

Laboratory Update

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The CPT Codes provided in this document are based on AMA guidelines and are for informational purposes only. CPT coding is the sole responsibility of the billing party. Please direct any questions regarding coding to the payor being billed. Any Profile/panel component may be ordered separately. Reflex tests are performed at an additional charge.

Summary of Test Changes

Page Number	Test Name	Test Code(s)	Re-direct(s) Performing Site	Test Code	Test Name	Specimen Requirements	Transport Temperature	Specimen Stability	Units of Measure	Reference Range	Methodology	CPT Codes	Reject Criteria	Other (see listing)
11	ColoVantage™ (methylated Septin 9)	16983				X								X
11	Sex Hormone Binding Globulin	30740								X				
11	GlycoMark®	19599											X	
12	Beta-2 Transferrin	10640												X
12	IGF-I, LC/MS	16293			X	X		X		X	X		X	X
13	Influenza A H1N1 (2009) Real-Time RT-PCR	16807				X								
13	Influenza Type A/B RT-PCR Reflex to Influenza A H1N1 (2009) RT-PCR	16861				X								
14	<i>Clostr. difficile</i> Cytotoxin Antibody, Neutralization	34403				X	X	X		X	X		X	
14	Herpes Simplex Virus Type 1 & 2 DNA, Quantitative Real-Time PCR	19502				X								
14	Porphobilinogen, Quantitative, 24-Hour Urine	726								X				
14	Porphobilinogen, Quantitative, Random Urine	6329								X				

Announcements

April 25, 2011

Dear Valued Client:

We are committed to providing the diagnostic insights and innovations that can help you improve the health of your patients.

In an effort to better meet your needs, we will be implementing more sensitive cutoff values for Opiates and Oxycodone for our **Pain Management Urine Drug Test** offering.

Effective April 25, 2011, we will update cutoff values as follows:

Pain Management - Screening Drug Tests (Urine)	Previous Cut Off Value (ng/mL)	New Cut Off Value (ng/mL)
Pain Management Opiate Screens (Urine) <ul style="list-style-type: none">• Pain Management <u>Profiles</u>: Base, Profile 1, 2, 3 and 4• Pain Management <u>Screen with Confirmations</u>: Opiates, Oxycodone• Pain Management <u>Screen Only</u>: Opiates, Oxycodone	300	100

Pain Management - Quantitative Drug Test (Urine)	Previous Cut Off Value (ng/mL)	New Cut Off Value (ng/mL)
Pain Management Opiates, Quantitative (Urine) <ul style="list-style-type: none">• Pain Management Opiates, Quantitative• Pain Management Opiates Expanded, Quantitative	100	50
Pain Management Oxycodone Quantitative (Urine) <ul style="list-style-type: none">• Pain Management Oxycodone, Quantitative	100	50

We thank you for your business and look forward to continuing to serve all of your laboratory needs.

If you have any questions regarding these changes, please feel free to contact us: 1-877-40 RX TOX (844-407-9869).

Sincerely,



Ron G. Schlesinger, M.D.
Medical Director, Diagnostic Laboratory of Oklahoma

Re-activations:

Hepatitis C Supplemental RIBA confirmation will again be available for testing effective April 28, 2011. Your patience and understanding during this nationwide vendor reagent outage has been appreciated. The following codes are impacted:

Hepatitis C Antibody w/Reflex HCV RIBA	
Effective Date:	April 28, 2011
Test Code:	37677

Hepatitis C Antibody Supplemental Testing (RIBA)	
Effective Date:	April 28, 2011
Test Code:	8739

