



Cleveland Clinic Laboratories

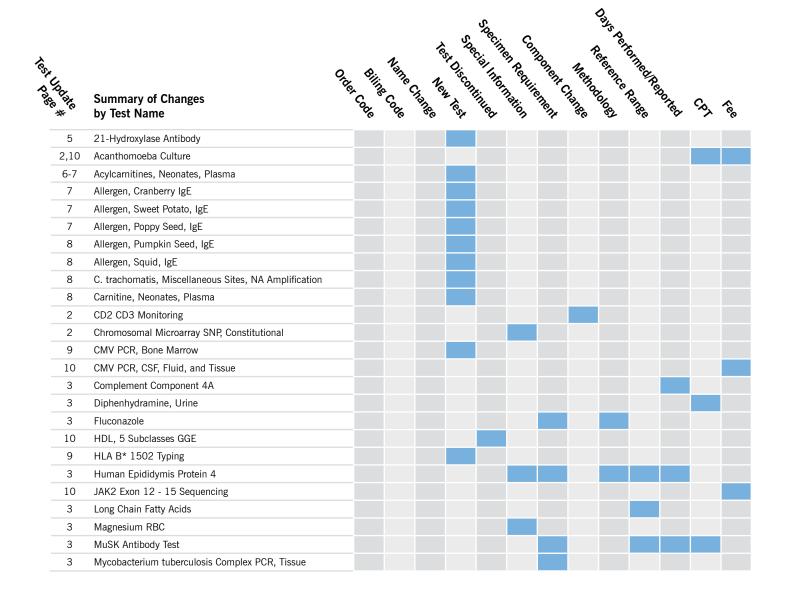
Technical Update • October 2013

Cleveland Clinic Laboratories is dedicated to keeping you updated and informed about recent testing changes. That's why we are happy to provide this technical update on a monthly basis.

Recently changed tests are **bolded**, and could include revisions to methodology, reference range, days performed or CPT code. For your convenience, tests are listed alphabetically and the order and billing codes are provided.

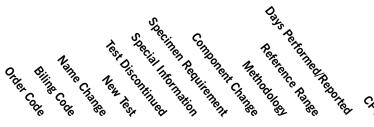
If you wish to compare the new information with previous test demographics, refer to the Test Directory, which can be accessed at clevelandcliniclabs.com.

Deleted tests and new tests are listed separately. Please update your database as necessary. For additional detail, contact Client Services at 216.444.5755 or 800.628.6816 or via email at clientservices@ccf.org.





Summary of Changes by Test Name



9	N. gonorrhoeae, Miscellaneous Sites, NA Amplification							
3	NMR Particle Analysis							
3	OncoChip Copy Number							
3	Oxalate, Serum							
10	Pancreatitis Panel							
3	Prometheus Celiac Genetics							
3	Prometheus Celiac PLUS							
9	Respiratory Viral Panel by PCR							
10	Respiratory Virus Panel							
3	Rheumatoid Factor IgM, IgG, and IgA							
4	Rufinamide							
4,10	Sex Hormone Binding Globulin							
10	Testosterone, Free, Adult Males by ED/LC-MS/MS							
4	Thyroid Stimulating Immunoglobulin							
4	Torch Antibodies, IgG & IgM							
4	Toxoplasma Antibody Evaluation, CSF							
10	Transplant Activation Markers							
4	TRAPS/Familial Hibernian Fever							
4,10	Trichomonas Prep							
5	Trichomonas vaginalis by Amplified Detection							
10	Trichomonas vaginalis RNA, Urine Qual, Males							
5	TSH Receptor Antibody							
5	VW Multimer Panel							

