer to the complete listing on the page numbers indicated.				
Test Code	Former Test Code	Test Name	Effective Date	Page #
<u>16379</u>	S51768	Chromogranin A, Electrochemiluminescence	7/13/2015	5
S41505		Euglobulin Clot Lysis Time	7/13/2015	6
<u>496</u>	4972	Hemoglobin A1c	7/13/2015	6
		Hemoglobin A1c Changes	7/13/2015	7
<u>8181</u>	4975	Hemoglobin A1c with Calculated Mean Plasma Glucose (MPG)	7/13/2015	7
<u>16802</u>	4971	Hemoglobin A1c with eAG	7/13/2015	7
<u>1468</u>		Human Anti-mouse Antibody (HAMA)	7/13/2015	8
<u>37673</u>	9421	Rubella Antibodies (IgG, IgM) Diagnostic	7/13/2015	8
<u>4422</u>	2475	Rubella Antibody (IgM)	7/13/2015	9
<u>802</u>	9416	Rubella Immune Status	7/13/2015	9
<u>90924</u>		Hepatitis C Viral RNA NS3 Genotype	7/27/2015	10
<u>4128X</u>	2366C	VDRL, CSF	7/27/2015	10

DISCONTINUED TESTS			
Please Note: Not all test codes assigned to each assay are listed in the table of contents. Please refer to the complete listing on the page numbers indicated.			
Test Code	Test Name	Effective Date	Page #
4972SR	Hemoglobin A1c w/Serial Reporting	7/13/2015	11
S52315	Chlamydia Trachomatis/Psittaci Culture	7/20/2015	11
S50047	HSV 1/2 IgM Antibody IFA CSF	7/20/2015	11
S49977	Weil Felix Da	7/20/2015	11

NY UPDATE		
Please Note: Not all test codes assigned to each assay are listed in the table of contents. Please refer to the complete listing on the page numbers indicated.		
Test Code	Test Name	Page #
	New York Approval Received for HPV Testing	11

New Test Offerings

The following tests will be available through Quest Diagnostics on the dates indicated below.

