

2018 Surveys and Anatomic Pathology Education Programs

Performance you can measure. Accuracy you can trust.



Performance you can measure. Accuracy you can trust.

Laboratory medicine is changing at a rapid pace. Our comprehensive range of programs is constantly evolving to keep you in step with these changes, enabling you to have more time for what matters most—accuracy in the laboratory.

Preview new innovations to help simplify your efforts.

Gain insight at a glance.

Monitor performance across all graded CAP proficiency testing programs using the Performance Analytics Dashboard. This complimentary web-based reporting solution gives you fast access to a single laboratory or an expansive network's performance.

Keep pace with advances in next-generation sequencing (NGS).

Add a level of quality assurance to ensure you deliver accurate and reliable test results with seven new NGS programs. Assess germline variants and improve diagnostic skills with the new Variant Interpretation Only (VIP), our first NGS educational program to provide CME/CE credit.

Submit your laboratory's slides and receive an expert evaluation of your tissue staining preparation.

Participate in the new HistoQIP programs evaluating gynecologic biopsy preparation (HQBX4), mismatch repair protein IHC staining (HQMMR), and IHC staining of non-small cell lung carcinoma (HQNSC).

Streamline your educational efforts using our improved process for claiming CME/CE credit.

Claiming CME/CE credit associated with the anatomic pathology educational programs has been simplified to save you time.

Purchase your Surveys and manage your account online. Visit cap.org and click on the SHOP tab.

Table of Contents

2018 Surveys and Anatomic Pathology Education Programs

1	New Developments 3-6	15	Microbiology 169–200
2	Continuing Education 7–20		Bacteriology170
-	Continuing Education Programs		Mycobacteriology 183
	Competency Assessment Program 15		Mycology184
	QM <i>Ed</i> ™ Online Educational Courses 18		Parasitology187
3	Quality Management Tools 21-40		Virology 191
	Q-PROBES™24		Molecular Microbiology 193
	Q-TRACKS [®]		Infectious Disease Serology 200
,		16	Immunology and Flow Cytometry 201-212
4	Quality Cross Check 41-50		Immunology 202
5	Point-of-Care Programs 51-54		Flow Cytometry 209
6	General Chemistry and Therapeutic Drug Monitoring 55-80	17	Transfusion Medicine, Viral Markers,
	General Chemistry and Therapeutic		and Parentage Testing 213-228
	Drug Monitoring 56		Transfusion Medicine 214
	Urine Chemistry 68		Viral Markers 224
	Special Chemistry 71		Parentage Testing
7	Endocrinology 81–90	18	Histocompatibility 229-234
8	Blood Gas, Critical Care, and Oximetry	19	Genetics and Molecular Pathology 235-254
	and Oximetry 91–94		Cytogenetics 236
9	Toxicology		Biochemical and Molecular Genetics 239
10	Accuracy-Based Programs 109–114		Next-Generation Sequencing 246
	Accuracy-Based Programs 110		Molecular Oncology—Solid Tumors 250
	Validated Materials 114		Molecular Oncology—Hematologic 254
11	Instrumentation Validation Tools 115-132	20	Anatomic Pathology 255–278
	Calibration Verification/Linearity		Surgical Pathology 256
	Management Programs 129		General Immunohistochemistry 267
			Predictive Markers 269
12	Hematology and		Specialty Anatomic Pathology 270
	Clinical Microscopy		Cytopathology272
	Clinical Microscopy		-
	• •	21	Forensic Sciences 279–284
13	Reproductive Medicine	22	Analyte/Procedure Index 285-332
14	Coagulation 157–168	23	Program Code Page Index 333-336

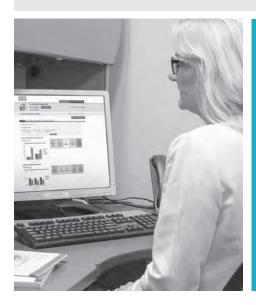
Insight at a glance.



In just seconds, the CAP's Performance Analytics Dashboard provides valuable insights into your laboratory's performance, letting you proactively focus energy on areas that need immediate attention while filtering out distractions. Updated daily, this complimentary Surveys and CAP accreditation performance monitoring tool reduces the stress of managing today's laboratory by giving you fast access to a single laboratory's or an expansive network's performance.

To view a demo, search Performance Analytics Dashboard at cap.org.

New Developments



Simplify analysis and reporting of PT and accreditation performance using the Performance Analytics Dashboard.

- Quickly identify trends/patterns to mitigate risk with the ability to access up to three years or three accreditation cycles of data.
- Benchmark your laboratory against your peers and CAP-wide performance.
- Consolidate multiple CAP numbers to view a single dashboard for an entire system.

New Developments

Quality Management Tools				
Subsection	Name	Program Code	Page(s)	
Q-PROBES™	Physician Satisfaction with Clinical Laboratory Services	QP181	25	
Q-PROBES	Laboratory Staff Turnover	QP182	26	
Q-PROBES	Technical Competency Assessment of Body Fluid Slide Review	QP183	27	
Q-PROBES	Laboratory Result Turnaround Time for Emergency Room Specimens	QP184	28	

Quality Cross Check				
Section	Name	Program Code	Page(s)	
Quality Cross Check	Quality Cross Check—Reticulocyte Series	RTQ, RT2Q, RT3Q, RT4Q	47	

General Chemistry and Therapeutic Drug Monitoring				
Subsection	Name	Program Code	Page(s)	
Special Chemistry	Trace Metals, Whole Blood	TMWB	79	

Endocrinology					
Section	Name	Program Code	Page(s)		
Endocrinology	Noninvasive Prenatal Testing	NIPT	87		

Toxicology						
Section			Name		Program Code	Page(s)
Toxicology			Trace Metals, Whole Blood		TMWB	103
Toxicology	Delayed	until 2019	Toxicology Quality Program		TQP	108

Hematology and Clinical Microscopy				
Subsection	Name	Program Code	Page(s)	
Hematology	Quality Cross Check—Reticulocyte	RTQ, RT2Q, RT3Q, RT4Q	140	

Microbiology					
Subsection	Name	Program Code	Page(s)		
Bacteriology	MRSA Screen, Molecular, 2 Challenge	MRS2M	179		
Bacteriology	MRSA Screen, Molecular, 5 Challenge	MRS5M	179		
Parasitology	Expanded Parasitology	PEX	189		
Molecular Microbiology	Vector-Borne Disease—Molecular	VBDM	195		

Immunology and Flow Cytometry					
Subsection	Name	Program Code	Page(s)		
Immunology	Alpha-2-Macroglobulin	A2MG	204		
Flow Cytometry	B-ALL Minimal Residual Disease	BALL	210		
Flow Cytometry	Flow Cytometry, Plasma Cell Neoplasms	PCNEO	211		

Coagulation					
Section	Name	Program Code	Page(s)		
Coagulation	Apixaban Anticoagulant Monitoring	APXBN	161		

Transfusion Medicine, Viral Markers, and Parentage Testing				
Subsection	Name	Program Code	Page(s)	
Viral Markers	Vector-Borne Disease—Molecular	VBDM	226	

Genetics and Molecular Pathology					
Subsection	Name	Program Code	Page(s)		
Biochemical and Molecular Genetics	Variant Interpretation Only	VIP/VIP1	244		
Biochemical and Molecular Genetics	Noninvasive Prenatal Testing	NIPT	245		
Next-Generation Sequencing	Next-Generation Sequencing Undiagnosed Disorders—Exome	NGSE	248		
Next-Generation Sequencing	Next-Generation Sequencing Bioinformatics Somatic Validated Material	NGSBV	249		
Molecular Oncology—Solid Tumors	IGHV Mutation Analysis	IGHV	250		
Molecular Oncology—Solid Tumors	Cell Free DNA	CFDNA	252		
Molecular Oncology—Solid Tumors	RNA Sequencing	RNA	252		

	Anatomic Pathology		
Subsection	Name	Program Code	Page(s)
Surgical Pathology	CAP/NSH HistoQIP Mismatch Repair IHC	HQMMR	264
Surgical Pathology	CAP/NSH HistoQIP Non-small Cell Lung Carcinoma IHC	HQNSC	265
Surgical Pathology	CAP/NSH HistoQIP Specialty Series Gynecologic Biopsy	HQBX4	266

Continuing Education



We support laboratory professionals. Maintain your certification with Surveys continuing education (CE).

- Offer your staff more than 100 CE credits.
- Enhance your learning with CE content that is tightly integrated with proficiency testing challenges.
- Meet certification and licensure requirements with CE across multiple disciplines.

Continuing Education

Continuing Education Programs	15
New Programs NEW	
Quality Culture QM <i>Ed</i> Course (ISOEDCL)	19

Continuing Education Programs

Your laboratory demonstrates its commitment to quality by choosing CAP Surveys programs. You'll find the same level of quality in the CAP Continuing Education Programs.



CME (Continuing Medical Education for Physicians)

Accreditation

The College of American Pathologists (CAP) is accredited by the Accreditation Council for Continuing Medical Education (ACCME) to provide continuing medical education for physicians.

CME Category 1

The CAP designates these enduring materials educational activities for a maximum of the stated number of *AMA PRA Category 1 Credits*™. Physicians should claim only the credit commensurate with the extent of their participation in the activity.

Note to CME participants of enduring* materials courses:

An AMA requirement mandates that all physicians wishing to claim CME credits must pass a scored assessment. All CAP enduring materials CME courses require participants to pass a scored assessment prior to claiming credit.

*Enduring courses are those courses that endure over time, such as print or online courses.



credit CE (Continuing Education for Nonphysicians)

The CAP designates these educational activities for a maximum of the stated number of credits of continuing education. Participants should claim only the credit commensurate with the extent of their participation in the activity.

The American Society for Clinical Pathology (ASCP) Board of Certification (BOC) Certification Maintenance Program (CMP) accepts these activities to meet its continuing education requirements. The states of California and Florida also approve these activities for continuing education credit.

Cytotechnologists may apply the credits from the PAP Education (PAPCE/PAPJE/PAPKE/PAPLE/PAPME), NGC, FNAG, FNA, and TICP programs toward the required educational activities for the American Society of Cytopathology (ASC) Continuing Education Credit Program (CECC) and the International Academy of Cytology (IAC).



credif This activity is eligible for continuing medical education (CME) credit or continuing education (CE) credit.

Surveys Continuing Education Activities

When your laboratory participates in CAP Surveys, every member of your team can enroll in education activities and earn continuing education (CE) credit at no additional charge. Simply follow these steps:

- 1. Establish a free Web account.
- 2. Complete a reading provided in the Participant Summary or Final Critique.
- 3. Answer online learning assessment questions.
- 4. Claim CE certificate.

Each member of your staff can access the Surveys education activities for a maximum of 12 months.

Surve	ys Educational Activi	ties	l
Program Name	Program Code	Discipline	Catalog Page(s)
General Chemistry and Therapeutic Drugs	C1, C3, C3X, C4, C7, CZ, CZX, CZ2X, Z	Chemistry	56-58
Quality Cross Check—Whole Blood Glucose	WBGQ	Chemistry	67
Endocrinology	Y, YY, DY, BU, BGS, RAP, ING, EPO	Chemistry	84-86, 8
Coagulation, Limited	CGL, CGB, CGDF	Coagulation	158
CAP/ACMG Cytogenetics	CY, CYBK	Cytogenetics	236
Basic Hematology	HE, HEP	Hematology and Clinical Microscopy	134
Blood Cell Identification, Limited	BCP, BCP2	Hematology and Clinical Microscopy	134
Hematology Automated Differentials FH Series	FH1-FH13, FH1P-FH13P	Hematology and Clinical Microscopy	135
Virtual Peripheral Blood Smear	VPBS	Hematology and Clinical Microscopy	141
Bone Marrow Cell Differential	BMD	Hematology and Clinical Microscopy	138
Clinical Microscopy	CMP, CMP1	Hematology and Clinical Microscopy	144
CAP/NSH HistoQIP	HQIP	Histology	263
Immunology	IG, IGX, ANA, ASO, CRP, HCG, IM, RF, RUB, IL, M/OLI, G, LPE/ SPE/UBJP, RDS/CCP, S2, S4, S5	Immunology	74, 76, 202-203 206-203
D-Dimer Calibration Verification/Linearity	LN42	Instrumentation	128
Bacteriology	D	Microbiology	170
Mycology and Aerobic Actinomycetes	F	Microbiology	184
Limited Bacteriology	D1, D2, D3, D4, D5, D6, D7, MC1, MC2, MC3, MC4, MC5	Microbiology	172-17
Parasitology	P, P3, P4, P5	Microbiology	187
Sperm Count, Motility, Morphology, and Viability	SMCD, SM1CD, SM2CD	Reproductive Medicine	154
Semen Analysis	SC, SC1, PV, SM, SV, ASA	Reproductive Medicine	154
Synthetic Cannabinoid/Designer Drugs	SCDD	Toxicology	105
CAP/AACC Urine Drug Testing, Screen	UDS, UDS6	Toxicology	98
Oral Fluid for Drugs of Abuse	OFD	Toxicology	100
Drug Monitoring for Pain Management	DMPM	Toxicology	106
Forensic Urine Drug Testing, Confirmatory	UDC	Toxicology	99
CAP/AACC Alcohol/Ethylene Glycol/Volatiles	AL1	Toxicology	101
Ethanol Biomarkers	ETB	Toxicology	102
Drug-Facilitated Crime	DFC	Toxicology	107
Serum Drug Screening	SDS	Toxicology	101
Trace Metals, Urine	TMU	Toxicology	103
Transfusion Medicine	J, J1, JE1, JAT, JATE1, EXM, EXM2	Transfusion Medicine	214-216

Surveys Self-Reported Training Opportunities

When your laboratory participates in CAP Surveys, every member of your team can receive self-reported training opportunities.