Test Changes

Test Name	Order Code	Billing Code	Change	Effective Date
Acanthomoeba Culture	AMBCUL	75752	CPT: 87081, 87015	11/7/2013
CD2 CD3 Monitoring	CD2MOP	82312	Includes: CD3+ Percent CD3+ Absolute Number CD2+ Percent CD2+ Absolute Number Test build will need to be changed; staff review will no longer be reported.	12/3/2013
Chromosomal Microarray SNP, Constitutional	CRMSNP	89612	Special Information: FISH to confirm CRMSNP will be added by Cytogeneticist when required for the completion of the Chromosomal Microarray test. Days performed excludes Cleveland Clinic observed holidays. Specimens are processed as soon as appropriate FISH probe(s) are received from the vendor. Patient will be charged based on the number of confirmatory probes performed. Fee is per probe and one probe is equivalent to \$3,600.00 (non-discountable) with CPT 88368x2. Focused Chromosome Study to confirm CRMSNP findings will be added by Cytogeneticist when required for the completion of the Chromosomal Microarray test. This may be added for both proband and family studies. No extra sample is required for proband; however, for parental or family studies additional 4 mL whole blood from a sodium heparin green top tube will be required. Additional fee will be \$447.00 (non-discountable) with CPT 88261, 88291.	10/8/2013

Test Changes (cont.)

Test Name	Order Code	Billing Code	Change	Effective Date
Complement Component	COMP4A	89288	Days Performed: Varies	10/3/2013
4A			Reported: 22 - 32 days	
Diphenhydramine, Urine	UDIPHN	87818	CPT: 83789	11/4/2013
Fluconazole	FLUC	80344	Specimen Requirement: 1 mL serum from a serum separator tube; Remove serum from cells within 2 hours of collection; Refrigerated	10/7/2013
			Methodology: Liquid Chromatography - Tandem Mass Spectrometry (LC-MS/MS)	
Human Epididymis Protein 4	HEP4	88459	Special Information: Electrochemiluminescence Immunoassay (ECLIA), effective 10/17/2013, will, on average, give results which are 28% higher than the previous method of Enzyme Linked Immunosorbent Assay (ELISA). The performing lab will save all samples, effective 9/14/2013, for re-baselining of positive results at no cost, if requested. Re-baselining will be available for 6 months starting 9/24/2013.	10/17/2013
			Specimen Requirement: 1 mL serum from a serum separator tube; Refrigerated	
			Method: Electrochemiluminescence Immunoassay (ECLIA)	
			Reference Range: Female: ≤ 140 pmol/L Male: Not applicable	
			Days Performed: Monday - Friday Reported: 2 - 5 days	
Long Chain Fatty Acids	LONFAT	75493	Reference Range: Phytanic Acid: < 3.00 μg/mL Pristanic Acid: < 0.3 μg/mL Ranges for all other components are unchanged	10/31/2013
Magnesium RBC	MAGRBC	82917	Special Information: This test is not approved for testing of patient samples from New York state.	11/7/2013
MuSK Antibody Test	MUSK	82592	Specimen Requirement: 2 mL serum from a red top tube; Refrigerated Reference Range: Negative: < 10 Borderline: 10 Positive: ≥ 20 Days Performed: Monday - Saturday Reported: 8 - 15 days CPT: 83519	10/31/2013
Mycobacterium tuberculosis Complex PCR, Tissue	MTPCRT	87922	Specimen Requirement: Tissue in sterile container; Specimen source must be included; Frozen Test build may need to be modified	10/24/2013
NMR Particle Analysis	NMRPAR	89316	Specimen Requirement: 2 mL serum from a red top tube; Do not use serum separator tubes. Allow specimen to clot at room temperature for 30 minutes. Separate serum from cells and refrigerate within 8 hours of collection; Refrigerated Days Performed: Varies Reported: 4 - 7 days	9/13/2013
OncoChip Copy Number	CNE	88012	CPT: 81406	10/15/2013
Oxalate, Serum	OXLATE	82970	Reference Range: 2.5 (SD 0.7) μ mol/L Days Performed: Wednesday Reported: 2 - 9 days	12/2/2013
Prometheus Celiac Genetics	CELIA	82969	Days Performed: Monday - Friday Reported: 4 - 8 days	10/17/2013
Prometheus Celiac PLUS	CEPLUS	84315	Specimen Requirement: THIS ASSAY REQUIRES MULTIPLE SPECIMEN TYPES; 2 mL serum from a serum separator tube *AND* 2 mL whole blood in an EDTA lavender top tube; Both tubes Ambient	10/17/2013
Rheumatoid Factor IgM, IgG, and IgA	RHEUMA	84429	Reference Range: Rheumatoid Factor IgG: ≤ 6 Units Rheumatoid Factor IgM: ≤ 6 Units Rheumatoid Factor IgA: ≤ 6 Units Days Performed: Monday, Wednesday, Friday Reported: 2 - 5 days	9/13/2013