Cortisol, LC/MS/MS, Saliva, 2 Samples																							
Effective Date	5/18/2015																						
Test Code	93020																						
CPT Codes	82530 (x2)																						
Specimen Requirements	0.5 mL (0.2 mL minimum) saliva collected in each of two separate Salivette® collection tubes																						
Reject Criteria	Hemolysis; any tubes other than Salivette®																						
Instructions	<p>Precautions:</p> <ol style="list-style-type: none"> 1. Saliva should be collected at the time(s) prescribed by your doctor. 2. No food or fluids for 30 minutes prior to collection. 3. Do not use any creams, lotions, or steroid inhalers immediately prior to collection. 4. Avoid any activity that can cause your gums to bleed, including brushing and flossing your teeth. Consult with your doctor if this is a chronic problem. 5. Do not use this kit on children under 3 years of age or any patient with increased risk of swallowing or choking. <p>Instructions for collection:</p> <ol style="list-style-type: none"> 1. Rinse mouth thoroughly with water and discard. Do not swallow. 2. Hold the Salivette® at the rim of the suspended insert and remove the stopper. 3. Remove the swab. 4. Place the swab under the tongue until well saturated, approximately 1 minute. 5. Return the saturated swab to the suspended insert and close the Salivette® firmly with the stopper. 6. Do not remove the tube holding the insert. The Salivette® should be sent to the lab with the swab inside. 7. Label the Salivette® with the patient name, date and time of collection, and any other identifying information. 																						
Transport Temperature	Refrigerated																						
Specimen Stability	Room temperature: 5 days Refrigerated: 7 days Frozen: 2 years																						
Set-up/Analytic Time	Set up: Sun-Thurs; Report available: 6-11 days																						
Reference Range	<table border="1"> <tbody> <tr> <td>8-10 AM:</td> <td>0.04-0.56 mcg/dL</td> </tr> <tr> <td>4-6 PM:</td> <td>< or = 0.15 mcg/dL</td> </tr> <tr> <td>10-11 PM:</td> <td>< or = 0.09 mcg/dL</td> </tr> </tbody> </table>		8-10 AM:	0.04-0.56 mcg/dL	4-6 PM:	< or = 0.15 mcg/dL	10-11 PM:	< or = 0.09 mcg/dL															
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4-6 PM:	< or = 0.15 mcg/dL																						
10-11 PM:	< or = 0.09 mcg/dL																						
Methodology	Liquid Chromatography/Tandem Mass Spectrometry																						
Performing Site	Quest Diagnostics Nichols Institute, San Juan Capistrano																						
Interface Mapping	<table border="1"> <thead> <tr> <th>Result Code</th> <th>Type</th> <th>Result Name</th> </tr> </thead> <tbody> <tr> <td>86008975</td> <td>Prompt-Result</td> <td>Draw Date 1</td> </tr> <tr> <td>86008976</td> <td>Prompt-Result</td> <td>Draw Time 1</td> </tr> <tr> <td>86006703</td> <td></td> <td>Cortisol,Salivary Sample 1</td> </tr> <tr> <td>86009038</td> <td>Prompt-Result</td> <td>Draw Date 2</td> </tr> <tr> <td>86009039</td> <td>Prompt-Result</td> <td>Draw Time 2</td> </tr> <tr> <td>86006704</td> <td></td> <td>Cortisol,Salivary Sample 2</td> </tr> </tbody> </table>		Result Code	Type	Result Name	86008975	Prompt-Result	Draw Date 1	86008976	Prompt-Result	Draw Time 1	86006703		Cortisol,Salivary Sample 1	86009038	Prompt-Result	Draw Date 2	86009039	Prompt-Result	Draw Time 2	86006704		Cortisol,Salivary Sample 2
Result Code	Type	Result Name																					
86008975	Prompt-Result	Draw Date 1																					
86008976	Prompt-Result	Draw Time 1																					
86006703		Cortisol,Salivary Sample 1																					
86009038	Prompt-Result	Draw Date 2																					
86009039	Prompt-Result	Draw Time 2																					
86006704		Cortisol,Salivary Sample 2																					

Herpes Simplex Virus 1/2 Antibody (IgM), IFA with Reflex to Titer, CSF																			
Clinical Significance	Detection of HSV IgM in CSF may aid in the diagnosis of encephalitis caused by HSV																		
Effective Date	7/20/2015																		
Test Code	90851																		
CPT Codes	86695, 86696																		
Specimen Requirements	1 mL (0.2 mL minimum) CSF collected in a sterile leak-proof container																		
Transport Temperature	Room temperature																		
Specimen Stability	Room temperature: 7 days Refrigerated: 14 days Frozen: 30 days																		
Set-up/Analytic Time	Set-up: Mon-Sat; Report available: 1-3 days																		
Reference Range	Negative																		
Methodology	Immunofluorescence Assay																		
Performing Site	Focus Diagnostics, Inc.																		
Interface Mapping	<table border="1"> <thead> <tr> <th>Result Code</th> <th>Result Name</th> </tr> </thead> <tbody> <tr> <td>86008220</td> <td>HSV 1 IgM Screen</td> </tr> <tr> <td>86008221</td> <td>HSV 2 IgM Screen</td> </tr> <tr> <td colspan="2"><i>This test is a true reflex. Please build the unit code below separately. Non-orderable Reflex: 90851-2-HSV 1 IgM Titer</i></td> </tr> <tr> <th>Result Code</th> <th>Result Name</th> </tr> <tr> <td>86008222</td> <td>HSV 1 IgM Titer</td> </tr> <tr> <td colspan="2"><i>This test is a true reflex. Please build the unit code below separately. Non-orderable Reflex: 90851-3-HSV 2 IgM Titer</i></td> </tr> <tr> <th>Result Code</th> <th>Result Name</th> </tr> <tr> <td>86008223</td> <td>HSV 2 IgM Titer</td> </tr> </tbody> </table>	Result Code	Result Name	86008220	HSV 1 IgM Screen	86008221	HSV 2 IgM Screen	<i>This test is a true reflex. Please build the unit code below separately. Non-orderable Reflex: 90851-2-HSV 1 IgM Titer</i>		Result Code	Result Name	86008222	HSV 1 IgM Titer	<i>This test is a true reflex. Please build the unit code below separately. Non-orderable Reflex: 90851-3-HSV 2 IgM Titer</i>		Result Code	Result Name	86008223	HSV 2 IgM Titer
Result Code	Result Name																		
86008220	HSV 1 IgM Screen																		
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Result Code	Result Name																		
86008223	HSV 2 IgM Titer																		
Additional Information	If HSV 1 IgM Screen is Positive, HSV 1 IgM Titer will be performed at an additional charge (CPT code: 86695). If HSV 2 IgM Screen is Positive, HSV 2 IgM Titer will be performed at an additional charge (CPT code: 86696).																		