New Test Offerings

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

Chromium, Blood	
Clinical Significance:	The assay is useful to: 1. Monitor exposure to chromium 2. Monitor progress of medical treatment 3. Determine nutritional status.
Effective Date:	March 28, 2011
Test Code:	6085
Specimen Requirements:	4 mL (EDTA) royal blue-top whole blood (2 mL minimum) To avoid contamination, use powderless gloves. DO NOT ALIQUOT SPECIMEN. Draw one Vacutainer of blood (1-2 mL) and discard. Draw second Vacutainer (2-4 mL in royal blue top, EDTA) for submission. See Specimen Collection Section, Toxicology, in Directory of Services, Quest Diagnostics Nichols Institute. Patient should refrain from taking mineral supplements, and multi-vitamin three days prior to specimen collection.
Rejection Criteria:	Moderate hemolysis; gross lipemia; clotted specimen
Transport Temperature:	Refrigerated
Specimen Stability:	Room temperature: 48 hours Refrigerated: 5 days Frozen: Unacceptable
Reference Ranges:	< or = 1.2 mcg/L
Methodology:	Inductively Coupled Plasma – Mass Spectrometry with Dynamic Reaction Cell. (DRC-ICP-MS)
Performing Site:	Quest Diagnostics Nichols Institute, San Juan Capistrano.

AccuType® IL28B	
Clinical Significance:	This AccuType genomic assay detects the rs12979860C/T variant upstream of the IL28B gene and will be used to help predict therapeutic response to interferon and ribavirin in patients infected with hepatitis C (HCV) genotype 1. Individuals with the mutation have been shown to exhibit a greater response to therapy than individuals without it (Nature 2009;461:399-401). The IL28B assay will detect the C polymorphism that affects response to interferon and ribavirin, which are associated with significant expense and side effects in the treatment of hepatitis C. The 48 week course of interferon and ribavirin has limited efficacy (in HCV genotype 1) and is often poorly tolerated due to side effects that prevent patients from finishing treatment. The discovery of the IL28B gene and its association with HCV therapy response is a significant breakthrough in understanding how genetic variants may influence outcomes. The IL28B assay can assist the physician in selecting the most appropriate therapy for each individual patient.
Effective Date:	April 4, 2011
Test Code:	90251
CPT Code(s):	83891, 83898, 83896 (x2), 83912
Specimen Requirements:	5 mL EDTA (lavender-top) whole blood
Transport Temperature:	Whole blood is shipped ambient (18-26 C). Do not freeze whole blood.
Specimen Stability:	Room temperature and Refrigerated: 8 days Frozen: Do Not Freeze
Reference Ranges:	Accompanies report
Methodology:	Real-Time Polymerase Chain Reaction
Performing Site:	Quest Diagnostics Nichols Institute, San Juan Capistrano

AccuType® Metformin									
Clinical Significance:	The SNP rs2289669 G>A in the SLC47A1 gene, coding for the MATE1 protein, is associated with glucose lowering effect of metformin in patients with diabetes. The A allele is associated with higher levels of reduction in HbA1c. Genotyping patients with respect to these polymorphisms will determine the therapeutic response to metformin.								
Effective Date:	May 2, 2011								
Test Code:	18979								
CPT Code(s):	83891, 83892, 83898, 83909, 83912, 83914								
Specimen Requirements:	5 mL EDTA (lavender-top) whole blood Normal phlebotomy procedure. EDTA is the preferred anticoagulant, but ACD is also acceptable. 2 mL saliva (min 1 mL) use ORAGENE DNA™ Self Collection Kit OG-500 or OG-510								
Rejection Criteria:	Blood: Frozen samples								
Transport Temperature:	Room temperature								
Specimen Stability:	<table border="0"> <tr> <td>Blood:</td> <td>Saliva:</td> </tr> <tr> <td>Room temperature: 7 days</td> <td>Room temperature: 8 days</td> </tr> <tr> <td>Refrigerated: 7 days</td> <td>Refrigerated: 8 days</td> </tr> <tr> <td>Frozen: Unacceptable</td> <td>Frozen: 8 days</td> </tr> </table>	Blood:	Saliva:	Room temperature: 7 days	Room temperature: 8 days	Refrigerated: 7 days	Refrigerated: 8 days	Frozen: Unacceptable	Frozen: 8 days
Blood:	Saliva:								
Room temperature: 7 days	Room temperature: 8 days								
Refrigerated: 7 days	Refrigerated: 8 days								
Frozen: Unacceptable	Frozen: 8 days								
Reference Ranges:	Accompanies report								
Methodology:	Real Time Polymerase Chain Reaction								
Performing Site:	Quest Diagnostics Nichols Institute, Valencia								