Self-Repor	ted Training Oppor	tunities*	
Program Name	Program Code	Source	Catalog Page(s)
Quality Management Tools			
QP181 - Physician Satisfaction with Clinical Laboratory Services NEW	QP181	Final Critique	25
QP182 - Laboratory Staff Turnover NEW	QP182	Final Critique	26
QP183 - Technical Competency Assessment of Body Fluid Slide Review NEW	QP183	Final Critique	27
QP184 - Laboratory Result Turnaround Time for Emergency Room Specimens NEW	QP184	Final Critique	28
Hematology and Clinical Microscopy			
Blood Cell Identification	BCP, BCP2	Participant Summary	134
Bone Marrow Cell Differential	BMD	Participant Summary	138
Extended Virtual Peripheral Blood Smear	EHE1	Participant Summary	142
Hematology Automated Differentials FH Series	FH1-FH13, FH1P-FH13P	Participant Summary	135
Basic Hematology	HE, HEP	Participant Summary	134
Hemoglobinopathy	HG	Participant Summary	139
Virtual Body Fluid	VBF	Participant Summary	146
Virtual Peripheral Blood Smear	VPBS	Participant Summary	141
Clinical Microscopy	CMP, CMMP, CMP1	Participant Summary	144- 145
Microbiology			
Blood Parasite	BP	Participant Summary/Final Critique	188
Expanded Bacteriology	DEX	Participant Summary/Final Critique	171
Mycobacteriology	E	Participant Summary/Final Critique	183
Yeast	F1	Participant Summary/Final Critique	184
Parasitology	Р	Participant Summary/Final Critique	187
Ticks, Mites, and Other Arthropods	TMO	Participant Summary	189
Worm Identification	WID	Participant Summary	189

^{*}Notes:

[•] CAP Self-Reported Training opportunities do not offer CE credit, but can be used toward fulfilling requirements for certification maintenance by agencies such as the American Society for Clinical Pathology (ASCP). Please verify with your certifying agency to determine your education requirements.

[•] These opportunities are subject to change. Refer to the Participant Summary/Final Critique for availability.

Maintenance of Certification (MOC)

Maintenance of Certification (MOC) is the board certification program that involves continuous professional development and ensures that an American Board of Pathology (ABP) board-certified pathologist is committed to lifelong learning and competency in a specialty and/or subspecialty.

There are six competency categories defined by the American Board of Medical Specialties (ABMS) and endorsed by the ABP to fulfill specific MOC requirements. They are listed below with their descriptions.

All CAP education activities providing CME credits meet the MOC Part II: Lifelong Learning requirements. Some programs will meet the requirements for Self-Assessment Module (SAM) and/or MOC Part IV at the laboratory or the individual levels. Programs that meet Part IV are identified within the description of the program. Visit the CAP website for the current list of programs that meet the requirements for MOC Part II and Part IV.

Interpersonal and Communication Skills

Demonstrate interpersonal and communication skills that result in effective information exchange and teaming with patients, patients' families, and professional associates.

Medical Knowledge

Demonstrate knowledge of established and evolving biomedical, clinical, and cognate sciences and the application of this knowledge to pathology.

Practice-Based Learning and Improvement

Demonstrate ability to investigate and evaluate diagnostic and laboratory practices in your own laboratory, appraise and assimilate scientific evidence, and improve laboratory practices and patient care.

Patient Care

Demonstrate a satisfactory level of diagnostic competence and provide appropriate and effective consultation in the context of pathology services.

Professionalism

Demonstrate a commitment to carrying out professional responsibilities, adherence to ethical principles, and sensitivity to diverse patient population.

Systems-Based Practice

Demonstrate understanding of and contribution to local, regional, and national health care systems, and support health care in systems-based practice definition.



Self Assessment Module: This activity fulfills the SAM credit requirements for MOC and is therefore eligible for SAM credit. Participants who successfully complete an online assessment may apply their earned credit(s) to the ABP's SAM requirements.

Note to CME/CE participants: The AMA mandates that all education providers (such as the CAP) require participants pass assessment questions in an enduring* program in order to earn and claim CME credits. All participants in any activity granting CME/CE will be required to complete and pass assessment questions before claiming their credits.

For CME/SAM activities ONLY: Participants have a total of three opportunities to take and pass the post-test, with feedback provided after each question. The AMA requires that participants pass the post-test in an enduring program to claim credit; therefore, if they do not pass, they cannot claim credit.

*Enduring programs are those courses that endure over time such as print or online courses.

	Educ	cation Programs			
Program Name	Program Code	Maximum AMA PRA CME Category 1 Credits™ Annually	Maximum CE Credits Annually	Format	Catalog Page
Autopsy Pathology	AUP/AUP1	12.5***	NA	Online	270
Clinical Pathology Improvement Program*	CPIP/CPIP1	15***	NA	Online	14
Digital Slide Program in Dermatopathology*	DPATH/DPATH1	15***	NA	Online (DigitalScope®)	259
Digital Slide Program in FNA*	FNA/FNA1	10	10	Online (DigitalScope)	277
Fine-Needle Aspiration Glass Slides	FNAG/FNAG1	10	10	Glass Slides	278
Forensic Pathology	FR/FR1	12	12	Online	281
Digital Slide Program in Hematopathology	HPATH/HPATH1	12***	12	Online (DigitalScope)	143
Nongynecologic Cytopathology Education**	NGC/NGC1	25	25	Glass Slides With Online Cases (DigitalScope)	276
Neuropathology Program	NP/NP1	10****	NA	Online (DigitalScope)	271
Gynecologic Cytopathology PAP Education Program***	PAPCE/APAPCE PAPUE/APAPUE PAPKE/APAPKE PAPLE/APAPLE PAPME/APAPME Series 1 or 2	8	8	Glass Slides	273
Glass Slide Cytopathology PAP PT Program (with Glass Slide PAP Education)***	PAPCPT/APAPCPT PAPUPT/APAPUPT PAPKPT/APAPKPT PAPLPT/APAPLPT PAPMPT/APAPMPT	8	8	Glass Slides	272
Cancer Staging Improvement Program	PCSP/PCSP1	5***	NA	Online (DigitalScope)	262

^{*}Program is available for purchase online. Go to cap.org and choose the Learning tab.

 $^{{\}tt **NGC\ provides\ up\ to\ 20\ CME/CE\ credits\ for\ the\ glass\ slides\ and\ 5\ CME/CE\ credits\ for\ the\ online\ slide\ portion\ of\ the\ program.}$

^{***}PAP provides up to 8 CME/CE credits for glass slides.

^{****}SAM credits are included in CME totals for the appropriate programs.

	Educ	cation Programs			
Program Name	Program Code	Maximum AMA PRA CME Category 1 Credits™ Annually	Maximum CE Credits Annually	Format	Catalog Page
Performance Improvement Program in Surgical Pathology	PIP/PIP1	40	NA	Glass Slides	256
Online Performance Improvement Program in Surgical Pathology*	PIPW/PIPW1	40	NA	Online (DigitalScope)	257
Nongynecologic Cytopathology Intraoperative Touch Imprint/ Crush Preparation Program*	TICP/TICP1	10****	10	Online (DigitalScope)	261
Variant Interpretation Only	VIP/VIP1 NEW	3	3	Online	244
Virtual Biopsy Program*	VBP/VBP1	25****	NA	Online (DigitalScope)	258

^{*}Program is available for purchase online. Go to cap.org and choose the Learning tab.

System Requirements

DigitalScope is a Web-based whole slide image (WSI) retrieval and viewing system. DigitalScope is supported with Microsoft Internet Explorer 11.0 (limited support for IE 9 and 10) or later, Firefox 4.0 or later, Safari 3, and the latest Google Chrome version.

For the most up-to-date information on system requirements, go to cap.org and select CONTACT & SUPPORT. The download speed and the appearance of the activity will vary depending on the type and speed of your Internet connection, computer's power, and browser.

Have you created or updated your CAP Profile?

Each laboratory staff member should have their own profile. Your profile is transferrable when you leave your current position. Use it to maintain information about yourself, including:

- · Business affiliations
- Personal contact information

Certifications

- · Specialties and skills
- Contact preferences
- Addresses
- Inspector-related information

To create or update your profile, visit cap.org, log in, and click on UPDATE MY PROFILE.



^{****}SAM credits are included in CME totals for the appropriate programs.

Clinical Pathology Improvement Program (CPIP)

The Clinical Pathology Improvement Program (CPIP) delivers 12 online clinical laboratory cases to study—one per month—and an opportunity to earn up to 15 CME/SAM credits annually. Assess and improve clinical pathology skills and fulfill Maintenance of Certification (MOC) requirements.

CPIP cases feature real-life case scenarios, including images and clinical background. Participants work through sequentially revealed information and a series of prompts to arrive at a resolution—just as in the laboratory.

Cases include thought-provoking questions with feedback and a multiple-choice post-test. Participants who earn passing scores on post-tests may apply their earned credits to the ABP's MOC SAM requirements.

Clinical Pathology II CPIP	mprovement Pr /CPIP1	ogram
Program Name	Program Code	Cases/Year
	CPIP/CPIP1	
Online cases in clinical pathology		12

Additional Information

Pathologists and residents can use CPIP online to assess and improve their skills in clinical pathology.

- Case topics may originate from the ABP's general listing suggested for MOC including laboratory administration and operations, transfusion medicine, chemistry, coagulation, hematology, immunology, microbiology, and molecular genetic pathology.
- Cases may include patient history, case-related static images, and whole slide images.
- Monthly individual CPIP cases can also be ordered online. Go to cap.org and choose the Learning tab. To order both CPIP and CPIP1, please call 800-323-4040 or 847-832-7000 option 1.

Program Information

- One online clinical laboratory case per month
- CPIP1 Additional pathologist (within the same institution) reporting option with CME/SAM credit; must order in conjunction with CPIP
- Earn a maximum of 15 CME/SAM credits (AMA PRA Category 1 Credits™) per year
- This activity meets the ABP MOC Part IV Practice Performance Assessment requirements
- Twelve cases per year; your CAP shipping contact will be notified <u>via email</u> when the activity is available





Competency Assessment Program

About one of every four laboratories is cited for a deficiency related to its competency assessment records. You can avoid becoming a part of this statistic.

Competency Assessment Program

The CAP's Competency Assessment Program helps keep you in compliance by managing your personnel's competency assessment performance and records. Use the CAP's Competency Assessment Program to track compliance to all six of the elements of competency assessment as defined by CLIA. Customizable to fit your specific laboratory's procedures, Competency Assessment Program offers benefits that simplify your documentation process.

- **Be organized.** Stay on top of your documentation and records with easy-to-use management reports, employee progress tracking, and individual employee transcripts so your laboratory is inspection-ready at all times.
- Obtain real-time results. Generate management reports with just a few clicks.
- Strengthen your learning. The program comes ready with multiple relevant, applicable courses already loaded, and new courses are added every six months. Plus, if employees need a refresher learning opportunity, reassessment courses are included.
- Customize training to your needs. If the wide selection of ready-made training courses (Pro Courses) doesn't meet your needs, customize them. You can match courses to your laboratory's exact standard procedures.
- Save time. Tools like ChecklistBuilder, CourseBuilder, and Competency Profiles allow your administrators easy, convenient methods to document all six areas of competency as defined by CLIA and the CAP Laboratory Accreditation Program.
- Access anywhere. The Competency Assessment Program is cloud based, so it's available 24/7 from any PC, laptop, or tablet—wherever you have an Internet connection. Courses are available for users throughout the subscription period.
- Stay focused. Use instrument-specific checklists for assessing competency and training.
- Remain in compliance. Many of the ready-made educational courses provide your staff the opportunity to earn CE credits.

Add Safety & Compliance Courses Especially Developed for the Laboratory

As an add-on option, Competency Assessment Program offers a package of seven non-credit, complementary safety and compliance courses—appropriate for annual laboratory-specific compliance training and for clinical laboratory science students prior to clinical rotations. These courses include:

- · OSHA Bloodborne Pathogens
- OSHA Hazard Communication and Chemical Hygiene
- · OSHA Electrical Safety
- · OSHA Fire Safety
- OSHA Formaldehyde
- Tuberculosis Awareness for Health Care Workers
- · Medical Error Prevention: Patient Safety

The CAP updates these courses as necessary to reflect changes in regulations or best practices.