Rickettsia Antibody Panel with Reflex to Titers	
Clinical Significance	Antigen-specific IgG and IgM titers allow rapid diagnosis of infection by organisms within either of the two major groups of Rickettsia (Rocky Mountain Spotted Fever) and R.akari (Rickettsial pox); the typhus group includes R.typhi (endemic/maurine typhus) and R.prowazeki (epidemic typhus).
Effective Date	7/20/2015
Test Code	37507
CPT Codes	86757 (x4)
Specimen Requirements	1 mL (0.2 mL minimum) serum
Reject Criteria	Gross hemolysis; gross lipemia
Transport Temperature	Room temperature

Specimen Stability	Room temperature: 7 days Refrigerated: 14 days Frozen: 30 days																																		
Set-up/Analytic Time	Set-up: Mon-Fri; Report available: 1-4 days																																		
Reference Range	Not Detected																																		
Methodology	Immunofluorescence Assay																																		
Performing Site	Focus Diagnostics, Inc.																																		
Interface Mapping	<table border="1"> <thead> <tr> <th>Result Code</th> <th>Result Name</th> </tr> </thead> <tbody> <tr> <td>70025900</td> <td>RMSF IgG</td> </tr> <tr> <td>70035700</td> <td>RMSF IgM</td> </tr> <tr> <td>70026000</td> <td>R. typhi IgG</td> </tr> <tr> <td>70026040</td> <td>R. typhi IgM</td> </tr> <tr> <td colspan="2"><i>This test is a true reflex. Please build the unit code below separately. Non-orderable Reflex: 6419-2-Rickettsia (RMSF) Antibody Titer, IgG</i></td> </tr> <tr> <th>Result Code</th> <th>Result Name</th> </tr> <tr> <td>70025906</td> <td>RMSF IgG Titer</td> </tr> <tr> <td colspan="2"><i>This test is a true reflex. Please build the unit code below separately. Non-orderable Reflex: 6419-3-Rickettsia (RMSF) Antibody Titer, IgM</i></td> </tr> <tr> <th>Result Code</th> <th>Result Name</th> </tr> <tr> <td>70035706</td> <td>RMSF IgM Titer</td> </tr> <tr> <td colspan="2"><i>This test is a true reflex. Please build the unit code below separately. Non-orderable Reflex: 37503-2-Rickettsia (Typhus Fever) Antibody Titer, IgG</i></td> </tr> <tr> <th>Result Code</th> <th>Result Name</th> </tr> <tr> <td>70026021</td> <td>R. typhi IgG Titer</td> </tr> <tr> <td colspan="2"><i>This test is a true reflex. Please build the unit code below separately. Non-orderable Reflex: 37503-3-Rickettsia (Typhus Fever) Antibody Titer, IgM</i></td> </tr> <tr> <th>Result Code</th> <th>Result Name</th> </tr> <tr> <td>70026071</td> <td>R. typhi IgM Titer</td> </tr> </tbody> </table>	Result Code	Result Name	70025900	RMSF IgG	70035700	RMSF IgM	70026000	R. typhi IgG	70026040	R. typhi IgM	<i>This test is a true reflex. Please build the unit code below separately. Non-orderable Reflex: 6419-2-Rickettsia (RMSF) Antibody Titer, IgG</i>		Result Code	Result Name	70025906	RMSF IgG Titer	<i>This test is a true reflex. Please build the unit code below separately. Non-orderable Reflex: 6419-3-Rickettsia (RMSF) Antibody Titer, IgM</i>		Result Code	Result Name	70035706	RMSF IgM Titer	<i>This test is a true reflex. Please build the unit code below separately. Non-orderable Reflex: 37503-2-Rickettsia (Typhus Fever) Antibody Titer, IgG</i>		Result Code	Result Name	70026021	R. typhi IgG Titer	<i>This test is a true reflex. Please build the unit code below separately. Non-orderable Reflex: 37503-3-Rickettsia (Typhus Fever) Antibody Titer, IgM</i>		Result Code	Result Name	70026071	R. typhi IgM Titer
Result Code	Result Name																																		
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70035700	RMSF IgM																																		
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Result Code	Result Name																																		
70026071	R. typhi IgM Titer																																		
Additional Information	If Rickettsia (RMSF) IgG or IgM screen is "Detected", then the appropriate IgG or IgM Titer will be performed at an additional charge (CPT code(s): 86757) . If Rickettsia (Typhus Fever) IgG or IgM screen is "Detected", then the appropriate IgG or IgM Titer will be performed at an additional charge (CPT code(s): 86757).																																		