Lyme Disease (<i>Borrelia</i> spp) DNA Qualitative Real-Time PCR, Blood	
Clinical Significance:	The diagnosis of Lyme disease is most often made by clinical examination combined with evidence of tick bite or exposure in endemic areas. Amplification of <i>Borrelia</i> genomic DNA from blood, fluids or tissues can support the diagnosis.
Effective Date:	May 2, 2011
Test Code:	15777
CPT Code(s):	87801
Specimen Requirements:	1 mL whole blood (EDTA or ACD)
Transport Temperature:	Refrigerated
Specimen Stability:	Room temperature: 48 hours Refrigerated: 7 days Frozen: Unacceptable
Reference Ranges:	Not Detected
Methodology:	Real-Time Polymerase Chain Reaction
Performing Site:	Focus Diagnostics, Inc.

Lyme Disease (<i>Borrelia</i> spp) DNA Qualitative Real-Time PCR, Synovial Fluid/CSF	
Clinical Significance:	The diagnosis of Lyme disease is most often made by clinical examination combined with evidence of tick bite or exposure in endemic areas. Amplification of <i>Borrelia</i> genomic DNA from blood, fluids or tissues can support the diagnosis.
Effective Date:	May 2, 2011
Test Code:	15564
CPT Code(s):	87801
Specimen Requirements:	1 mL synovial fluid or CSF
Transport Temperature:	Refrigerated
Specimen Stability:	Room temperature: 48 hours Refrigerated: 7 days Frozen: 30 days
Reference Ranges:	Not Detected
Methodology:	Real-Time Polymerase Chain Reaction
Performing Site:	Focus Diagnostics, Inc.

Lyme Disease (<i>Borrelia</i> spp) DNA Qualitative Real-Time PCR, Tick	
Clinical Significance:	The diagnosis of Lyme disease is most often made by clinical examination combined with evidence of tick bite or exposure in endemic areas. Amplification of <i>Borrelia</i> genomic DNA from blood, fluids, or tissues can support the diagnosis.
Effective Date:	May 2, 2011
Test Code:	15510
CPT Code(s):	87801
Specimen Requirements:	1 deer tick in 70% ethanol or in wet tissue in a sterile screw cap container
Transport Temperature:	Room temperature
Specimen Stability:	Room temperature, Refrigerated and Frozen: 14 days
Reference Ranges:	Not Detected
Methodology:	Real-Time Polymerase Chain Reaction
Performing Site:	Focus Diagnostics, Inc.

Lyme Disease (<i>Borrelia</i> spp) DNA Qualitative Real-Time PCR, Urine	
Clinical Significance:	The diagnosis of Lyme disease is most often made by clinical examination combined with evidence of tick bite or exposure in endemic areas. Amplification of <i>Borrelia</i> genomic DNA from blood, fluids or tissues can support the diagnosis.
Effective Date:	May 2, 2011
Test Code:	15868
CPT Code(s):	87801
Specimen Requirements:	4 mL random urine
Transport Temperature:	Frozen
Specimen Stability:	Room temperature: 48 hours Refrigerated: 7 days Frozen: 30 days
Reference Ranges:	Not Detected
Methodology:	Real-Time Polymerase Chain Reaction
Performing Site:	Focus Diagnostics, Inc.