Please see next pages for all course descriptions. For more information, visit cap.org and choose Learning for Laboratory Professionals via the Learning tab.

	Assessment Course Schedu	le
Discipline	January 2018 Release	July 2018 Release
Blood Banking/Transfusion Medicine—Generalist	Antibody screen and ID	Transfusion reactions
Blood Banking/Transfusion Medicine—Specialist	Antibody screen and ID	Transfusion reactions
Chemistry	Liver and renal testing	Chemistry QC, calibration, and reportable range
Hematology and Coagulation	Common coagulation tests	Platelet testing, morphology, and disorders
Histology	Safety issues in the histology laboratory	Special stains
Immunology	Qualitative HIV testing	Molecular amplification methods for detection of infectious diseases
Microbiology—Generalist	Gram stain: organism detection and differentiation	Urine and body fluid cultures
Microbiology—Specialist	Gram stain: organism detection and differentiation	Urine and body fluid cultures
Phlebotomy/Specimen Processing	Challenges of phlebotomy: pediatric blood collection, alternate sites, and difficult draws	Specimen collection for workplace urine drug testing programs and forensic drug and alcohol testing
Point-of-Care Testing	Whole blood prothrombin time and INR (PT/INR) testing	Cardiac biomarkers
Quality Programs/Management	Laboratory management: monitoring the quality control program	Competency evaluation
Safety	Fire and electrical safety	Ergonomics
Urinalysis/Body fluids	Cerebrospinal fluid analysis	Semen analysis

	Pro Course Schedule	
Discipline	January 2018 Release	July 2018 Release
Blood Banking/Transfusion Medicine	Direct antiglobulin test	ABO typing discrepancies
Chemistry	Clinical toxicology	Electrolytes, acid base and anion gap
Hematology and Coagulation	Erythrocyte morphology	White blood cell inclusions
Histology	Immunohistochemistry, part 2	Histology specimen handling
Immunology	Monitoring the testing process in immunology	Human chorionic gonadotropin and fetal fibronectin
Microbiology	Genital tract pathogens	Microbiology of wounds
Phlebotomy/Specimen Processing	Phlebotomy professionalism and ethics	Venipuncture
Point-of-Care Testing	Provider performed testing	Urine dipstick
Quality Programs/Management	Document control	New instrument method validation
Safety	Hazardous chemicals	Laboratory waste and spill management
Urinalysis/Body Fluids	Microscopic urinalysis, part 2—crystals and casts	Serous and synovial fluids

Safety & Compliance Courses

OSHA Bloodborne Pathogens. Addresses the OSHA Bloodborne Pathogens standard as it applies to clinical and medical laboratories. Covers major bloodborne pathogens, including hepatitis B and HIV. Focuses on proper handling of sharps, personal protective equipment (PPE), engineering controls such as microbiological safety cabinets, and proper work practices like handwashing.

OSHA Hazard Communication and Chemical Hygiene. Describes the OSHA Chemical Hygiene Standard and helps satisfy OSHA requirements for annual training. Explains Haz-Com, the National Fire Protection Agency diamond, the Safety Data Sheet, and common-sense laboratory safety rules applied to clinical laboratory practice.

OSHA Electrical Safety. Addresses electrical safety and electrical hazards commonly found in the clinical laboratory. Covers prevention and safety measures, fighting electrical fires, and treatment of electrical injuries.

OSHA Fire Safety. Teaches the basics of fire safety in the clinical laboratory, including classes of fire and key acronyms, such as PASS and RACE. Addresses fire prevention, drills, and firefighting techniques.

OSHA Formaldehyde. Covers essentials for any laboratory that uses formaldehyde or formalin. Shares facts about formaldehyde, safety risks, proper handling procedure, monitoring, spill clean-up, and personal protective equipment.

Tuberculosis Awareness for Health Care Workers. Provides background information about spread of tuberculosis, purified protein derivative (PPD) testing procedures, CDC guidelines, and methods of control.

Medical Error Prevention: Patient Safety. Includes potential causes of medical errors in the clinical laboratory, important legislation and definitions, and steps laboratory professionals can take to reduce the impact of medical errors in their workplace. Serves as an ideal part of an effective medical error reduction program. Appropriate for both experienced and newer laboratory personnel.

Note: The Safety & Compliance courses are not available for purchase separately. The courses listed above do not offer CE credit.

So you're going to collect a blood specimen

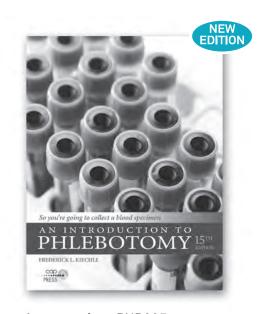
Up to 70% of laboratory errors occur prior to sample analysis and testing. Ensure everyone on your team is equipped to procure a quality blood specimen with this modern update to the classic reference guide.

- Step-by-step instructions for venipuncture, skin puncture, and infant heelstick
- Best practices for collection, transporting, processing, and storage
- Procedures for blood smears, blood cultures, and neonatal screening
- Special considerations for the difficult venipuncture
- · Four ways to inspire confidence in your patient

Buy multiple copies and save. Call 800-323-4040 option 1 (Country Code 001).

Or, view sample pages and order online:

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Item number: PUB225 Spiral bound; 84 pages; 30+ images and tables; 2017

QMEd[™] Online Educational Courses

Learn quality tools and techniques with case examples from medical laboratories.



Program information

- CAP Quality Management Educational Resources (QMEd) courses help you improve your processes and eliminate waste.
- CAP QMEd courses help you build a quality management system—one piece at a time—that sustains your continuous improvement and Lean efforts.
- CAP QMEd courses are delivered online via a highly interactive user interface that allows you to learn at your own pace.
- All CAP QMEd courses are licensed for one year, allow sharing of logins, and include continuing education (CE) credit.

CAP online interactive QMEd courses will help you:

- · Understand the concept of a quality management system
- · Self-assess your current QMS against international quality standards
- · Plan and resource for the development of your QMS
- Interpret ISO 15189 requirements
- · Improve your document control system
- · Perform internal audits using tracer audit and process audit methods
- · Implement and refine occurrence management with root cause analysis
- · Write an effective quality manual
- · Measure, analyze, and set goals with senior management

About the Courses

15189 Walkthrough

Order ISOEDWT

Designed for laboratory quality managers (along with your medical and administrative decision makers) considering implementation of an ISO 15189 program. Summarizes each section of the standard, while clarifying its intent and key requirements. Hear directly from the CAP's ISO 15189 assessors who offer context and examples of how technical problems relate to more fundamental deficiencies in the quality management system.

2 CE credits available

QMS Implementation Roadmap

Order ISOEDRM

Outlines the practical steps necessary to build, implement, and maintain a quality management system that meets the ISO 15189 standard. Gain perspective on practices (and pitfalls) straight from the CAP's ISO 15189 assessors, as well as ISO 15189-accredited laboratories. Designed for laboratory quality managers, plus your implementation team members.

2 CE credits available

Root Cause Analysis

Order ISOEDRC

Learn real-world methodology to conduct a root cause analysis, along with the tools necessary to implement it. Learn from actual examples of complete root cause analysis based on projects in laboratories like yours. You will even perform key steps based on a participant case study. The course is designed for laboratory quality managers and implementation team members.

6 CE credits available

Internal Auditing

Order ISOEDIA

Increase your capabilities for internal auditing with a proven methodology for process audits, tracer audits, and laser audits. Learn from the CAP's ISO 15189 assessors how to prepare for interviews, communicate findings to your quality management team, and use audits to drive process improvements.

3 CE credits available

Document Control

Order ISOEDDC

This "how-to" course on document control systems details how to control documents in a way that meets ISO 15189 requirements, how to accomplish document control even with minimal resources (such as spreadsheets), and how document control contributes to cost containment. All this from the CAP's ISO 15189 assessors who give examples and commentary on common pitfalls and issues.

2 CE credits available

Quality Manual Development

Order ISOEDQM

This course provides guidance on how to go beyond a quality plan to develop a manual that organizes and communicates your laboratory's quality management system. You will see an example of an effectively structured and written manual so you can organize and create your own. Plus, the CAP's ISO 15189 assessors show you approaches to link your quality policy to quality objectives and metrics.

2 CE credits available

Management Review

Order ISOEDMR

This course interprets the ISO 15189 requirements for management review. The CAP's ISO 15189 assessors discuss how to structure the review meeting, communicate results of quality assessments, and prompt strategic decisions from management—all in the context of the overall health of your organization.

2 CE credits available

Mistake Proofing

Order ISOEDMP

Increase your ability to design new processes, modify existing processes, minimize mistakes, and manage your risks. This course provides a methodology focused on five main categories of mistake-proofing tactics and shows examples of these tactics from the domain of laboratory medicine. It includes commentary by CAP member pathologists with experience using Lean and other process improvement techniques.

4 CE credits available

Quality Culture *Order ISOEDCL*



Designed for laboratory medical directors, administrative directors, quality managers, and other leaders who can affect the culture of their laboratory through their decisions and actions. The course provides an adaptable program for proactively shaping culture. It includes commentary by CAP member pathologists. *This course will be released in January 2018.*

2 CE credits available

Make sure your laboratory team is ready to meet the challenges ahead. Sign up now for this comprehensive set of learning tools

For more information, visit cap.org and search QMEd or call 800-323-4040 or 847-832-7000 option 1.

How current is your laboratory quality management plan?

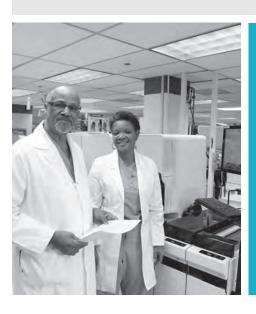


Created specifically for the needs of the anatomic pathology laboratory, this comprehensive manual can help you develop, implement, and maintain a comprehensive quality program. Learn valuable tips for designing your own laboratory quality plan that documents regulatory compliance. Text includes cross-references to the CAP's Laboratory Accreditation Program checklists, Joint Commission standards, and CLIA '88.

View sample pages and order online:

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Quality Management Tools



Engage in quality measures with our new Q-PROBES™ programs.

- Assess physician satisfaction with laboratory services to target quality improvement activities (QP181).
- Benchmark institutional turnover rates for laboratory staff and determine human resource practices for lower rates of turnover (QP182).
- Focus your laboratory education needs in its body fluid competency program (QP183).
- Examine your turnaround times on critical laboratory tests needed by the emergency department (QP184).

Quality Management Tools

Q-PROBES™Q-TRACKS®	
Clinical Pathology MonitorsAnatomic Pathology Monitors	31
Q-MONITORS®	
New Programs NEW	
Physician Satisfaction with Clinical Laboratory Services (QP181)	25
Physician Satisfaction with Clinical Laboratory Services (QP181)	26

Discontinued Programs

Utilization of Red Blood Cell Transfusions (QP171)
Workflow Process Mapping (QP172)
Phlebotomy Staffing Ratios (QP173)
Preanalytic Errors Competency (QP174)
Mislabeled Cases, Specimens, Blocks, and Slides in Surgical Pathology (QT19)

Quality Management Tools

Use the CAP's Quality Management Tools (QMT) to **improve the Total Testing Process** by identifying quality improvement opportunities of selected key processes in the clinical and anatomic pathology laboratories, examining preanalytical, analytical, and postanalytical phases:

- Establish realistic goals by comparing performance against similar institutions with comparable demographics
- Monitor progress through unique and robust quality indicators on a periodical basis
- Make effective quality management decisions based on practical and in-depth individual reports provided to participants
- Improve efficiencies to allow time for more patient-centric activities
- Easily integrate quality management into your daily work processes with predesigned monitoring tools developed by laboratory professionals and scientists

Q-PROBES™ A One-Time Opportunity to Perform In-Depth Quality Assessment

Q-TRACKS® A Program for Continuous Quality Monitoring

Q-MONITORS® Customized Quality Monitors Program

Q-PROBES, Q-TRACKS, and Q-MONITORS activities meet the American Board of Pathology MOC Part IV Practice Performance Assessment requirements.