Zinc Transporter 8 (ZnT8) Antibody	
Clinical Significance	Patient population is primarily children and adolescents to confirm a diagnosis of type 1 diabetes.
Effective Date	7/20/2015
Test Code	93022
CPT Codes	86341
Specimen Requirements	1 mL (0.5 mL minimum) serum
Reject Criteria	Gross hemolysis; lipemia; icterus; specimens other than serum
Transport Temperature	Room temperature
Specimen Stability	Room temperature and Refrigerated: 7 days

	Frozen: 28 days					
Set-up/Analytic Time	Set up: Wed, Sat; Report available: 2-3 days					
Reference Range	<15 U/mL					
Methodology	Enzyme Linked Immunosorbent Immunoassay					
Performing Site	Quest Diagnostics Nichols Institute, San Juan Capistrano					
Interface Mapping	<table border="1"> <thead> <tr> <th>Result Code</th> <th>Result Name</th> </tr> </thead> <tbody> <tr> <td>86013368</td> <td>ZnT8 Ab</td> </tr> </tbody> </table>		Result Code	Result Name	86013368	ZnT8 Ab
Result Code	Result Name					
86013368	ZnT8 Ab					

Leptospira DNA, Qualitative Real-Time PCR											
Clinical Significance	PCR can be used to rapidly diagnose Leptospirosis, an infection caused by a waterborne spirochete of the genus Leptospira. In addition, testing of blood, CSF and urine may give an indication of the stage of infection.										
Effective Date	7/27/2015										
Test Code	17875										
CPT Codes	87798										
Specimen Requirements	5 mL (1 mL minimum) frozen urine collected in a sterile, plastic, leak-proof container										
Transport Temperature	Frozen										
Specimen Stability	Room temperature and Refrigerated: Unacceptable Frozen: 30 days										
Set-up/Analytic Time	Set up: Mon, Wed, Fri; Report available: 1-3 days										
Reference Range	Not detected										
Methodology	Real-Time Polymerase Chain Reaction										
Performing Site	Focus Diagnostics, Inc.										
Interface Mapping	<table border="1"> <thead> <tr> <th>Result Code</th> <th>Type</th> <th>Result Name</th> </tr> </thead> <tbody> <tr> <td>86007404</td> <td>Prompt-Result</td> <td>Specimen Source:</td> </tr> <tr> <td>86012010</td> <td></td> <td>Leptospira DNA, QL</td> </tr> </tbody> </table>		Result Code	Type	Result Name	86007404	Prompt-Result	Specimen Source:	86012010		Leptospira DNA, QL
Result Code	Type	Result Name									
86007404	Prompt-Result	Specimen Source:									
86012010		Leptospira DNA, QL									