FISH, Melanoma, Deletion 9p21 (CDKN2A/P16)	
Clinical Significance:	The deletion of 9p21 (CDKN2A/p16) is present in both early and late stages of the primary melanoma. Studies suggest that 9p21 deletion might be a highly informative marker in common and dysplastic nevi with high risk of malignant transformation (Casorzo, et al. Melanoma Res. 2005;15:155-160)
Effective Date:	May 2, 2011
Test Code:	16863
CPT Code(s):	88271(x2), 88275, 88291
Specimen Requirements:	Formalin-fixed, paraffin-embedded tissue Properly fixed and paraffin-embedded human tissue blocks will keep indefinitely if stored in a cool place. Slides with paraffin-embedded tissue sections can be kept 1-3 years if stored at 2-8C.
Transport Temperature:	Room temperature
Specimen Stability:	See Instructions
Reference Ranges:	FISH, Melanoma, del 9p21: Accompanies report
Methodology:	Fluorescence in situ Hybridization
Performing Site:	Quest Diagnostics Nichols Institute, San Juan Capistrano

Aspergillus DNA, Qualitative Real-Time PCR			
<i>This test includes: Aspergillus spp, A.fumigatus and A. terreus</i>			
Clinical Significance:	Detection of <i>Aspergillus</i> species DNA in clinical specimens can be useful in the diagnosis of invasive aspergillosis. Identification of the particular <i>Aspergillus</i> species can be useful in determining treatment.		
Effective Date:	May 9, 2011		
Test Code:	18873		
CPT Code(s):	87798 (x3)		
Specimen Requirements:	3 mL bronchoalveolar lavage, sputum, or whole blood (EDTA, ACD) or 3mm tissue		
Transport Temperature:	Frozen: Tissue Refrigerated: All others		
Specimen Stability:	Bronchoalveolar lavage or sputum	Whole blood	Tissue
	Room temperature: 48 hours	48 hours	Unacceptable
	Refrigerated: 7 days	7 days	7 days
	Frozen: 30 days	Do not Freeze	30 days
Reference Ranges:	Not Detected		
Methodology:	Real-Time Polymerase Chain Reaction		
Performing Site:	Focus Diagnostics, Inc.		

Dabigatran	
Clinical Significance:	Dabigatran is a new oral direct anti thrombin drug and an alternative to Warfarin. The drug received FDA clearance in 2010 for the indication of stroke prevention in patients with atrial fibrillation. Routinely, the drug does not need to be monitored, however, there are instances where monitoring is needed: (1) Determination of failure of therapy vs. poor compliance (2) Potential dose adjustment required for renal or hepatic dysfunction (3) Titration.
Effective Date:	June 13, 2011
Test Code:	15972
CPT Code(s):	80299
Specimen Requirements:	1 mL sodium citrate (light blue-top tube) plasma Whole blood collected in 3.2% citrate (light blue top tube). Centrifuge light blue-top tube 15 minutes at approx. 1500 g within 60 minutes of collection. Using a plastic pipette, remove plasma, taking care to avoid the wbc/platelet buffy layer and place into a plastic vial. Centrifuge a second time and transfer platelet-poor plasma into a new plastic vial. Plasma must be free of platelets (<10,000/mcl). Freeze immediately and ship on dry ice.
Rejection Criteria:	Gross hemolysis, visible fibrin clot
Transport Temperature:	Frozen
Specimen Stability:	Room temperature and Refrigerated: Unacceptable Frozen: 30 days
Reference Ranges:	64 - 443 ng/mL
Methodology:	Clotting Time
Performing Site:	Quest Diagnostics Nichols Institute, Chantilly

Modified Hodge Test	
Clinical Significance:	<i>Klebsiella pneumoniae</i> carbapenemase (KPC)-producing organisms can be a difficult therapeutic challenge, due to their broad spectrum of resistance to beta-lactams. The profile of KPC beta-lactam resistance is similar to that of extended-spectrum beta-lactamases (ESBLs) with the addition of resistance to the carbapenems (imipenem, meropenem, doripenem, and ertapenem). In addition, KPC producers may be cross-resistant to a wide range of antibiotic classes including the aminoglycosides, fluoroquinolones, and sulfa drugs.
Effective Date:	June 13, 2011
Test Code:	18869
CPT Code(s):	87184
Specimen Requirements:	1 pure <i>Enterobacteriaceae</i> slant (preferred), swab or plate Organism identification must be supplied
Rejection Criteria:	Mixed isolates, organisms other than <i>Enterobacteriaceae</i>, nonviable organisms, and frozen samples are all unacceptable for this test.
Transport Temperature:	Room temperature
Specimen Stability:	Room temperature: Varies with species and transport system Refrigerated: Varies with species and transport system Frozen: Unacceptable
Reference Ranges:	Negative
Methodology:	Modified Disk Diffusion
Performing Site:	Focus Diagnostics, Inc. and Quest Diagnostics Nichols Institute, Chantilly