Test Changes (cont.)

Test Name	Order Code	Billing Code	Change	Effective Date
Rufinamide	RUFIN	88110	CPT: 83789	11/4/2013
Sex Hormone Binding Globulin	SHBG2 79803		Specimen Requirement: 1 mL serum from a serum separator tube; Refrigerated Methodology: Electrochemiluminescence Immunoassay (ECLIA)	12/2/2013
			Reference Range: Male: 20 - 49 years: 16.5 - 55.9 nmol/L ≥ 50 years: 19.3 - 76.4 nmol/L Female: 21 - 49 years: 24.6 - 122 nmol/L ≥ 50 years: 17.3 - 125 nmol/L	
			Days Performed: Sunday - Saturday Reported: 2 - 4 days	
Thyroid Stimulating Immunoglobulin	TSIG	41070	Special Information: TSH levels > 6 mU/L may produce a weakly positive Thyroid Stimulating Immunoglobulin result.	10/3/2013
			Methodology: Chemiluminescence Immunoassay (CLIA), Bioassay	
			Reference Range: Negative: ≤ 122% of normal response Positive: ≥ 123% of normal response	
			Days Performed: Monday - Saturday Reported: 3 - 6 days	
Torch Antibodies, IgG & IgM	TORCH	79189	Days Performed: Tuesday, Thursday, Saturday Reported: 2 - 5 days	11/14/2013
Toxoplasma Antibody Evaluation, CSF	CSFTOX	81721	Reference Range: IgG: < 0.90 Index IgM: < 0.80 Index	9/30/2013
			Days Performed: Tuesday, Friday	
TRAPS/Familial	TRAPS	82892	Reported: 2 - 6 days Specimen Requirement: 4 mL whole blood in an EDTA lavender top tube;	11/14/2013
Hibernian Fever	TRAFS	02092	Refrigerated Methodology: Capillary Sequencing	11/14/2013
Trichomonas Prep	TRICHO	77786	Special Information: 1. The OSOM® Trichomonas Rapid test has not been approved for urine samples. The test has only been validated for qualitative detection of T. vaginalis antigen from vaginal swabs. 2. A negative result may be obtained if the specimen is inadequate or if the antigen concentration is below the sensitivity of the test. 3. Samples contaminated with preparations containing iodine or by the immediate prior use of vaginal lubricant are not recommended. 4. The test does not differentiate between viable and non-viable organisms nor does it differentiate between acute infection and carrier status. 5. Staph aureus in specimens at concentrations higher than 1 x 10(8) cfu/mL may interfere with the test results in negative samples. These concentrations are higher than would be expected to be present in normal patient samples. 6. Limit of Detection - The OSOM® Trichomonas Rapid test is reported to detect as little as 2,500 organisms/mL.	11/11/2013
			Specimen Requirement: Prostatic fluid Copan swab or in Stuarts medium; Ambient, if sent same day; Refrigerated, if sent next day *OR* Urethral Copan swab or in Stuarts medium; Ambient, if sent same day; Refrigerated, if sent next day *OR* Vaginal Copan swab or in Stuarts medium; Ambient, if sent same day; Refrigerated, if sent next day Urines are no longer acceptable for this test, please order Trichomonas vaginalis, Urine Qual, Females, by TMA (UTRICF) or Trichomonas vaginalis RNA, Urine Qual, Males (UTRICM) for urine samples.	
			Methodology: Immunochromatographic CPT: 87808	

Test Changes (cont.)