Test Changes

The following test changes will be effective on the dates indicated below. **Please note information that is changing appears in bold text in this update.** *Former test names and test codes have been italicized.*

Chromogranin A, Electrochemiluminescence	
Clinical Significance	Chromogranin A (CgA) is the major protein within the catecholamine storage vesicles of the adrenal medulla. CgA has been shown to be co-released with polypeptide hormones from the following endocrine tissues: pancreatic islet cells, enteroendocrine cells, parathyroid chief cells, thyroid parafollicular C cells, and anterior pituitary cells. CgA concentrations may be elevated in patients with various endocrine neoplasms, e.g., anterior pituitary adenoma, parathyroid adenoma, medullary thyroid carcinoma, carcinoid tumor, and pancreatic islet cell tumor.
Effective Date	7/13/2015
<i>Former Test Name</i>	<i>Chromogranin A</i>

Former Test Code	S51768					
Test Code	16379					
Reject Criteria	Gross hemolysis; grossly lipemic; grossly icteric					
Instructions	Collect blood in red-top or serum separator tube. Centrifuge and separate the serum from the cells.					
Transport Temperature	Room temperature					
Set-up/Analytic Time	Set up: Sun-Fri; Report available: 3-5 days					
Reference Range	< or = 15 ng/mL					
Methodology	Electrochemiluminescence					
Performing Site	Quest Diagnostics Nichols Institute, San Juan Capistrano					
Interface Mapping	<table border="1"> <thead> <tr> <th>Result Code</th> <th>Result Name</th> </tr> </thead> <tbody> <tr> <td>86006273</td> <td>Chromogranin A, ECL</td> </tr> </tbody> </table>	Result Code	Result Name	86006273	Chromogranin A, ECL	
Result Code	Result Name					
86006273	Chromogranin A, ECL					
Pricing Message	Negotiated pricing on S51768 will be applied to code 16379.					

Euglobulin Clot Lysis Time	
Clinical Significance	ECLT provides an overall assessment of the fibrinolysis system by measuring the time for an in vitro clot to dissolve in the absence of the normal plasmin inhibitors. ECLT is useful in assessing the fibrinolytic system and monitoring patients on urokinase or streptokinase fibrinolytic therapy.
Effective Date	7/13/2015
Former Test Name	Euglobulin Lysis Time
Test Code	S41505
Specimen Requirements	2 mL (1 mL minimum) frozen plasma collected in 3.2% sodium citrate (light blue-top) tube
Instructions	Specimen collection container is blue-top (3.2% sodium citrate) tube. To avoid release of plasminogen activator, do not massage vein vigorously, pump first excessively or leave tourniquet in place for a prolonged period. Centrifuge within 30 minutes after collection to get platelet-poor plasma and freeze on dry ice. Ship specimens frozen on dry ice. Keep samples in a -60 to -80 degree C freezer if they cannot be shipped promptly. Prohibit exercise prior to drawing sample. Jacobs D., et al, "Laboratory Test Handbook", 4th Edition, Lexi-Comp Inc., Hudson (Cleveland), 1996, p238.
Specimen Stability	Room temperature: Unacceptable Refrigerated: 4 hours Frozen: 21 days
Units Of Measure	min
Methodology	Clot Dissolution
Performing Site	Quest Diagnostics Nichols Institute, San Juan Capistrano

Hemoglobin A1c	
Clinical Significance	Assesses long term diabetic control in diabetes mellitus.
Effective Date	7/13/2015
Former Test Code	4972
Test Code	496
Specimen Requirements	1 mL (0.5 mL minimum) whole blood collected in an EDTA (lavender-top) tube Lithium heparin (green-top) tubes are no longer acceptable