Test Changes

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

ColoVantage™ (methylated Septin 9)	
Effective Date:	March 11, 2011
Test Code:	16983
Specimen Requirements:	10 mL EDTA (lavender-top tube) plasma (5 mL minimum) This test requires 10mL of plasma (minimum volume 5 mL. Samples with less than 5 mL of plasma will be rejected). To obtain this volume of plasma collect blood in two (2) 10 mL EDTA (lavender-top) tubes or five (5) standard EDTA (lavender-top) tubes. Centrifuge the blood samples, separate plasma and combine them. Send plasma in a single 10 mL pour-off tube. If a 10 mL pour-off tube is not available, two (2) 5 mL pour-off tubes can be used.
Performing Site:	Quest Diagnostics Nichols Institute, Chantilly

Sex Hormone Binding Globulin			
Effective Date:	June 6, 2011		
Test Code:	30740		
Rejection Criteria:	Gross hemolysis		
Reference Ranges:	< 3 years:	Not Established	
		Male	Female
	3 - 9 years:	32-158 nmol/L	32-158 nmol/L
	10 – 13 years:	20-166 nmol/L	24-120 nmol/L
	14 – 17 years:	20-87 nmol/L	12-150 nmol/L
	Tanner Stages (7-17 years)	Male	Female
	Tanner I	47-166 nmol/L	47-166 nmol/L
	Tanner II	23-168 nmol/L	25-129 nmol/L
	Tanner III	23-168 nmol/L	25-129 nmol/L
	Tanner IV	21-79 nmol/L	30-86 nmol/L
	Tanner V	9-49 nmol/L	15-130 nmol/L
	18 – 55 years	10-50 nmol/L	17-124 nmol/L
	> 55 years	22-77 nmol/L	14-73 nmol/L
Additional Information:	Please note this test is included in the following code: 14966 Testosterone, Free, Bioavailable, and Total LC/MS/MS		

GlycoMark®	
Effective Date:	June 6, 2011
Test Code:	19599
Rejection Criteria:	Gross hemolysis
Additional Information:	Please note this test is reflexed from the following codes: 16320 - Hemoglobin A1c w/eAG with Reflex to GlycoMark® 16715- Hemoglobin A1c w/Reflex to GlycoMark®

Beta-2 Transferrin	
Effective Date:	June 13, 2011
Test Code:	10640
Assay Category:	FDA Approved
Performing Site:	Quest Diagnostics Nichols Institute, Chantilly