Q-PROBES, Q-TRACKS, and Q-MONITORS

offer a comprehensive collection of tools to complement your quality management program needs.*

Select Q-PROBES, Q-TRACKS, and Q-MONITORS studies to support your quality improvement initiatives.	Preanalytic	Analytic	Postanalytic	Anatomic Pathology	Clinical Pathology	Turnaround Time	Patient Safety	Microbiology	Transfusion Medicine	Chemistry/ Hematology	Customer Satisfaction
Q-PROBES											
Physician Satisfaction with Clinical Laboratory Services (QP181) NEW	1				1						
Laboratory Staff Turnover (QP182) NEW											
Technical Competency Assessment of Body Fluid Slide Review (QP183) NEW		ı			1		ı				
Laboratory Result Turnaround Time for Emergency Room Specimens (QP184) NEW	•	•			•	•	•		•		•
Q-TRACKS							,				
Patient Identification Accuracy (QT1)											
Blood Culture Contamination (QT2)											
Laboratory Specimen Acceptability (QT3)											
In-Date Blood Product Wastage (QT4)											
Gynecologic Cytology Outcomes: Biopsy Correlation Performance (QT5)	1	ı		•			•				
Satisfaction With Outpatient Specimen Collection (QT7)	•										•
Stat Test Turnaround Time Outliers (QT8)		1									
Critical Values Reporting (QT10)			•								
Turnaround Time of Troponin (QT15)		ı								1	
Corrected Results (QT16)			•		ı						•
Outpatient Order Entry Errors (QT17)										•	
Q-MONITORS											
Monitoring of Troponin Metrics for Suspected MI (QM1)	ı	ı									

^{*}The CAP requires accredited laboratories to have a quality management plan that covers all areas of the laboratory and includes benchmarking key measures of laboratory performance (GEN.13806, GEN.20316, COM.04000). The Joint Commission requires accredited hospitals to regularly collect and analyze performance data (PI.01.01.01, PI.02.01.01). CLIA requires laboratories to monitor, assess, and correct problems identified in preanalytic, analytic, and postanalytic systems (§493.1249, §493.1289, §493.1299).

Q-PROBES

A One-Time Opportunity to Perform In-Depth Quality Assessment

Implement quality monitoring—Use Q-PROBES short-term comprehensive quality studies¹ to learn how to start monitoring and measuring key processes that you may not have followed in the past or that are not commonly monitored in most laboratories. Q-PROBES studies analyze hot topics and industry trends to keep the laboratory current.

Gain experience in data collection and analysis—Participants will collect data during predetermined dates. Based on submitted data, the CAP provides personalized reports with the individual participant's performance compared against other participants.

Strengthen your quality assessment expertise—The CAP's pathologist experts provide in-depth discussion and identify best practices for laboratories to strive for. In addition, consolidated results of the studies are carefully reviewed and analyzed to be published in the form of scientific articles. Such articles give participants an extra layer of information to be utilized for further analysis.

Participants in the Q-PROBES program receive:

- User guide
- Templates and instructions for data collection
- Individual report, how to interpret the results guide, overall aggregated data
- Data Analysis and Critique that includes data distributions and analysis of laboratory practices and commentaries from pathologist experts on improvement opportunities

Quanty mana			aiviauai	Report o	n nesuits			
Performance Indicator	Your Result	10th	utions Per 50th (vertical line)	centiles 90th (right edge of box)	Perfo	mance Dis	tribution	
ABO mislabeled specimen rate (per 1000 specimens) (n=30)	3.68	0.00	4.04	18.19		10	15]
Performance note: The bar graph ranges from th Lower percentiles (shaded an					epresents the n			_
			etter relativ		epresents the n			
	ea and lowe	r) represent b	All Inst	ve performan	epresents the noce.			
Additional Information ABO typing result discrepancy rate (per 1000	ea and lowe	Your Result	All Inst	titutions Pe	epresents the noce.			

Access to the scientific articles that are published with the results of the studies

Q-PROBES activities meet the American Board of Pathology MOC Part IV Practice Performance Assessment requirements.

¹ Q-PROBES studies are available only one time annually and may not be repeated in the future.



Physician Satisfaction with Clinical Laboratory Services QP181

Introduction

Assessing physician satisfaction with laboratory services provides valuable information for targeting quality improvement activities. The CAP's Laboratory Accreditation Program requires that institutions measure customer satisfaction (eg, physicians, patients, nurses) with laboratory services. This Q-PROBES study will assist your organization in meeting these requirements while helping to identify areas for improvement, and furthering understanding of client needs to ensure future physician satisfaction with your services. Enrollment in QP181 can help meet College of American Pathologist (Laboratory General Checklist Statements GEN.20316, GEN.20335) and Joint Commission requirements for laboratory accreditation.

Objectives

This study will assess physician satisfaction with laboratory services and correlate this with laboratory workload, performance improvement activities, and customer support services.

Data Collection

Clinicians will be asked to complete a satisfaction survey regarding their experience across various clinical laboratory service categories, including turnaround time, critical value notification, diagnostic accuracy, communication, accessibility, responsiveness, and courtesy. They will also indicate whether they would recommend the laboratory to another physician.

The surveys will be available in two formats: electronic distribution with direct survey submission to the CAP, or paper response forms. The CAP can accept up to 50 paper surveys, but the number of electronic submissions is unlimited.

Performance Indicators

- · Overall mean satisfaction score with clinical laboratory services
- · Mean satisfaction scores for the specific service categories
- Laboratory recommendation rate (%)

This is a one-time study conducted in the first quarter.



Laboratory Staff Turnover QP182

Introduction

The national vacancy rate for medical technologists is 10.4%. The turnover rate for all laboratory employees is not known. Laboratory medical directors are keenly interested in laboratory staff turnover rates since licensing and accreditation agencies hold them responsible for ensuring that clinical laboratories are adequately staffed. Hospital and laboratory administrators share that interest, as their boards of trustees hold them responsible for the successful operation and financial solvency of their institutions.

Objective

This study will measure national institutional turnover rates for laboratory staff and determine human resources practices associated with lower rates.

Data Collection

Laboratories will provide data on full-time equivalent (FTE) employees, job status changes, and job vacancies. From this data, turnover rates for several classes of laboratory workers will be calculated. Laboratories will be asked to complete a detailed questionnaire concerning human resources practices in order to identify characteristics that are associated with lower and higher turnover rates.

Performance Indicators

- Overall laboratory employee turnover rate
- · Turnover rate by personnel category
- · Vacancy rate

This is a one-time study conducted in the second quarter.

NEW

Technical Competency Assessment of Body Fluid Slide Review QP183

Introduction

Technologists receive a variety of body fluids for examination in the laboratory and must maintain their identification skills of these specimens. Laboratories are required to provide education and assess competency in this area on a regular basis.

Participation in this Q-PROBES study helps laboratories meet CLIA personnel requirements (Subpart M, 42 CFR §493.1), the CAP Laboratory Accreditation Program Checklist Statement requirement (GEN.55500 Competency Assessment of Testing Personnel), and Joint Commission HR.01.06.01 for competency assessment of nonwaived testing personnel.

Objective

This study will assess the effectiveness of educational and practical experience policies and procedures dedicated to the laboratory's efforts in maintaining technologist skills in the performance of accurate body fluid cell counts and identification of other body fluid features. The individual competency assessment will be performed using whole slide images that were evaluated by hematology experts. Results of this study will assist individuals, the laboratory director, and manager with areas to focus on for improvement.

Online whole slide images are powered by DigitalScope technology. See system requirements on page 13.

Data Collection

Information will be collected from each site regarding minimum qualifications and experience requirements of their technologists, their ongoing educational programs and requirements, as well as relevant procedures and policies.

A series of whole slide images will be available online to each participating institution to assess their technologists' ability to perform a cell differential on Wright Stained body fluid and identify miscellaneous cells and inclusions in cytocentrifuged preparations. Each program ordered will include input forms for use by up to 10 technologists. Laboratories that need forms for more than 10 individuals should order one additional program for each 10 additional technologists.

Participants will provide additional information about their competency assessment programs, continuing education, and professional background.

Reports

Laboratory Summary

- Institution performance shown by case and by technologist
- Institution performance compared to all institutions
- Overall laboratory performance based on the facility's individual technologist performance(s)

Technologist Summaries

- Individual technologist performance for each case and an overall performance
- Technologist's ability to identify various WBC types, red blood cells, and other items present in normal and abnormal cases in comparison to consensus responses

This is a one-time study conducted in the third quarter.



Laboratory Result Turnaround Time for Emergency Room Specimens QP184

Introduction

Fast delivery of test results is important for patient care in the emergency department (ED). Therefore, monitoring turnaround time (TAT) is an important quality measure. Timeliness of reporting results for high volume, automated tests such as serum potassium or troponin is commonly used as the key performance indicator. However, other equally important tests may inherently take longer to complete due to processing or analysis requirements. Factors involved in TATs could account for bottlenecks in diagnosis, treatment, or patient flow if impacted by any of these tests.

Objective

The objective of this study is to provide a more complete assessment of TAT testing for emergency room care by examining a representative variety of critically important tests from various functional areas of the laboratory. Tests to be studied include those that may be critical for patient care or subsequent workup, and include blood type and screen, D-dimer, influenza A virus, microscopic urine leukocyte count, serum potassium, urine drug screen, and urine pregnancy. In addition, the study will examine operational and administrative factors which may influence performance.

Data Collection

Laboratories will retrospectively record accession and report times for each designated test over a variety of days of the week and shifts.

Laboratories will provide their expected TAT goals, if available, for each test examined in the study.

Laboratories will complete a survey about their ED STAT test menu.

Performance Indicators

- · Accession to reporting TAT for STAT ordered ED tests
- Compliance rate with expected TAT goals for STAT ordered ED tests

This is a one-time study conducted in the fourth quarter.

Q-TRACKS

A Program for Continuous Quality Monitoring

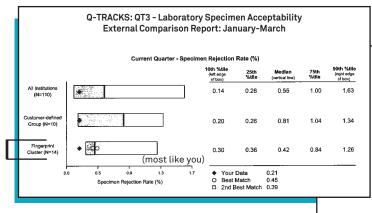
Identify and monitor opportunities for quality improvement over time

Use established Q-TRACKS programs to identify opportunities to quantitate your quality improvement measures.

Evaluate quality improvements

Measure the effectiveness and impact of implemented changes in key processes. The individual reports include performance of quality indicators over time, benchmarking information, trends, and suggested areas for improvement.

Step 1: Establish realistic benchmarks by comparing your laboratory to others like yours.



Step 2: Identify improvement opportunities.

Specimen Rejection Reasons	Your Data (%)	Aggregate Percent*
Specimen lost/not received	0.0	12.1
Unlabeled specimen	6.4	2.2
Mislabeled specimen	4.5	3.0
Incompletely labeled specimen or Inadequately filled-out form	0.0	1.6
Specimen hemolyzed	40.0	29.3
Specimen clotted	29.1	17.9
Insufficient specimen quantity	16.4	15.1
Unacceptable variance (delta check)	0.0	3.1
Wrong container	3.6	2.5
Wrong temperature	0.0	0.4
Other reason	0.0	12.7

Q-TRACKS: QT3 - Laboratory Specimen Acceptability
Trend Analysis Report: January-March

Specimen Rejection Rate (%)

Step 3:

Monitor improvement over time to ensure accurate results, patient safety, and quality patient care.

Participants in the Q-TRACKS program receive:

- User Guide
- · Templates and instructions for data collection
- Quarterly reports that include: fingerprint clusters, customer-defined groups, and all institution comparisons
- · Peer directory

Q-TRACKS activities meet the American Board of Pathology MOC Part IV Practice Performance Assessment requirements.

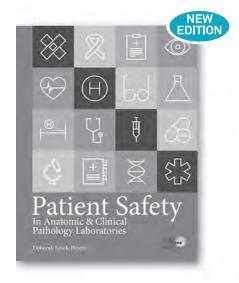
Enhance the culture of patient safety in your laboratory

Patient Safety in Anatomic & Clinical Pathology Laboratories enables you to connect the patient safety culture in your laboratory to the overall mission and goals of your health care enterprise.

- Prevent errors in communication, handoffs, and transitions
- Use technology to improve laboratory patient safety
- · Learn how cognitive bias can contribute to patient safety errors
- Build high-reliability teams
- Engage the patient navigator to address safety issues through continuity and coordination of care
- Develop and implement a patient safety curriculum for the laboratory
- Understand how new accreditation milestones advance patient safety initiatives

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Item number: PUB316 Softcover; 128 pages; 2017

Q-TRACKS Clinical Pathology Monitors

Patient Identification Accuracy QT1

In order to report accurate laboratory results and meet The Joint Commission National Patient Safety Goal #1: "Identify patients correctly," institutions must properly identify patients. Since most laboratories perform testing away from the patient, patient identification, and labeling of specimens and coordination with test requisitions must be performed accurately and completely. By continuously monitoring for wristband errors, participants can promptly identify and correct problems that may interfere with patient care services. Use this monitor to help meet CAP General Checklist statements GEN.20316, GEN 40490, and GEN.40825.

Objectives

Assess the incidence of wristband errors within individual institutions, compare performance between participating institutions, and identify improvement opportunities.

Data Collection

On six predetermined days per month, participants will monitor patient wristband identification for all phlebotomies performed at their institution. Phlebotomists will tally the total number of wristbands checked, the number of errors found, and the types of wristband errors. This monitor includes all routinely wristbanded patients. Include emergency department patients only if the emergency department routinely applies wristbands to these patients.