Test Name	Order Code	Billing Code	Change	Effective Date
Trichomonas vaginalis by Amplified Detection	VAGAMD	89383	Special Information: Urine specimen MUST be from females only; To reduce the potential for contamination ThinPrep® specimens should be poured off, using sterile technique, into the APTIMA® specimen transfer tube prior to cytology testing; This test is intended for medical purposes only and is not valid for the evaluation of suspected sexual abuse or for other forensic purposes.	10/24/2013
			Specimen Requirement: Endocervical or vaginal swab in APTIMA® Combo2 Transport Media; Refrigerated *OR* Cervical brush in Thin Prep media; Vortex Pap media and transfer 1 mL aliquot to APTIMA Combo2 transport media; Refrigerated *OR* 2 mL random urine in APTIMA® Combo2 Transport Media; For Females Only, The patient should not have urinated for at least one hour prior to specimen collection. Patient to provide a first-catch urine (approximately 20 - 30 mL of the initial urine stream) into a urine collection cup free of any preservatives. Collection of larger volumes of urine may result in specimen dilution that may reduce test sensitivity. Patients should not cleanse the labial area prior to providing the specimen. Within 24 hours of collection, transfer 2 mL of urine, using a disposable pipette, into Aptima® urine transport medium. The correct volume of urine has been added when the fluid level is between the black fill lines on the specimen transport tube label. Urine specimens must be refrigerated pending transfer into Aptima® urine transport medium. Aptima® urine medium is available through Client Services at 800.628.6816 or 216.444.5755. When complete, store and transport Refrigerated Methodology: Transcription Mediated Amplification	
TSH Receptor Antibody	TRAB	41202	Specimen Requirement: THIS ASSAY REQUIRES MULTIPLE ALIQUOT TUBES; 2 mL serum from a serum separator tube split into two separate aliquot tubes; Refrigerated Methodology: Chemiluminescent Immunoassay, Bioassay Reference Range: Thyroid Stimulating Immunoglobulins: Negative: ≤ 122% of normal response Positive: ≥ 123% of normal response TSH Binding Inhibition: < 1.0 U/L Days Performed: Thyroid Stimulating Immunoglobulins: Monday - Saturday TSH Binding Inhibition: Monday, Thursday Reported: 3 - 6 days	10/3/2013
VW Multimer Panel	VWMULP	89323	Specimen Requirement: 3 mL platelet-poor plasma from a sodium citrate light blue top tube; Frozen	9/16/2013

New Tests

Test Name	Order Code	Billing Code	Test Information	Effective Date
21-Hydroxylase Antibody	210HAB	89715	Specimen Requirement: 1 mL serum from a red top tube; Refrigerated	11/7/2013
			Methodology: Radioimmunoassay (RIA)	
			Reference Range: ≤ 1.0 U/mL	
			Days Performed: Tuesday	
			Reported: 3 - 11 days	
			CPT: 83519	
			Price: \$99.00	