IGF-I, LC/MS					
Effective Date:	June 13, 2011				
Test Code:	16293				
<i>Former Test Name:</i>	<i>IGF-I, Electrochemiluminescence</i>				
Specimen Requirements:	0.5 mL serum (0.3 mL minimum) Sodium heparin (green-top) and EDTA (lavender-top) plasma are acceptable.				
Rejection Criteria:	SST (red-top/glass) tubes				
Specimen Stability:	Room temperature: 48 hours Refrigerated: 5 days Frozen: 72 hours -70 degrees: 21 days				
Reference Ranges:	Pediatric:				
	Age (Years)	Female	Age (Years)	Male	
	<1*	17-185 ng/mL	<1*	< or =142 ng/mL	
	1-1.9*	15-175 ng/mL	1-1.9*	< or =134 ng/mL	
	2-2.9*	16-178 ng/mL	2-2.9*	< or =135 ng/mL	
	3 - 3.9	38-214 ng/mL	3 - 3.9	30-155 ng/mL	
	4 - 4.9	34-238 ng/mL	4 - 4.9	28-181 ng/mL	
	5 - 5.9	37-272 ng/mL	5 - 5.9	31-214 ng/mL	
	6 - 6.9	45-316 ng/mL	6 - 6.9	38-253 ng/mL	
	7 - 7.9	58-367 ng/mL	7 - 7.9	48-298 ng/mL	
	8 - 8.9	76-424 ng/mL	8 - 8.9	62-347 ng/mL	
	9 - 9.9	99-483 ng/mL	9 - 9.9	80-398 ng/mL	
	10 - 10.9	125-541 ng/mL	10 - 10.9	100-449 ng/mL	
	11 - 11.9	152-593 ng/mL	11 - 11.9	123-497 ng/mL	
	12 - 12.9	178-636 ng/mL	12 - 12.9	146-541 ng/mL	
	13 - 13.9	200-664 ng/mL	13 - 13.9	168-576 ng/mL	
	14 -14.9	214-673 ng/mL	14 -14.9	187-599 ng/mL	
	15 -15.9	218-659 ng/mL	15 -15.9	201-609 ng/mL	
	16 - 16.9	208-619 ng/mL	16 - 16.9	209-602 ng/mL	
	17 - 17.9	185-551 ng/mL	17 - 17.9	207-576 ng/mL	
*Brabant G, et al. Serum insulin-like growth factor I reference values for an automated chemiluminescence immunoassay system: Results from a multicenter study. Horm. Res. 2003;60:53-60.					
Female Tanner Stages (based on breast stage)					
Age (Years)	1	2	3	4,5	UOM
8-8.9	80-307	84-414	197-642	388-871	ng/mL
9-9.9	92-332	91-432	197-642	358-823	ng/mL
10 -10.9	105-359	99-451	197-642	330-776	ng/mL
11 -11.9	118-387	107-470	197-642	304-731	ng/mL
12 -12.9	133-416	115-490	197-642	278-688	ng/mL
13 -13.9	148-447	123-510	197-642	254-646	ng/mL
Male Tanner Stages (based on on testicular volume)					
Age (Years)	1	2,3	4,5	UOM	
10 - 10.9	84-315	78-418	349-817	ng/mL	

11 - 11.9	96-341	101-478	318-765	ng/mL
12 - 12.9	109-368	127-543	289-716	ng/mL
13 - 13.9	123-396	158-614	262-668	ng/mL
14 - 14.9	138-426	192-689	236-622	ng/mL
15 - 15.9	153-457	230-769	212-578	ng/mL
Adults:	Age (Years)			
	18 – 19.9		108-548	ng/mL
	20 – 24.9		83-456	ng/mL
	25 - 29.9		63-373	ng/mL
	30 - 39.9		53-331	ng/mL
	40 - 49.9		52-328	ng/mL
	50 - 59.9		50-317	ng/mL
	60 - 69.9		41-279	ng/mL
	70 - 79.9		34-245	ng/mL
>80		34-246	ng/mL	
Z-Score (Male):			-2.0 - +2.0	SD
Z-Score (Female):			-2.0 - +2.0	SD
Methodology:	Liquid Chromatography Mass Spectrometry			
Performing Site:	Quest Diagnostics Nichols Institute, San Juan Capistrano			

Influenza A H1N1 (2009) Real-Time RT-PCR	
Effective Date:	June 20, 2011
Test Code:	16807
Specimen Requirements:	Nasal swab, Nasopharyngeal swab, Nasal aspirate or throat swab in Multi Microbe Media (M4) or V-C-M medium (green-cap) tube or equivalent (UTM). Swabs must be sterile Dacron, nylon, or rayon with plastic shafts. Place swab in sterile viral transport media containing protein stabilizer, antibiotics to inhibit bacterial fungal growth, and buffer solution (e.g. UTM, VCM, M4, M5, M6 and other media intended to transport chlamydia, mycoplasma or viruses).
Performing Site:	Focus Diagnostics, Inc.
Additional Information:	Please note this test is included in the following group code: 16255 – Respiratory Virus PCR Panel with 2009 H1N1