Performance Indicator

Wristband error rate (%)

Performance Breakdown

• Breakdown of wristband error types (%)

Blood Culture Contamination QT2

Despite advances in blood culture practices and technology, false-positive blood culture results due to contaminants continue to be a critical problem. Blood culture contamination rate, the primary indicator of preanalytic performance in microbiology, is associated with increased length of hospital stay, additional expense, and the administration of unnecessary antibiotics. The CAP and other accrediting organizations require you to monitor and evaluate key indicators of quality for improvement opportunities. Use this monitor to help meet CAP Checklist statement note MIC.22630: "It is recommended that blood culture statistics, including number of contaminated cultures, be maintained and reviewed regularly by the laboratory director. The laboratory should establish a threshold for an acceptable rate of contamination. Tracking the contamination rate and providing feedback to phlebotomists or other persons drawing cultures has been shown to reduce contamination rates."

Objective

Determine the rate of blood culture contamination using standardized criteria for classifying contaminants.

Data Collection

On a monthly basis, participants will tabulate the total number of blood cultures processed and the total number of contaminated blood cultures. Blood cultures from neonatal patients are tabulated separately. For the purposes of this study, participants will consider a blood culture to be contaminated if they find one or more of the following organisms in only one of a series of blood culture specimens: Coagulase-negative *Staphylococcus*; *Micrococcus*; Alpha-hemolytic viridans group streptococci; *Propionibacterium acnes*; *Corynebacterium* sp. (diptheroids); or *Bacillus* sp. Participants have the option to monitor institution-specific subgroups, for example, a specific department or patient population.

Performance Indicators

- Neonatal contamination rate (%)
- Other contamination rate (%)
- Overall contamination rate (%)

Laboratory Specimen Acceptability QT3

A substantial amount of rework, diagnostic and therapeutic delay, and patient inconvenience can result from specimen rejection. Patient redraws may result from unlabeled, mislabeled, and incompletely labeled specimens; clotted and/or hemolyzed specimens; or insufficient specimen quantity. By continuously monitoring specimen acceptability, collection, and transport, laboratories can promptly identify and correct problems. Enrollment in this Q-TRACK may assist the laboratory in monitoring compliance with Laboratory General Checklist statement GEN.40825: "There is a system to positively identify all patient specimens, specimen types, and aliquots at all times."

Objective

Identify and characterize unacceptable blood specimens that are submitted to the chemistry and hematology/coagulation sections of the clinical laboratory for testing.

Data Collection

This monitor includes all blood specimens submitted for testing to the chemistry and hematology departments of the clinical laboratory. On a weekly basis, participants will record the total number of specimens received, the number of rejected specimens, and the primary reason each specimen was rejected.

Performance Indicator

• Specimen rejection rate (%)

Performance Breakdown

• Breakdown of reasons for rejection (%)

In-Date Blood Product Wastage QT4

Blood for transfusion is a precious resource. At a minimum, wastage of blood that is not out-of-date represents a financial loss to the health care system. More ominously, systemic wastage of blood may reflect an environment of care that is out of control and could pose risks to patient safety.

Objective

Compare the rates of blood product wastage (ie, units discarded in-date) in participating hospitals and track rates of improvement over time.

Data Collection

On a monthly basis, participants will use blood bank records to obtain information on the total number of units transfused for each type of blood component. Participants will track the number and type of blood units that are wasted in-date and the circumstances of wastage. This monitor includes the following types of blood components: red blood cells (allogeneic), frozen plasma, platelet concentrates, single donor platelets, and cryoprecipitate.

Performance Indicators

- Overall blood wastage rate (%)
- Wastage rates by blood component type (%)

Performance Breakdown

• Breakdown of circumstances of wastage (%)

Satisfaction With Outpatient Specimen Collection QT7

Specimen collection is one of the few areas of laboratory medicine that involves direct outpatient contact. As a result, patient satisfaction with this service is a vital indicator of quality laboratory performance. The CAP's Laboratory Accreditation Program requires measurement of patient satisfaction with laboratory services (GEN.20335). Use this monitor to help meet this requirement.

Objective

Assess patient satisfaction with outpatient phlebotomy services by measuring patients' assessments of waiting time, discomfort level, courteous treatment, and overall satisfaction.

Data Collection

On a monthly basis, participants will provide copies of a standardized questionnaire to a minimum of 25 outpatients (maximum of 99 outpatients) using predetermined data collection criteria. This monitor includes any outpatient undergoing venipuncture. This monitor excludes patients seen in the emergency department, ambulatory surgery area, urgent care facility, chest pain center, 23-hour short-stay facility, employee health department, outpatient health screening fair/promotion, dialysis center, nursing home, or extended care facility.

Performance Indicators

- Satisfaction scores and satisfaction rates (% of patients rating 4 or 5) for the following categories:
 - o Overall experience

o Courtesy

o Waiting time

o Patient privacy

o Patient comfort

o Laboratory hours of operation

Stat Test Turnaround Time Outliers QT8

The stat test turnaround time (TAT) outlier rate, expressed as a percentage of tests missing target reporting times, is a measure of outcomes that evaluates how well the laboratory meets patient and clinician needs. This monitor helps meet CAP Checklist requirement GEN.20316: "The QM program includes monitoring key indicators of quality in the preanalytic, analytic, and postanalytic phases."

Objective

Monitor the frequency that stat test TAT intervals exceed institutional stat test TAT expectations.

Data Collection

Before beginning data collection, participants will establish a specimen receipt-to-report deadline for emergency department (ED) stat potassium tests. On six predetermined days per month, participants will monitor the TAT of up to 10 randomly selected ED stat potassium tests on each of three, eight-hour shifts (up to 180 tests per month) and track the number of ED stat potassium results reported later than the established reporting deadline. This monitor includes stat potassium tests ordered as part of a panel and excludes stat potassium levels that are requested on body fluids other than blood, as part of timed or protocol studies, or after the specimen arrives in the laboratory.

Performance Indicator

• Stat test TAT outlier rate (%)

Performance Breakdowns

- Breakdown of outliers by shift (%)
- Breakdown of outliers by day of week (%)

Critical Values Reporting QT10

Laboratories commonly refer to critical values as results requiring immediate notification to the physician or caregiver for necessary patient evaluation or treatment. Regulations from agencies and accreditors such as the CMS, The Joint Commission, and the CAP (GEN.20316, COM.30000) mandate that laboratories develop and implement an alert system for critical values. Use this monitor to document compliance with your laboratory's alert plan.

Objective

Evaluate the documentation of successful critical values reporting in the general laboratory for inpatients and outpatients.

Data Collection

On a monthly basis, participants will evaluate 120 inpatient and 120 outpatient critical values. Data collection will include general chemistry, hematology, and coagulation analytes on the critical values list. Retrospectively, participants will record the total number of critical values monitored and the number with documentation of successful notification. In addition, participants will provide the number of critical values that were not communicated within three hours, the number of failed notifications due to laboratory oversight, and the number of successful notifications to licensed caregivers. This monitor will exclude critical values for cardiac markers, drugs of abuse, therapeutic drug levels, urinalysis, blood gases, point-of-care tests, and tests performed at reference laboratories.

Performance Indicators

- Total critical values reporting rate (%)
- Inpatient critical values reporting rate (%)
- Outpatient critical values reporting rate (%)
- Failed notification (<3 hours) rate (%)

Turnaround Time of Troponin QT15

The swiftness with which physicians establish diagnoses of acute myocardial infarction (AMI) in patients presenting to the emergency department (ED) with chest pain may determine the type and predict the outcome of therapy those patients will receive. Included in the total time consumed in establishing diagnoses of AMI are the component intervals required to measure biochemical markers of myocardial injury. One of the most critical biochemical markers is troponin. Use this monitor to help meet CAP Checklist requirement GEN.20316 QM Indicators of Quality.

Objective

Determine the median order-to-report turnaround time (TAT) of troponin (I or T) ordered to rule out myocardial infarction and the percent of troponin results reported by each institution's established deadline.

Data Collection

On six predetermined days per month, participants will record TATs (in minutes) for three randomly selected troponin specimens obtained from ED patients on each of three traditional shifts, a total of nine measurements. Participants will measure TATs from the time of test order to the time results are available to ED personnel.

Performance Indicators

- · Median troponin order-to-report TAT (minutes)
- Troponin TAT compliance rate (%)

Corrected Results QT16

The CAP developed this Q-TRACKS monitor in recognition of the importance of timely detection and correction of erroneous laboratory results. Accuracy in laboratory results is critical to the effectiveness of a physician's plan of care for a patient. An erroneous result can delay or alter patient treatment; therefore, detection of erroneous results should be a priority in every laboratory and should be monitored as a key quality indicator. Help measure your compliance with CLIA 493.1299, Postanalytic Systems Quality Assessment, and help meet CAP Checklist statement GEN.20316 with this monitor.

Objective

Monitor the number of corrected test results within individual institutions and compare performance with that of all institutions and those institutions similar to yours.

Data Collection

On a monthly basis, participants will monitor the number of corrected test results and the total number of billable tests for that month. Include test results for all patients in all care settings with the following exclusions: anatomic pathology tests, narrative physician-interpreted tests (eg, bone marrow biopsies and peripheral smear reports), and point-of-care tests.

Performance Indicator

• Test result correction rate (per 10,000 billable tests)

Outpatient Order Entry Errors QT17

Order accuracy bears an obvious relationship to the quality of laboratory testing. When the laboratory fails to complete a requested test, it delays the diagnostic evaluation, potentially extending a patient's hospital stay and prolonging therapy. When the laboratory completes a test that was not requested, the cost of care increases, patients may be subjected to unnecessary phlebotomy, and laboratory efficiency declines. Use this monitor to help meet CAP Checklist statement GEN.20136 for test order accuracy.

Objective

Measure the incidence of incorrectly interpreted and entered outpatient physician test orders into the laboratory information system, compare performance across institutions, and track performance over time.

Data Collection

On six preselected weekdays per month, participants will compare eight outpatient requisitions or order sheets to the orders entered into the laboratory's information system to determine if any order entry errors occurred.

This monitor includes test order review from ambulatory outpatients seen in offices and clinics operated by your laboratory services, private physician offices, nursing homes, extended care facilities, and free-standing phlebotomy areas. Also included are send-out tests, chemistry, hematology, microbiology, immunology, toxicology, and urinalysis tests on outpatients. Order entry error categories include requesting physician errors; incorrect, missing, and extra test errors; test priority errors; and copy or fax result errors.

This monitor excludes tests performed in transfusion medicine or anatomic pathology and also excludes tests from the following patient care settings: inpatient, emergency department, ambulatory surgery, urgent care, chest pain center, 23-hour short-stay facility, employee health department, outpatient screening fair/promotion, and dialysis center.

Performance Indicators

- Overall outpatient order entry error rate (%)
- Order entry error rates by type (%)

Performance Breakdown

Breakdown of error types (%)

Q-TRACKS Anatomic Pathology Monitors

Gynecologic Cytology Outcomes: Biopsy Correlation Performance QT5

The correlation of cervicovaginal cytology (Pap test) findings with cervical biopsy results is a significant part of the cytopathology laboratory's quality assurance program. By monitoring this correlation, the laboratory can identify and address potential problems requiring improvement, thereby ensuring better patient results.

Objective

Quantify the correlation between the findings of cervicovaginal cytology and corresponding histologic material.

Data Collection

On a monthly basis, participants will record the number of true-positive, false-positive, and false-negative cytology-biopsy correlations. The false-negative correlations will be classified into four error categories: screening errors, interpretive errors, screening and interpretive errors, and adequacy determination errors. Participants will also record the biopsy diagnoses for Pap tests with an interpretation of atypical squamous cells (ASC-US and ASC-H) or atypical glandular cells (AGC). This monitor includes cervical biopsy specimens submitted to the laboratory that have a corresponding satisfactory or satisfactory but limited Pap test within three months of the biopsy.

Performance Indicators

- Predictive value of positive cytology (%)
- Sensitivity (%)
- Screening/interpretation sensitivity (%)
- Sampling sensitivity (%)
- · Percent positive for ASC-US interpretations
- · Percent positive for ASC-H interpretations
- Percent positive for AGC interpretations

Q-MONITORS

A Program for a Customized Comprehensive Assessment

Evaluate quality improvements in your laboratory

With today's focus on reducing medical errors, achieving and maintaining excellence is key to success. Using continuous monitoring, Q-MONITORS provide a comprehensive assessment of key processes in your institution.

Structure your data collection and analysis for success

Use Q-MONITORS to help build and improve data collection and analyze processes that contribute to quality of care, patient safety, and outcomes. Observe performance trends over time to identify and monitor opportunities for quality improvement through quantitative quality measures.

Establish realistic laboratory benchmarks and performance goals

Q-MONITORS offer customized programs that address process-, outcome-, and structureoriented quality assurance issues. Establish benchmarks through external database comparisons and compare your performance to establish goals for performance improvement.

Q-MONITORS Customized Quality Monitoring Program

Monitoring of Troponin Metrics for Suspected MI QM1

Patients presenting to the emergency department (ED) with chest pain must be evaluated quickly. Rapid serum troponin measurement is an important part of ED practice that can provide decisive information for patient management. Reducing delays in troponin testing has been reported to result in shorter length of stay in the ED and more rapid initiation of anti-ischemic treatment. Emergency departments and chest pain centers should, therefore, have effective procedures for ensuring optimal turnaround time (TAT) for troponin and a process for ongoing monitoring to ensure that performance meets expectations.