Reject Criteria	Sodium fluoride/oxalate (grey-top) tube; heparinized plasma					
Transport Temperature	Room temperature					
Reference Range	<5.7 % of total Hgb					
Performing Site	Quest Diagnostics Nichols Institute, Valencia					
Interface Mapping	<table border="1"> <thead> <tr> <th>Result Code</th> <th>Result Name</th> </tr> </thead> <tbody> <tr> <td>50026400</td> <td>Hemoglobin A1c</td> </tr> </tbody> </table>		Result Code	Result Name	50026400	Hemoglobin A1c
Result Code	Result Name					
50026400	Hemoglobin A1c					

Hemoglobin A1c Changes										
Effective Date	7/13/2015									
Reject Criteria	Sodium fluoride/oxalate (grey-top) tube; heparinized plasma									
Reference Range	<5.7 % of total Hgb									
Performing Site	Quest Diagnostics Nichols Institute, San Juan Capistrano									
Tests Affected	<table border="1"> <thead> <tr> <th>Test Codes:</th> <th>Name:</th> </tr> </thead> <tbody> <tr> <td>92062</td> <td>Diabetes and ASCVD Risk Panel with Scores</td> </tr> <tr> <td>92027</td> <td>Diabetes Risk Panel with Score</td> </tr> <tr> <td>91920</td> <td>Diabetes Risk Panel without Score</td> </tr> </tbody> </table>		Test Codes:	Name:	92062	Diabetes and ASCVD Risk Panel with Scores	92027	Diabetes Risk Panel with Score	91920	Diabetes Risk Panel without Score
Test Codes:	Name:									
92062	Diabetes and ASCVD Risk Panel with Scores									
92027	Diabetes Risk Panel with Score									
91920	Diabetes Risk Panel without Score									

Hemoglobin A1c with Calculated Mean Plasma Glucose (MPG)							
Clinical Significance	Assesses long term diabetic control in diabetes mellitus.						
Effective Date	7/13/2015						
Former Test Name	<i>Hemoglobin A1c w/MPG</i>						
Former Test Code	4975						
Test Code	8181						
Specimen Requirements	1 mL (0.5 mL minimum) whole blood collected in an EDTA (lavender-top) tube Lithium heparin (green-top) tubes are no longer acceptable						
Reject Criteria	Sodium fluoride/oxalate (grey-top) tube; heparinized plasma						
Transport Temperature	Room temperature						
Reference Range	Hemoglobin A1c: <5.7 % of total Hgb						
Performing Site	Quest Diagnostics Nichols Institute, Valencia						
Interface Mapping	<table border="1"> <thead> <tr> <th>Result Code</th> <th>Result Name</th> </tr> </thead> <tbody> <tr> <td>50026400</td> <td>Hemoglobin A1c</td> </tr> <tr> <td>50067805</td> <td>Mean Plasma Glucose</td> </tr> </tbody> </table>	Result Code	Result Name	50026400	Hemoglobin A1c	50067805	Mean Plasma Glucose
Result Code	Result Name						
50026400	Hemoglobin A1c						
50067805	Mean Plasma Glucose						

Hemoglobin A1c with eAG	
Clinical Significance	Assesses long term diabetic control in diabetes mellitus.

Effective Date	7/13/2015									
<i>Former Test Code</i>	4971									
Test Code	16802									
Specimen Requirements	1 mL (0.5 mL minimum) whole blood collected in an EDTA (lavender-top) tube Lithium heparin (green-top) tubes are no longer acceptable									
Reject Criteria	Sodium fluoride/oxalate (grey-top) tube; heparinized plasma									
Transport Temperature	Room temperature									
Reference Range	Hemoglobin A1c: <5.7 % of total Hgb									
Performing Site	Quest Diagnostics Nichols Institute, Valencia									
Interface Mapping	<table border="1"> <thead> <tr> <th>Result Code</th> <th>Result Name</th> </tr> </thead> <tbody> <tr> <td>50026400</td> <td>Hemoglobin A1c</td> </tr> <tr> <td>50067820</td> <td>eAG (mg/dL)</td> </tr> <tr> <td>50067830</td> <td>eAG (mmol/L)</td> </tr> </tbody> </table>		Result Code	Result Name	50026400	Hemoglobin A1c	50067820	eAG (mg/dL)	50067830	eAG (mmol/L)
Result Code	Result Name									
50026400	Hemoglobin A1c									
50067820	eAG (mg/dL)									
50067830	eAG (mmol/L)									