Influenza Type A/B RT-PCR Reflex to Influenza A H1N1 (2009) RT-PCR	
Effective Date:	June 20, 2011
Test Code:	16861
Specimen Requirements:	Nasal swab, Nasopharyngeal swab, Nasal aspirate or throat swab in Multi Microbe Media (M4) or V-C-M medium (green-cap) tube or equivalent (UTM). Swabs must be sterile Dacron, nylon, or rayon with plastic shafts. Place swab in sterile viral transport media containing protein stabilizer, antibiotics to inhibit bacterial fungal growth, and buffer solution (e.g. UTM, VCM, M4, M5, M6 and other media intended to transport chlamydia, mycoplasma or viruses).
Performing Site:	Focus Diagnostics, Inc.

<i>Clostridium difficile</i> Cytotoxin Antibody, Neutralization	
Effective Date:	June 27, 2011
Test Code:	34403
Specimen Requirements:	2 mL serum
Rejection Criteria:	Stool, other sterile body fluids, specimens beyond stability or received in inappropriate container.
Transport Temperature:	Refrigerated
Specimen Stability:	Room temperature: Unacceptable Refrigerated: 14 days Frozen: 30 days
Reference Ranges:	< or = 1:2 titer
Methodology:	Antibody Neutralization
Performing Site:	Focus Diagnostics, Inc.

Herpes Simplex Virus Type 1 & 2 DNA, Quantitative Real-Time PCR	
Effective Date:	June 27, 2011
Test Code:	19502
Specimen Requirements:	Add vaginal swab as an acceptable specimen type. Submit in Aptima Vaginal Swab Collection Kit.
Performing Site:	Focus Diagnostics, Inc.

Porphobilinogen, Quantitative, 24-Hour Urine	
Effective Date:	June 27, 2011
Test Code:	726
Reference Range:	Porphobilinogen, 24 hr Ur: <2.4 mg/24 h
Performing Site:	Quest Diagnostics Nichols Institute, San Juan Capistrano

Porphobilinogen, Quantitative, Random Urine	
Effective Date:	June 27, 2011
Test Code:	6329
Reference Ranges:	1-8 years: 0.9-2.8 mg/g creat 9-17 years: 0.5-2.0 mg/g creat >=18 years: <2.0 mg/g creat
Performing Site:	Quest Diagnostics Nichols Institute, San Juan Capistrano

Redirects

Effective June 27, 2011 the following tests previously performed at ViraCor-IBT Laboratories, Inc., will now be performed at Quest Diagnostics Nichols Institute, Valencia.

Test Name:	Test Code:
Allergen - Apple IgG	38257
Allergen - Avocado IgG	10639
Allergen - Broccoli IgG	38124
Allergen - Cabbage IgG	38118
Allergen - Carrot IgG	38122
Allergen - Cashew Nut IgG	10323
Allergen - Celery IgG	38101
Allergen - Codfish IgG	38110
Allergen - Coffee IgG	38560
Allergen - Halibut IgG	38103
Allergen - Lettuce IgG	10318
Allergen - Lobster IgG	38107
Allergen - Onion IgG	38123
Allergen - Pea IgG	38127
Allergen - Salmon IgG	38144
Allergen - Sesame Seed IgG	38136
Allergen - Spinach IgG	38120

The following information is common for all tests:

CPT Codes	86001
Specimen Requirements:	1 mL serum (0.3 mL min)
Specimen Stability:	Room temperature: 7 days Refrigerated: 14 days Frozen: 30 days
Reference Ranges:	< 2.0 mcg/mL
Assay Category:	Investigational Use Only
Performing Site:	Quest Diagnostics Nichols Institute, Valencia