Objective

Determine and monitor troponin TATs for patient arrival to result availability and/or up to six time intervals within the total testing process for patients presenting to the ED with chest pain.

Data Collection

Six days per month, collect data from nine patients presenting to the ED with chest pain and tested for troponin level. Data includes time of patient arrival, troponin test order, specimen collection, laboratory receipt, and result availability. It is not necessary to provide data from each TAT component. Participants select which TAT metrics to monitor, with the option to monitor all metrics.

Participants will also complete a questionnaire about clinical and laboratory practices related to troponin testing.

Metrics

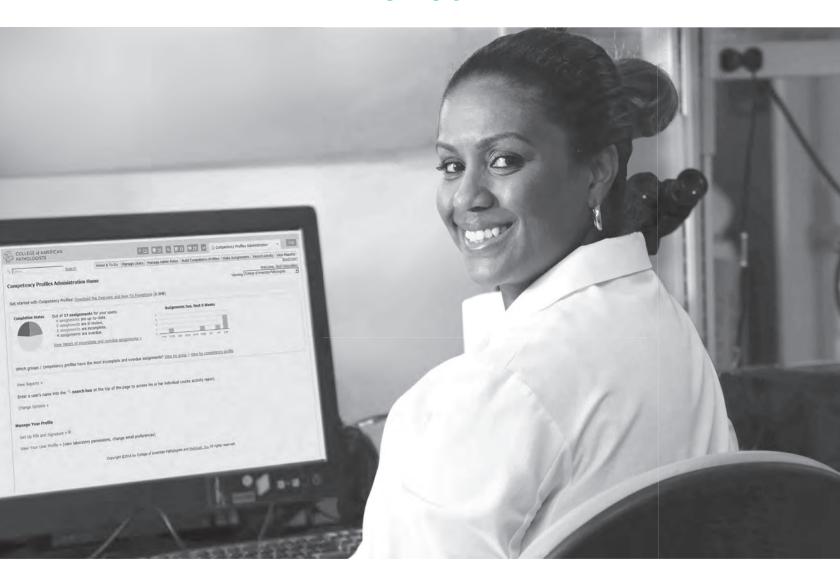
Depending on the data submitted, the following metrics will be provided. In addition, TAT benchmarking, as compared to all institutions, will be provided for both point-of-care and clinical laboratory testing for patient arrival to result availability and specimen collection to result availability.

- · Patient arrival to result availability
- · Specimen collection to result availability
- · Test order to result availability
- · Patient arrival to test order
- · Test order to specimen collection
- · Specimen collection to laboratory receipt
- · Laboratory receipt to result availability

Performance Indicators

- Median TAT for troponin testing intervals (monthly)
- Test order to result availability compliance rate (if applicable)
- Specimen collection to result availability compliance rate (if applicable)

If it's not documented, it's not compliant. Period.



Documenting the competency assessment of your staff is the #1 deficiency cited by major laboratory accreditors. It's true—one in four laboratories do not fully meet the documentation requirements of competency assessment.

You may know that your team follows all CLIA regulations to the letter. But when inspection time comes—if it's not documented, it's considered a deficiency. Learn how to align your CLIA competency assessment plan with the quality assurance processes you already perform regularly.

Discover how the CAP's Competency Assessment Program can improve your laboratory's readiness for inspection.

Learn more at cap.org and search for Competency Assessment or email competency@cap.org.

4

Quality Cross Check



Simplify biannual instrument comparability studies with Quality Cross Check.

- Receive custom reports with peer group evaluations and instrument comparability statistics.
- Monitor up to 30 glucose meters with Quality Cross-Check—Whole Blood Glucose program (WBGQ).



Perform instrument comparability and stay in compliance

Quality Cross Check is a convenient solution to monitor instrument performance and assess comparability across multiple instruments in your laboratory and to identify potential issues before they affect patient results.

How It Works

- Receive three challenges in each of two mailings a year.
- Report up to three instruments for each challenge (and report up to 30 instruments for Quality Cross Check—Whole Blood Glucose).
- Receive a custom report package that includes peer group comparison and instrument comparability statistics for each reported analyte.

Stay in Compliance

In August 2015, the Centers for Medicare & Medicaid Services (CMS) reiterated that laboratories are not permitted to test proficiency testing samples on multiple instruments unless that is how the laboratory tests patient specimens.

The CMS interpretation was expanded beyond regulated analytes to include analytes not listed in Subpart I of the Clinical Laboratory Improvement Amendments regulations, including waived methods.

Quality Cross Check complements your existing CAP Surveys to monitor multiple instrument performance and is compliant with the CMS directive.

Monitoring Performance of Glucose Meters

Beginning in 2017, PT for waived whole blood glucose on glucose meters is no longer required for laboratories accredited by the CAP. Laboratories are required to perform alternative performance assessment.

In response to this change, the CAP introduced the Quality Cross Check—Whole Blood Glucose program (WBGQ). Participants in this program will enjoy the benefits of Quality Cross Check and have the ability to report up to 30 instruments for each challenge.

General Chemistry and Therapeutic Drug Monitoring

Quality Cross Check—Chemistry and Therapeutic Drug Monitoring CZQ					
Analyte	Program Code Challenges/Shipment				
	CZQ				
See Survey CZ analytes on pages 56-58	ı	3			

This program does not meet regulatory requirements for proficiency testing; see Survey CZ on pages 56-58. For additional information about the CAP Quality Cross Check program, see page 42.

Quality Cross Check—BNP BNPQ						
Analyte	Program Code Challenges/Shipment					
	BNPQ					
BNP	I	3				
NT-proBNP	I	3				

This program does not meet regulatory requirements for proficiency testing; see Survey BNP or BNP5 on page 61. For additional information about the CAP Quality Cross Check program, see page 42.

Quality Cross Check—Whole Blood Glucose WBGQ					
Analyte	Program Code Challenges/Shipment				
	WBGQ				
Glucose	■ 3				

The CAP Accreditation Program requires all accredited laboratories performing waived whole blood glucose testing using glucose meters to perform alternative performance assessment. This program can be used to meet alternative performance assessment requirements.

Program Information

- Three 5.0-mL liquid serum specimens in duplicate
- · Report up to three instruments
- · Conventional and International System of Units (SI) reporting offered
- Two shipments per year

Program Information

- Three 1.5-mL liquid specimens
- · Report up to three instruments
- · Conventional and International System of Units (SI) reporting offered
- · Two shipments per year

- Three 2.0-mL whole blood specimens
- Report up to 30 instruments
- Two shipments per year



Quality Cross Check—Body Fluid Chemistry FLDQ Analyte **Program Code** Challenges/Shipment **FLDQ** 3 Albumin 3 Amylase 1 CA19-9 1 Carcinoembryonic antigen (CEA) 3 Cholesterol 3 Creatinine 3 Glucose 3 Lactate 3 Lactate dehydrogenase (LD) 3 рΗ Protein, total 3 Triglycerides 3 Urea nitrogen

This program does not meet regulatory requirements for proficiency testing; see Survey FLD on page 72. For additional information about the CAP Quality Cross Check program, see page 42.

Quality Cross Check—Hemoglobin A _{1c} GHQ				
Analyte	Program Code Challenges/Shipme			
	GHQ			
Hemoglobin A _{1c}		3		

This program does not meet regulatory requirements for proficiency testing; see Survey GH5 on page 63. For additional information about the CAP Quality Cross Check program, see page 42.

Program Information

- Three 3.0-mL specimens in duplicate
- · Report up to three instruments
- Two shipments per year

- · Three 0.8-mL liquid specimens in triplicate
- · Report up to three instruments
- Two shipments per year

Endocrinology

Quality Cross Check-	-Parathyroid Ho	rmone PTHQ
Analyte	Program Code	Challenges/Shipment
	PTHQ	
Parathyroid hormone (PTH)	ı	3

This program does not meet regulatory requirements for proficiency testing; see Survey ING on page 86. For additional information about the CAP Quality Cross Check program, see page 42.

Program Information

- Three 5.0-mL lyophilized serum specimens in duplicate
- Report up to three instruments
- Two shipments per year

World-class recognition deserves to be displayed.



Let your peers, patients, and the public know you've earned the CAP accreditation certification mark.

Proudly display the mark. It distinguishes you as one of almost 8,000 laboratories worldwide that have attained CAP accreditation, the most respected and recognized laboratory accreditation in the world.

Blood Gas, Critical Care, and Oximetry

Quality Cross Check—Blood Oximetry SOQ				
Analyte	Program Code	Challenges/Shipment		
	SOQ			
Carboxyhemoglobin	I	3		
Hematocrit, estimated		3		
Hemoglobin, total		3		
Methemoglobin		3		
Oxyhemoglobin		3		

This program does not meet regulatory requirements for proficency testing; see Survey SO on page 94. For additional information about the CAP Quality Cross Check program, see page 42.

Program Information

- Three 1.2-mL liquid specimens in triplicate
- Report up to three instruments
- Conventional and International System of Units (SI) reporting offered
- Two shipments per year

Quality Cross Check—Blood Gas AQQ, AQ2Q, AQ3Q, AQ4Q					
Analyte		Progra	m Code		Challenges/Shipment
	AQQ	AQ2Q	AQ3Q	AQ4Q	
Calcium, ionized		•			3
Chloride		ı			3
Hematocrit		1			3
Hemoglobin, estimated		1			3
Lactate		1			3
Magnesium, ionized		1			3
PCO ₂		1			3
рН		ı			3
PO ₂		1			3
Potassium		1			3
Sodium		1			3
tCO ₂		1		•	3
Creatinine		1			3
Glucose		1			3
Urea nitrogen (BUN)		1		•	3

These programs do not meet regulatory requirements for proficiency testing; see Surveys AQ and AQ2-AQ4 on page 92. For additional information about the CAP Quality Cross Check program, see page 42.

- AQQ, AQ2Q Three 2.5-mL specimens in triplicate and three 2.5-mL specimens for hematocrit testing in triplicate; appropriate for all methods except i-STAT®
- AQ3Q, AQ4Q Three
 1.7-mL specimens in triplicate for i-STAT methods only
- Report up to three instruments
- Conventional and International System of Units (SI) reporting offered
- Two shipments per year

Hematology and Clinical Microscopy

Quality Cross Check—Hematology Series FH3Q, FH4Q, FH6Q, FH9Q					
Analyte/Procedure	Program Code Challenges/ Shipment				
	FH3Q	FH4Q	FH6Q	FH9Q	
Hematocrit		ı	ı		3
Hemoglobin	•	ı	ı		3
Immature granulocyte parameter					3
Large unstained cells (LUC)		ı			3
MCV, MCH, MCHC	•	ı	ı		3
MPV	•	ı	ı		3
Nucleated red blood cell count (nRBC)					3
Platelet count		ı	ı		3
RDW	•	ı	ı		3
Red blood cell count	•	ı	ı		3
White blood cell count	•	ı			3
WBC differential	•	ı	ı		3

These programs do not meet regulatory requirements for proficiency testing; see the FH Series on page 135. For additional information about the CAP Quality Cross Check program, see page 42.

Quality Cross Check—Reticulocyte RTQ, RT2Q, RT3Q, RT4Q					NEW
Instrument/Method		Program Code			Challenges/ Shipment
	RTQ	RT2Q	RT3Q	RT4Q	
Abbott Cell-Dyn 4000, Sapphire, Siemens ADVIA 120/2120, and all other automated and manual methods	•				3
Abbott Cell-Dyn 3200, 3500, 3700, Ruby					3
Coulter GenS, HmX, LH500, LH700 series, MAXM, STKS, Unicel DxH			•		3
Sysmex XE-2100, XE-2100C, XE-5000, XN Series, XT-2000i, XT-4000i				ı	3

These programs do not meet regulatory requirements for proficiency testing; see the RT Series on page 139. For additional information about the CAP Quality Cross Check program, see page 42.

Program Information

- Three 2.5-mL whole blood specimens with pierceable caps
- · Report up to three instruments
- · For method compatibility, see instrument matrix on page 137
- Two shipments per year

- RTQ, RT2Q Three 1.0-mL stabilized red blood cell specimens
- RT3Q Three 3.0-mL stabilized red blood cell specimens
- RT4Q Three 2.0-mL stabilized red blood cell specimens
- · Includes percentage and absolute result reporting
- · Report up to three instruments
- Two shipments per year

Quality Cross Check—Urinalysis CMQ				
Analyte	Program Code Challenges/Shipn			
	CMQ			
Bilirubin		3		
Blood or hemoglobin		3		
Glucose	ı	3		
hCG urine, qualitative		3		
Ketones		3		
Leukocyte esterase	I	3		
Nitrite		3		
Osmolality		3		
рН	I	3		
Protein, qualitative		3		
Reducing substances		3		
Specific gravity		3		
Urobilinogen		3		

This program does not meet regulatory requirements for proficiency testing; see Surveys CMP and CMP1 on page 144. For additional information about the CAP Quality Cross Check program, see page 42.