Human Anti-mouse Antibody (HAMA)	
Effective Date	7/13/2015
Test Code	1468
Reference Range	< or = 74 ng/mL
Performing Site	Quest Diagnostics Nichols Institute, San Juan Capistrano

Rubella Antibodies (IgG, IgM) Diagnostic					
Clinical Significance	<p>Rubella (German Measles) is a common and usually benign viral infection of children. IgM Antibody is detectable 11-25 days after the onset of exanthem, in most individuals 15-25 days after vaccination, and in 90-97% of infants with congenital rubella between 2 weeks and 3 months after birth. Acute and convalescent IgG titers may be interpreted to identify a >30% rise in the antibody index. A positive IgG antibody indicates successful immunization or past exposure.</p>				
Effective Date	7/13/2015				
<i>Former Test Name</i>	<i>Rubella Antibodies (IgG, IgM)</i>				
<i>Former Test Code</i>	9421				
Test Code	37673				
Reference Range	<table border="1"> <tbody> <tr> <td>Rubella Antibody (IgG), Dx</td> <td> < or = 0.90 Negative 0.91-1.09 Equivocal > or = 1.10 Positive </td> </tr> <tr> <td>Rubella Antibody (IgM)</td> <td> <0.90 Negative 0.90-1.09 Equivocal > or = 1.10 Positive </td> </tr> </tbody> </table>	Rubella Antibody (IgG), Dx	< or = 0.90 Negative 0.91-1.09 Equivocal > or = 1.10 Positive	Rubella Antibody (IgM)	<0.90 Negative 0.90-1.09 Equivocal > or = 1.10 Positive
Rubella Antibody (IgG), Dx	< or = 0.90 Negative 0.91-1.09 Equivocal > or = 1.10 Positive				
Rubella Antibody (IgM)	<0.90 Negative 0.90-1.09 Equivocal > or = 1.10 Positive				
Units Of Measure	Remove IU/mL				
Methodology	Immunoassay				
Performing Site	Quest Diagnostics Nichols Institute, Valencia				

Interface Mapping	Result Code	Result Name
	86010949	Rubella Antibody (IgG), Dx
	70026600	Rubella Antibody (IgM)
Pricing Message	Negotiated pricing on 9421 will be applied to code 37673.	

Rubella Antibody (IgM)		
Effective Date	7/13/2015	
<i>Former Test Code</i>	2475	
Test Code	4422	
Transport Temperature	Room temperature	
Performing Site	Quest Diagnostics Nichols Institute, Valencia	
Interface Mapping	Result Code	Result Name
	70026600	Rubella Antibody (IgM)
Pricing Message	Negotiated pricing on 2475 will be applied to code 4422.	
Tests Affected	Test Codes:	Name:
	9901	Torch IgG and IgM Antibodies Evaluation
	2231	Torch IgM Antibodies Evaluation