Test Name	Order Code	Billing Code	Test Information	Effective Date
Acylcarnitines, Neonates, plasma	ACYLN	89673	Includes: Acylcarnitine (C2) Propionylcarnitine (C3) Isos/Butyrylcarnitine (C4) Isovaleryl-/2-Methylbutyrylcarnitine (C5) Hexanoylcarnitine (C66) 3-OH-hexanoylcarnitine (C66) 1-Octanoylcarnitine (C81) Decenoylcarnitine (C10:1) Decanoylcarnitine (C10:1) Decanoylcarnitine (C10:1) Decanoylcarnitine (C10:1) Dodecanoylcarnitine (C12:1) Dodecanoylcarnitine (C12:1) Dodecanoylcarnitine (C12:1) Dodecanoylcarnitine (C12:1) Tetradecadienoylcarnitine (C14:1) Tetradecadienoylcarnitine (C14:1) Tetradecanoylcarnitine (C16:1) Hexadecanoylcarnitine (C16:1) Hexadecanoylcarnitine (C16:1) Hexadecanoylcarnitine (C16:1) Hexadecanoylcarnitine (C16:1) S-OH-hexadecanoylcarnitine (C16:1-OH) 3-OH-hexadecanoylcarnitine (C16:1-OH) 3-OH-hexadecanoylcarnitine (C18:2) Oleylcarnitine (C18:1) Searoylcarnitine (C18:1) Separamitine (C18:1) Speciame Requirement: 0.1 ml plasma from an EDTA lavender top tube; Draw specimen just prior to a scheduled meal or feeding; Frozen Methodology: Flow Injection Analysis - Tandem Mass Spectrometry (FIA-MS/MS) Reference Range: (nmol/mL) Acylcarnitine (C2): 1 - 7 days: 2.14 - 15.89 Methodology: Flow Injection Analysis - Tandem Mass Spectrometry (FIA-MS/MS) Reference Range: (nmol/mL) Acylcarnitine (C2): 1 - 7 days: 2.14 - 15.89 Methodology: Flow Injection Analysis - Tandem Mass Spectrometry (FIA-MS/MS) Reference Range: (nmol/mL) Acylcarnitine (C3): 1 - 7 days: < 0.15 Methodology: Flow Injection Analysis - Tandem Mass Spectrometry (FIA-MS/MS) Reference Range: (nmol/mL) Acylcarnitine (C3): 1 - 7 days: < 0.16 B days - 1 month: < 1.06 Isovaleryl-/2-Methylbutyrylcarnitine (C5): 1 - 7 days: < 0.18 Methodology: Flow Injection Analysis - 1.78 B days - 1 month: < 0.23 3-OH-hexanoylcarnitine (C8:1): 1 - 7 days: < 0.18 B days - 1 month: < 0.19 Octenoylcarnitine (C8:1): 1 - 7 days: < 0.19 B days - 1 month: < 0.10 Dodecenoylcar	10/24/2013

Test Name	Order Code	Billing Code	Test Information	Effective Date
Acylcarnitines, Neonates, plasma continued			Reference Range continued: Tetradecadienoylcarnitine (14:2): 1 - 7 days: < 0.09	
Allergen, Cranberry IgE	CRANBY	89665	Specimen Requirement: 0.1 mL serum from a serum separator tube; Refrigerated Methodology: Fluorescent Enzyme Immunoassay (FEIA) by ImmunoCAP Reference Range: < 0.35 kU/L Days Performed: Sunday - Saturday Reported: 1 - 2 days CPT: 86003 Price: \$33.00	10/29/2013
Allergen, Sweet Potato, IgE	SWEPOT	89666	Specimen Requirement: 0.1 mL serum from a serum separator tube; Refrigerated Methodology: Fluorescent Enzyme Immunoassay (FEIA) by ImmunoCAP Reference Range: < 0.35 kU/L Days Performed: Sunday - Saturday Reported: 1 - 2 days CPT: 86003 Price: \$33.00	10/29/2013
Allergen, Poppy Seed, IgE	POPSED	89667	Specimen Requirement: 0.1 mL serum from a serum separator tube; Refrigerated Methodology: Fluorescent Enzyme Immunoassay (FEIA) by ImmunoCAP Reference Range: < 0.35 kU/L Days Performed: Sunday - Saturday Reported: 1 - 2 days CPT: 86003 Price: \$33.00	10/29/2013