Quality Cross Check—Occult Blood OCBQ					
Analyte	Program Code Challenges/Shipment				
	OCBQ				
Occult blood	I	3			

This program does not meet regulatory requirements for proficiency testing; see Survey OCB on page 149. For additional information about the CAP Quality Cross Check program, see page 42.

Program Information

- Three 10.0-mL liquid urine specimens for use with all instruments
- · Report up to three instruments
- Two shipments per year

- Three 2.0-mL simulated fecal specimens
- · Report up to three instruments
- Two shipments per year

Coagulation

Quality Cross Check—Coagulation CGLQ				
Analyte	Challenges/ Shipment			
	CGLQ			
Activated partial thromboplastin time	I	3		
Fibrinogen	I	3		
International normalized ratio (INR)	I	3		
Prothrombin time	I	3		
D-dimer	I	1		
Fibrin(ogen) degradation products, plasma	ı	1		
Fibrin(ogen) degradation products, serum	ı	1		

This program does not meet regulatory requirements for proficiency testing; see Survey CGL on page 158. For additional information about the CAP Quality Cross Check program, see page 42.

- Three 1.0-mL lyophilized plasma specimens in triplicate, one 1.0-mL plasma specimen, and one 2.0-mL serum specimen
- · Report up to three instruments
- Two shipments per year

Quality Cross Check— Activated Clotting Time Series CTQ, CT1Q, CT2Q, CT3Q, CT5Q

Instrument/Cartridge	Program Code				Challenges/ Shipment	
	CTQ	CT1Q	CT2Q	CT3Q	CT5Q	
Helena Actalyke®						3
Helena Cascade POC						3
IL Gem® PCL ACT						3
IL Gem PCL ACT-LR						3
IL GEM PCL Plus ACT						3
IL GEM PCL Plus ACT-LR						3
ITC Hemochron® CA510/FTCA510						3
ITC Hemochron FTK-ACT						3
ITC Hemochron Jr. Signature/ACT+						3
ITC Hemochron Jr. Signature/ACT-LR						3
ITC Hemochron P214/P215	•					3
i-STAT Celite® and Kaolin ACT						3
Medtronic HemoTec ACT/ACTII/ACT Plus HR-ACT						3
Medtronic HemoTec ACT/ACTII/ACT Plus LR-ACT		•				3
Medtronic HemoTec ACT/ACTII/ACT Plus R-ACT						3
Medtronic Hepcon HMS, HMS Plus						3
Sienco Sonoclot®	•					3

These programs do not meet regulatory requirements for proficiency testing; see Surveys CT-CT3 and CT5 on page 162. For additional information about the CAP Quality Cross Check program, see page 42.

- CTQ Three 3.0-mL lyophilized whole blood specimens in triplicate with corresponding diluents
- CT1Q Three 1.7-mL lyophilized whole blood specimens in triplicate with corresponding diluents
- CT2Q Three 0.5-mL lyophilized whole blood/ diluent ampules in triplicate
- CT3Q Three 0.5-mL lyophilized whole blood/ diluent ampules in triplicate
- CT5Q Three 1.7-mL lyophilized whole blood specimens in triplicate with corresponding diluents
- Report up to three instruments
- Two shipments per year

5

Point-of-Care Programs



The CAP broadens its network of laboratory experts through its collaborations.

Among the organizations we partner with:

- American Association for Clinical Chemistry (AACC)
- American College of Medical Genetics and Genomics (ACMG)
- Association for Molecular Pathology (AMP)
- National Society for Histotechnology (NSH)

Point-of-Care Programs

POC Competency Challenges are designed to improve waived test results. These programs evaluate instrument and method performance, troubleshoot, assess staff competency, and provide information to train staff. Expected results will be provided. These programs are not proficiency testing programs and participants will not return results to the CAP.

POC Competency Challenges may have limited availability and stability.

POC Competency Challenges POC1, POC2, POC3, POC4						
Program Name	Program Code Challenges/ Shipment					
	POC1	POC2	POC4			
hCG Competency					10	
Glucose Competency					10	
Urine Dipstick Competency					10	
Strep Screen Competency					10	

Program Information

- POC1 One positive 10.0-mL liquid urine specimen
- POC2 One abnormal 2.0-mL whole blood specimen
- POC3 One abnormal 10.0-mL liquid urine specimen
- POC4 One 1.0-mL positive liquid specimen
- Each program provides material to test up to 10 staff
- Shipments available upon request

POC Competency Challenges POC6, POC7, POC8, POC9						
Program Name					Challenges/ Shipment	
	POC6	POC7	POC8	POC9		
PT/INR, CoaguChek XS Plus and XS Pro Competency					10	
Waived Chemistry, Glucose and HgB Competency					10	
Influenza A/B Antigen Detection Competency					10	
Fecal Occult Blood Competency					10	

- POC6 One abnormal 0.3-mL lyophilized plasma specimen (five vials) and five corresponding diluents
- POC7 One abnormal 2.5-mL whole blood specimen compatible with the HemoCue® B, HemoCue 201, and Stanbio HemoPoint® H2 instruments
- POC8 One 1.5-mL positive liquid specimen for influenza A; one 1.5-mL positive liquid specimen for influenza B
- POC9 One positive 2.0-mL fecal specimen
- Each program provides material to test up to 10 staff
- Shipments available upon request

POC Competency Challenges POC10, POC11, POC12						
Program Name	Program Code Challenges/ Shipment					
	POC10					
Blood Gases Competency				10		
Blood Gases, i-STAT® Competency				10		
Plasma Cardiac Markers Competency			•	10		

Program Information

- POC10 One abnormal 2.5-mL aqueous blood gas specimen (ten vials) and one 2.5-mL hematocrit/ hemoglobin specimen (ten vials)
- POC11 One abnormal 2.5-mL aqueous specimen (ten vials) for blood gas and hematocrit/hemoglobin testing
- POC12 One 1.5-mL plasma specimen (two vials); compatible with plasma based tests, such as Alere Triage® and i-STAT instruments
- Programs provide material to test up to 10 staff
- Shipments available upon request

Guide your point-of-care testing with confidence

Point-of-Care Testing (POCT) Toolkit

POCT implementation requires a systematic approach that involves all stakeholders. This toolkit serves as a resource for any member of the POCT team who wants to learn about POCT or who has responsibility to guide or direct POCT. Pathologists may also use the toolkit to guide other members of their POCT teams, including POCT coordinators and medical technologists who are involved in POCT.

The toolkit covers:

- POCT advantages and disadvantages
- Current and projected technology
- Pathologist, laboratory director, and POCT coordinator roles in POCT
- Selection of appropriate test methods
- Validation and verification protocols
- Quality control and data management
- Patient safety
- · POCT training and competency

Purchase the ebook at ebooks.cap.org.



POC Competency Challenges POC14, POC15, POC16						
Program Name	Program Code Challenges/ Shipment					
	P0C14					
Medtronic ACT/ACT, i-STAT Competency				5		
Hemochron Jr IL GEM PCL ACT-LR Competency		•		5		
Hemochron Jr Signature IL GEM PCL ACT Competency			1	5		

Program Information

- POC14 Five abnormal
 1.7-mL lyophilized whole
 blood specimens with five
 corresponding diluents
 and one calcium chloride
 diluent vial; compatible with
 Medtronic HemoTect ACT/
 ACTII/ACT Plus, Medtronic
 Hepcon HMS/HMS Plus, and
 i-STAT Celine and Kaolin ACT
- POC15 Five abnormal 0.5-mL lyophilized whole blood/diluent ampules; compatible with IL GEM PCL Plus ACT-LR and ITC Hemochron Jr./Signature ACT-LR
- POC16 Five abnormal
 0.5-mL lyophilized whole
 blood/diluent ampules;
 compatible with IL GEM PCL
 Plus ACT and ITC Hemochron
 Jr./Signature ACT+
- Programs provide material to test up to five staff
- Shipments available upon request

We are here to help. Fast Focus on Compliance—the inspector's quick guide

Multiply inspector confidence in just 10 minutes with the CAP's Fast Focus on Compliance mini-training vignettes. Using real world examples, inspectors will arm themselves with practical approaches to handle new and perplexing topics:

- Inspecting Personnel Records
- 12 Inspector Tools to Make Your Inspection Go More Smoothly
- Competency Assessment
- Identifying Systemic Issues—Critical Role of the Inspection Team Leader
- Documenting Your Inspection Findings
- IQCP—What It Means to the Inspector

Access these concentrated topics online by searching Inspector Training at cap.org.

General Chemistry and Therapeutic Drug Monitoring



Standardize hemoglobin A_{1c} testing with our Hemoglobin A_{1c} (GH2/GH5) Surveys.

- Mimic patient testing using specimens from human donors with levels that reflect clinical decision points.
- Ensure accuracy of testing—these Surveys are evaluated against the National Glycohemoglobin Standardization Program (NGSP) reference method targets.

General Chemistry and Therapeutic Drug Monitoring

General Chemistry and Therapeutic Drug Monitoring	
Urine Chemistry	68
Special Chemistry	71
New Programs NEW	
Trace Metals in Whole Blood (TMWB)	79
New Analyte/Drug Additions	
Follicle-stimulating hormone (FSH)	77
Luteinizing hormone (LH)	77
Pregabalin	60
0	••••••••••••••••••••••••••••••

Discontinued Programs

Lung Maturity (LM, LM1) TDM-Special Double Volume (ZZT) Urine Chemistry, Special (NVM)

General Chemistry and Therapeutic Drug Monitoring

Analytes/procedures in **bold** type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

General Chemistry and Therapeutic Drugs C1, C3/C3X, C4, CZ/CZX/CZ2X, Z

C1, C3/C3X,	C4,	CZ/CZ	-X/ (SZZX, Z		
Analyte		Program Code				Challenges/ Shipment
	C1	C3/C3X	C4	CZ/CZX/ CZ2X	z	
Alanine aminotransferase (ALT/SGPT)	•	•				5
Albumin				ı		5
Alkaline phosphatase				ı		5
Amylase				1		5
Aspartate aminotransferase (AST/SGOT)	•			ı		5
Bilirubin, direct				ı		5
Bilirubin, total*			•	•		5
Calcium						5
Chloride				ı		5
Cholesterol, total			•	1		5
Cortisol						5
Creatine kinase (CK)				ı		5
Creatinine			•	1		5
Glucose			•			5
HDL cholesterol				ı		5
Human chorionic gonadotropin (hCG), quantitative	ı					5
Iron				ı		5
Lactate dehydrogenase (LD)				ı		5
LDL cholesterol				ı		5
Lipoprotein (a)				1		5
Magnesium				ı		5
Pancreatic amylase				1		5
Potassium						5
Protein, total				ı		5
Sodium				ı		5
T3, free (triiodothyronine, free)						5
T3, total (triiodothyronine, total)				ı		5
T3, uptake and related tests				ı		5
Continued on the next page						

- C1, C3, C4, CZ, Z Five 5.0-mL liquid serum specimens
- C3X, CZX Five 5.0-mL liquid serum specimens in duplicate
- CZ2X Five 5.0-mL liquid serum specimens in triplicate
- Conventional and International System of Units (SI) reporting offered
- Three shipments per year
- For second instrument reporting options, see the Quality Cross Check program, CZQ, on page 59



^{*}General Chemistry and Therapeutic Drugs Surveys do not fulfill the CAP accreditation requirements for neonatal bilirubin proficiency testing. See Surveys NB, NB2 on page 65.

General Chemistry and Therapeutic Drugs C1, C3/C3X, C4, CZ/CZX/CZ2X, Z continued Challenges/ Analyte **Program Code** Shipment CZ/CZX/ C1 C3/C3X C4 CZ2X Z 5 T4, free (thyroxine, free) T4, total (thyroxine, total) 5 Thyroid-stimulating 5 П hormone (TSH) **Triglycerides** П 5 Urea nitrogen (BUN) 5 Uric acid 5 Acid phosphatase 5 ı П 5 Ammonia Apolipoprotein A1 5 П ī 5 Apolipoprotein B Calcium, ionized 5 Ī 5 Carbon dioxide (CO₂) Ī Ferritin 5 П 5 Gamma glutamyl transferase (GGT) Iron binding capacity, 5 total (measured) Iron binding capacity, ī 5 unsaturated (measured) Iron saturation (%) ı 5 Lactate Ī 5 Lipase ī 5 5 Osmolality Ī Phosphorus (inorganic) 5 Prealbumin 5 Transferrin 5 5 Lithium П Ī Acetaminophen 5 5 Amikacin Ī Caffeine П 5 Carbamazepine П 5 Carbamazepine, free 5 5 Digoxin

ī

5

Digoxin, free

Continued on the next page

- C1, C3, C4, CZ, Z Five 5.0-mL liquid serum specimens
- C3X, CZX Five 5.0-mL liquid serum specimens in duplicate
- CZ2X Five 5.0-mL liquid serum specimens in triplicate
- Conventional and International System of Units (SI) reporting offered
- Three shipments per year
- For second instrument reporting options, see the Quality Cross Check program, CZQ, on page 59



General Chemistry and Therapeutic Drugs C1, C3/C3X, C4, CZ/CZX/CZ2X, Z continued Challenges/ Analyte **Program Code** Shipment CZ/CZX/ C1 C3/C3X C4 CZ2X Ζ Disopyramide 5 5 Ethosuximide Gentamicin 5 Lidocaine 5 П 5 Methotrexate Ī Ī 5 N-acetylprocainamide (NAPA) Phenobarbital 5 П П Phenytoin 5 П Phenytoin, free 5 Primidone ī ī 5 Procainamide 5 Quinidine 5 5 Salicylate П П 5 Theophylline 5 **Tobramycin** П 5 Valproic acid П П 5 Valproic acid, free 5 Vancomycin

- C1, C3, C4, CZ, Z Five 5.0-mL liquid serum specimens
- C3X, CZX Five 5.0-mL liquid serum specimens in duplicate
- CZ2X Five 5.0-mL liquid serum specimens in triplicate
- Conventional and International System of Units (SI) reporting offered
- Three shipments per year
- For second instrument reporting options, see the Quality Cross Check program, CZQ, on page 59



Quality Cross Check—Chemistry and Therapeutic Drug Monitoring CZQ Analyte Program Code Challenges/Shipment CZQ See Survey CZ analytes on pages 56-58

This program does not meet regulatory requirements for proficiency testing; see Survey CZ on pages 56-58. For additional information about the CAP Quality Cross Check program, see page 42.