Rubella Immune Status		
Clinical Significance	Rubella is an acute exanthematous viral infection of children and adults. Rash, fever and lymphadenopathy characterize the illness. While many infections are subclinical, this virus has the potential to cause fetal infection with resultant birth defects. Diagnosis of a Rubella infection is best made serologically. In the absence of a current or recent infection, a demonstration of specific IgG on a serum sample is evidence of immunity to Rubella.	
Effective Date	7/13/2015	
<i>Former Test Name</i>	Rubella IgG Antibodies	
<i>Former Test Code</i>	9416	
Test Code	802	
Specimen Requirements	1 mL (0.1 mL minimum) serum	
Reject Criteria	Gross hemolysis; grossly lipemic, plasma	
Transport Temperature	Room temperature	
Specimen Stability	Room temperature: 4 days Refrigerated: 7 days Frozen: 30 days	
Reference Range	< or = 0.90	Not consistent with immunity
	0.91-1.09	Equivocal
	> or = 1.10	Consistent with immunity

Units Of Measure	Remove IU/mL									
Methodology	Immunoassay									
Performing Site	Quest Diagnostics Nichols Institute, Valencia									
Interface Mapping	<table border="1"> <thead> <tr> <th>Result Code</th> <th>Result Name</th> </tr> </thead> <tbody> <tr> <td>70012000</td> <td>Rubella Antibody (IgG)</td> </tr> </tbody> </table>	Result Code	Result Name	70012000	Rubella Antibody (IgG)					
Result Code	Result Name									
70012000	Rubella Antibody (IgG)									
Pricing Message	Negotiated pricing on 9416 will be applied to code 802.									
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Test Codes:	Name:									
1341	Immune Status Panel									
9901	Torch IgG and IgM Antibodies Evaluation									
9911	Torch IgG Antibodies Evaluation									

Hepatitis C Viral RNA NS3 Genotype	
Effective Date	7/27/2015
Test Code	90924
Specimen Requirements	Preferred: 2 mL (0.6 mL minimum) plasma collected in an EDTA (lavender-top) tube Acceptable: Plasma collected in a PPT potassium EDTA (white-top) tube
Transport Temperature	Refrigerated
Specimen Stability	Room temperature: 72 hours Refrigerated: 14 days Frozen: 42 days
Set-up/Analytic Time	Set up: Mon; Report available: 4-11 days
Performing Site	Focus Diagnostics, Inc.

VDRL, CSF	
Clinical Significance	Positive results are suggestive of neurosyphilis.
Effective Date	7/27/2015
Former Test Code	2366C
Test Code	4128X
Specimen Requirements	1 mL (0.2 mL minimum) CSF collected in a plastic, screw-cap vial
Reject Criteria	Hemolysis; lipemia
Transport Temperature	Refrigerated
Specimen Stability	Room temperature: 4 days Refrigerated: 14 days Frozen: 6 months
Methodology	Slide Micro-flocculation
Performing Site	Quest Diagnostics Nichols Institute, Valencia

Interface Mapping	Result Code	Result Name
	40010700	VDRL, CSF
Pricing Message	Negotiated pricing on 2366C will be applied to code 4128.	

Discontinued Tests

Hemoglobin A1c w/Serial Reporting	
Effective Date	7/13/2015
Test Code	4972SR
Additional Information	The recommended alternative is test code 496-Hemoglobin A1c

Chlamydia Trachomatis/Psittaci Culture	
Effective Date	7/20/2015
Test Code	S52315
Additional Information	The recommended alternative is test code 2400-Chlamydia Trachomatis Culture

HSV 1/2 IgM Antibody IFA CSF	
Effective Date	7/20/2015
Test Code	S50047
Additional Information	The recommended alternative is test code 90851-Herpes Simplex Virus 1/2 Antibody (IgM), IFA with Reflex to Titer, CSF

Weil Felix Da	
Effective Date	7/20/2015
Test Code	S49977
Additional Information	The recommended alternative is test code 37507- Rickettsia Antibody Panel with Reflex to Titers

New York Patient Testing Update

New York Approval Received for HPV Testing													
Message	**The following test codes are now available for New York patient testing.**												
Performing Site	Focus Diagnostics, Inc.												
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	Test Codes:	Name:											
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92211	HPV mRNA E6/E7 SurePath(TM) with Reflex												

91932	HPV mRNA E6/E7, Rectal
19863	HPV, High Risk, with Reflex to Genotypes 16 and 18