Test Name	Order Code	Billing Cod	Test Information	Effective Date
Allergen, Pumpkin Seed, IgE	PUMKSD	89668	Specimen Requirement: 0.1 mL serum from a serum separator tube; Refrigerated Methodology: Fluorescent Enzyme Immunoassay (FEIA) by ImmunoCAP	10/29/2013
			Reference Range: < 0.35 kU/L Days Performed: Sunday - Saturday	
			Reported: 1 - 2 days	
			CPT: 86003	
			Price: \$33.00	
Allergen, Squid, IgE	SQUID	89669	Specimen Requirement: 0.1 mL serum from a serum separator tube; Refrigerated	10/29/2013
			Methodology: Fluorescent Enzyme Immunoassay (FEIA) by ImmunoCAP	
			Reference Range: < 0.35 kU/L	
			Days Performed: Sunday - Saturday	
			Reported: 1 - 2 days	
			CPT: 86003	
0 1 1 1	NAAOT	00717	Price: \$33.00	11/14/0010
C. trachomatis, Miscellaneous Sites, NA Amplification	NAACT	89717	Special Information: This assay is intended for use in clinical monitoring or management of patients; it is not intended for use in medico-legal applications. In general, this assay should not be used to assess therapeutic success or failure, since nucleic acids from these organisms may persist for 3 weeks or more following antimicrobial therapy.	11/14/2013
			Specimen Requirement: One APTIMA Collection Unisex swab from ocular, oral, anal, or rectal sites; Refrigerated	
			Methodology: Transcription Mediated Amplification	
			Reference Range: Negative	
			Days Performed: Monday - Saturday	
			Reported: 2 - 4 days CPT: 87491	
			Price: \$158.00 (non-discountable)	
Carnitine, Neonates, Plasma	NCARN	89674	Includes: Total Carnitine Free Carnitine Acylcarnitine Acylcarnitine/Free Carnitine ratio	10/24/2013
			Special Information: This test is ONLY for neonates ≤ 1 month old.	
			Specimen Requirement: 0.5 mL plasma from an EDTA lavender top tube; Frozen	
			Methodology: Flow Injection Analysis - Tandem Mass Spectrometry (FIA-MS/MS)	
			Reference Range: Total Carnitine: 1 day: 23 - 68 nmol/mL 2 - 7 days: 17 - 41 nmol/mL 8 - 31 days: 19 - 59 nmol/mL	
			Free Carnitine: 1 day: 12 - 36 nmol/mL 2 - 7 days: 10 - 21 nmol/mL 8 - 31 days: 12 - 46 nmol/mL Acylcarnitine: 1 day: 7 - 37 nmol/mL	
			2 - 7 days: 3 - 24 nmol/mL 8 - 31 days: 4 - 15 nmol/mL Acylcarnitine/Free Carnitine ratio: 1 day: 0.4 - 1.7 2 - 7 days: 0.2 - 1.4 8 - 31 days: 0.1 - 0.7	
			Days Performed: Monday - Friday	
			Reported: 3 - 6 days	
			CPT: 82379	
			Price: \$70.00 (non-discountable)	

Test Name	Order Code	Billing Code	Test Information	Effective Date
CMV PCR, Bone Marrow	CMVBM	CMVBM 89659	Special Information: This assay is only to be used for patients with a clinical history and symptoms consistent with CMV infection, and must be interpreted in the context of the clinical picture. This test should not be used to screen asymptomatic patients.	10/10/2013
			Specimen Requirement: 0.5 mL bone marrow in an EDTA lavender top tube; Refrigerated	
			Methodology: Real-Time Polymerase Chain Reaction (PCR) DNA Probe Hybridization	
			Reference Range: Negative	
			Days Performed: Monday - Saturday	
			Reported: 2 - 4 days	
			CPT: 87496	
			Price: \$366.00 (non-discountable)	
HLA B* 1502 Typing	B1502	89713	Specimen Requirement: 5 mL whole blood in an ACD A or B yellow top tube; Ambient	11/7/2013
			Methodology: Polymerase Chain Reaction (PCR) Sequence Specific Oligonucleotide Probes	
			Reference Range: Refer to report	
			Days Performed: Monday - Friday	
			Reported: 9 - 10 days	
			CPT: 81381	
			Price: \$250.00 (non-discountable)	
N. gonorrhoeae, Miscellaneous Sites, NA Amplification	NAAGC	NAAGC 89712	Special Information: This assay is intended for use in clinical monitoring or management of patients; it is not intended for use in medico-legal applications. In general, this assay should not be used to assess therapeutic success or failure, since nucleic acids from these organisms may persist for 3 weeks or more following antimicrobial therapy.	11/14/2013
			Specimen Requirement: One APTIMA Collection Unisex swab from ocular, oral, anal, or rectal sites; Refrigerated	
			Methodology: Transcription Mediated Amplification	
			Reference Range: Negative	
			Days Performed: Monday - Saturday	
			Reported: 2 - 4 days	
			CPT: 87591	
			Price: \$160.00 (non-discountable)	
Respiratory Viral Panel by PCR	RVPPCR	89720	Includes: Influenza A, Influenza A H1, Influenza A H3, Influenza A H1N1 2009, Influenza B, Respiratory Syncytial Virus A, Respiratory Syncytial Virus B, Parainfluenza Virus 1, Parainfluenza Virus 2, Parainfluenza Virus 3, Human Metapneumovirus, Rhinovirus, Adenovirus B/E, Adenovirus C	12/2/2013
			Specimen Requirement: Nasopharyngeal swab in M4; Refrigerated	
			Methodology: Reverse Transcription/Polymerase Chain Reaction (RT/PCR)	
			Reference Range: Negative	
			Days Performed: Sunday - Saturday, not performed on Cleveland Clinic holidays	
			Reported: 2 - 3 days	
			CPT: 87633	
			Price: \$440.00	

Test Name	Order Code	Billing Code	Test Information	Effective Date
Trichomonas vaginalis RNA, Urine Qual, Males	UTRICM	89708	Specimen Requirement: 2 mL random urine in a clean container; The patient should not have urinated for at least one hour prior to specimen collection. Patient to provide a first-catch urine (approximately 20 - 30 mL of the initial urine stream) into a urine collection cup free of any preservatives. Collection of larger volumes of urine may result in specimen dilution that may reduce test sensitivity. Two mL of urine specimen MUST be transferred into the Gen-Probe Aptima® Urine transport medium ASAP or within 24 hours of collection and before being assayed. Urine specimens must be refrigerated pending transfer into Aptima® urine transport medium. Aptima® Urine transport medium is available through Client Services at 800.628.6816 or 216.444.5755. Transport Refrigerated	10/24/2013
			Methodology: Transcription Mediated Amplification (TMA)	
			Reference Range: Not detected	
			Days Performed: Monday - Saturday	
			Reported: 4 - 6 days	
			CPT: 87798	
			Price: \$136.00 (non-discountable)	

Fee Increases

Test Name	Order Code	Billing Code	List Fee	CPT Code
Acanthomoeba Culture	AMBCUL	75752	\$65.00 (non-discountable)	87081, 87015
Pancreatitis Panel	PANCPL	83724	\$3,800.00 (non-discountable)	81223, 81404x2, 81224
Trichomonas Prep	TRICHO	77786	\$54.00	87808

Fee Reductions

Test Name	Order Code	Billing Code	List Fee	CPT Code
CMV PCR, CSF, Fluid, and Tissue	CMVCSF	79779	\$105.00 (non-discountable)	87496
JAK2 Exon 12 - 15 Sequencing	JAKNON	87775	\$500.00 (non-discountable)	81403
Sex Hormone Binding Globulin	SHBG2	79803	\$49.00 (discountable)	84270
Testosterone, Free, Adult Males by ED/LC-MS/MS	FTESAM	89350	\$300.00 (discountable)	84402

Discontinued Tests

Test Name	Order Code	Billing Code	Test Information	Effective Date
HDL, 5 Subclasses GG	HDLGGE	89561	This test will no longer be available.	10/3/2013
Respiratory Virus Panel	RVPAN	83312	This test will no longer be available. Suggest ordering Respiratory Viral Panel by PCR (RVPPCR)	12/2/2013
Transplant Activation Markers	TAM	38	This test will no longer be available.	12/3/2013