The Quality Cross Check Program:

- Provides a solution for monitoring performance across multiple instruments, and is in compliance with the CMS directive regarding proficiency testing on multiple instruments.
- Simplifies instrument comparability efforts by providing custom reports with both peer group comparison and instrument comparability statistics.

CAP/AACC Immunosuppressive Drugs CS						
Analyte Program Code Challenges/Shipment						
	cs					
Cyclosporine	I	3				
Sirolimus (rapamycin)	I	3				
Tacrolimus	I	3				

Mycophenolic Acid MPA					
Analyte	Program Code Challenges/Shipmen				
	MPA				
Mycophenolic acid	I	3			

Program Information

- Three 5.0-mL liquid serum specimens in duplicate
- Report up to three instruments
- Conventional and International System of Units (SI) reporting offered
- Two shipments per year

Program Information

- Three 5.0-mL whole blood specimens
- For laboratories monitoring cyclosporine, sirolimus, and tacrolimus in transplant patients
- Two shipments per year



Program Information

- Three 4.0-mL whole blood specimens
- Two shipments per year

- Three 5.0-mL lyophilized serum specimens
- Two shipments per year

Therapeutic Drug Monitoring—Extended ZE						
Analyte	Program Code Challenges/Ship					
	ZE					
Clozapine	ı	3				
Gabapentin	ı	3				
Lacosamide	ı	3				
Lamotrigine	ı	3				
Levetiracetam	ı	3				
Oxcarbazepine metabolite	ı	3				
Pregabalin NEW	ı	3				
Rufinamide		3				
Teriflunomide	•	3				
Topiramate	ı	3				
Zonisamide		3				

Program Information

- Three 5.0-mL serum specimens
- Two shipments per year

Therapeutic Drug Monitoring—Special ZT					
Analyte	Program Code Challenges/Shipr				
	ZT				
Amitriptyline	I	3			
Desipramine		3			
Imipramine		3			
Nortriptyline	I	3			
Tricyclics, total (qualitative/ quantitative)		3			

Program Information

- Three 5.0-mL lyophilized serum specimens
- Two shipments per year

Have you created or updated your CAP Profile?

Each laboratory staff member should have their own profile. Your profile is transferrable when you leave your current position. Use it to maintain information about yourself, including:

- Business affiliations
- Personal contact information

Certifications

- Specialties and skills
- Contact preferences
- Addresses
- Inspector-related information

To create or update your profile, visit cap.org, log in, and click on UPDATE MY PROFILE.



Accuracy-Based Lipids ABL					
Analyte Program Code Challenges/Ship					
	ABL				
Apolipoprotein A1*	1	3			
Apolipoprotein B*	1	3			
Cholesterol*	1	3			
HDL cholesterol*	1	3			
Non-HDL cholesterol		3			
LDL cholesterol		3			
Lipoprotein (a)	1	3			
Triglycerides*		3			

^{*}This analyte will be evaluated against the reference method.

B-Type Natriuretic Peptides BNP, BNP5			
Analyte	Challenges/Shipment		
	Program Code		
	BNP BNP5		
BNP	2	5	
NT-proBNP	2	5	

Additional Information

- The College of American Pathologists Accreditation Program requires all accredited laboratories performing non-waived testing for BNP and NT-proBNP to complete 15 PT challenges per year.
- For i-STAT®, use Plasma Cardiac Markers programs PCARM or PCARMX.
- For second instrument reporting options, see the Quality Cross Check program, BNPQ, below.

Quality Cross Check—BNP BNPQ		
Analyte	Program Code	Challenges/Shipment
	BNPQ	
BNP	•	3
NT-proBNP		3

This program does not meet regulatory requirements for proficiency testing; see Survey BNP or BNP5 above. For additional information about the CAP Quality Cross Check program, see page 42.

The Quality Cross Check Program:

- Provides a solution for monitoring performance across multiple instruments, and is in compliance with the CMS directive regarding proficiency testing on multiple instruments.
- Simplifies instrument comparability efforts by providing custom reports with both peer group comparison and instrument comparability statistics.

Program Information

- Three 1.0-mL human serum specimens
- Two shipments per year

Program Information

- BNP Two 1.0-mL liquid plasma specimens
- Conventional and International System of Units (SI) reporting offered; two shipments per year
- BNP5 Five 1.0-mL liquid plasma specimens
- Conventional and International System of Units (SI) reporting offered; three shipments per year

- Three 1.5-mL liquid specimens
- Report up to three instruments
- Conventional and International System of Units (SI) reporting offered
- · Two shipments per year

Harmonized Thyroid ABTH		
Analyte	Program Code	Challenges/Shipment
	ABTH	
T3, free (triiodothyronine, free)	ı	3
T3, total (triiodothyronine, total)		3
T4, free (thyroxine, free)	I	3
T4, total (thyroxine, total)	ı	3
Thyroid-stimulating hormone (TSH)	ı	3

Program Information

- Three 1.0-mL frozen human specimens
- Two shipments per year

Additional Information

- Analytes will be evaluated using harmonization.
- Specimens are collected by a modified application of Clinical Laboratory and Standards Institute Guideline CLSI C37-A, Preparation and Validation of Commutable Frozen Human Serum Pools as Secondary Reference Materials for Cholesterol Measurement Procedures; Approved Guideline.

Cardiac Markers CRT, CRTI, TNT, TNT5 Challenges/ Analyte **Program Code** Shipment **CRT CRTI TNT** TNT5 ī 5 CK-MB, immunochemical CK isoenzymes (CK-BB, CK-MB, П 5 CK-MM), electrophoretic LD1, LD2, LD3, LD4, LD5, Ī 5 electrophoretic LD1/LD2 ratio calculation 5 and interpretation Myoglobin 2 5 Troponin I Ī 2 Troponin T, two challenges ī 5 Troponin T, five challenges

The College of American Pathologists Accreditation Program requires all accredited laboratories performing non-waived testing for Troponin I and Troponin T to complete 15 PT challenges per year.

- CRT Five 2.0-mL liquid serum specimens
- CRTI Ten 2.0-mL liquid serum specimens
- TNT Two 2.0-mL liquid serum specimens
- TNT5 Five 2.0-mL liquid serum specimens
- · Three shipments per year

Hemoglobin A _{1c} GH2, GH5		
Analyte	Challenges	/Shipment
	Program Code	
	GH2	GH5
Hemoglobin A _{1c}	3	5

Additional Information

- These Surveys will be evaluated against the National Glycohemoglobin Standardization Program (NGSP) reference method.
- The College of American Pathologists Accreditation Program requires all accredited laboratories performing non-waived testing for Hemoglobin A_{1c} to complete 15 PT challenges per year.
- For second instrument reporting options, see the Quality Cross Check program, GHQ, below.

Quality Cross Check—Hemoglobin A _{1c} GHQ		
Analyte	Program Code	Challenges/Shipment
	GHQ	
Hemoglobin A _{1c}		3

This program does not meet regulatory requirements for proficiency testing; see Survey GH5 above. For additional information about the CAP Quality Cross Check program, see page 42.

The Quality Cross Check Program:

- Provides a solution for monitoring performance across multiple instruments, and is in compliance with the CMS directive regarding proficiency testing on multiple instruments.
- Simplifies instrument comparability efforts by providing custom reports with both peer group comparison and instrument comparability statistics.

Hemoglobin A _{1c} GH5I		
Analyte	Program Code	Challenges/Shipment
	GH5I	
Hemoglobin A _{1c}		5

Additional Information

- This program meets the CAP's Accreditation Program requirements for proficency testing.
- This Survey will not be evaluated against the National Glycohemoglobin Standardization Program (NGSP) reference method. See Survey GH5 to be evaluated against the NGSP reference method.

Program Information

- GH2 Three 0.8-mL liquid human whole blood specimens; two shipments per year
- GH5 Five 0.8-mL liquid human whole blood specimens; three shipments per year

Program Information

- Three 0.8-mL liquid specimens in triplicate
- Report up to three instruments
- · Two shipments per year

- Five 0.5-mL lyophilized specimens with a 3.0-mL dropper-tipped vial of diluent
- Designed for international laboratories that have experienced significant shipping and receiving issues and require longer specimen stability
- Three shipments per year

Glycated Serum Albumin GSA		
Analyte	Program Code	Challenges/Shipment
	GSA	
Glycated serum albumin	I	3

Program Information • Three 1.0-mL liquid serum

specimens

 Two shipments per ye 	ar
--	----

High-Sensitivity C-Reactive Protein HSCRP			
Analyte	Program Code	Challenges/Shipment	
	HSCRP		
High-sensitivity C-reactive protein	I	3	

Program Information

- Three 0.5-mL liquid serum specimens
- Two shipments per year

Homocysteine HMS		
Analyte	Program Code	Challenges/Shipment
	HMS	
Homocysteine		3

Program Information

- Three 1.0-mL serum specimens
- Two shipments per year

Ketones KET		
Analyte	Program Code	Challenges/Shipment
	KET	
Beta-hydroxybutyrate	I	2
Total ketones	ı	2

Program Information

- Two 2.0-mL serum specimens
- For use with Acetest® and other qualitative/semiquantitative methods using the nitroprusside reaction for total ketones testing
- Two shipments per year

Chemistry—Limited, Waived LCW		
Analyte	Program Code	Challenges/Shipment
	LCW	
Cholesterol		3
Glucose	•	3
HDL cholesterol		3
LDL cholesterol		3
Triglycerides	I	3

- Three 3.0-mL liquid serum specimens
- For use with waived methods such as the Cholestech LDX® and Roche ACCU-CHEK® Instant Plus
- The glucose specimens are not appropriate for use on other whole blood glucose meters
- Two shipments per year

Neonatal Bilirubin NB, NB2			
Analyte Challenges/Shipment			
Program Code			
NB NB2			
Bilirubin, direct 2 2		2	
Bilirubin, total	5	2	

One human-based serum specimen will offer the value assigned using the reference method procedure (*Clin Chem.* 1985;31:1779-1789).

Program Information

- NB Five 1.0-mL human serum specimens; three shipments per year
- NB2 Two 1.0-mL human serum specimens; must order in conjunction with a five-challenge total bilirubin proficiency testing program to meet regulatory requirements; two shipments per year
- Conventional and International System of Units (SI) reporting offered

Plasma Cardiac Markers PCARM, PCARMX				
Analyte	Progra	Program Code Challenges/Shipment		
	PCARM	PCARMX		
BNP	1		5	
CK-MB	ı		5	
D-dimer	1		2	
Myoglobin	ı		2	
Troponin I			5	

The College of American Pathologists Accreditation Program requires all accredited laboratories performing non-waived testing for BNP and Troponin I to complete 15 PT challenges per year.

Program Information

- PCARM Five 1.5-mL liquid EDTA plasma specimens
- PCARMX All Survey PCARM specimens in duplicate
- For Point-of-Care instruments such as Alere Triage[®] and i-STAT
- Three shipments per year

Test your diagnostic skills as a pathologist with CPIP

Online, hands-on and interactive, the Clinical Pathology Improvement Program (CPIP) enables pathologists to sharpen their diagnostic skills in real time by working through an actual case. Each month, you will receive a new scenario, including slide images and clinical background. As the case unfolds, more information is revealed, just as in the laboratory. Participants who successfully complete the posttest may apply their earned credits to their MOC SAM requirements. Enjoy a full year of CPIP and earn up to 15 CME/SAM credits.

Choose code CPIP/CPIP1 on your Surveys order form.

Whole Blood Chemistry Compatibility Matrix

Whole Blood Analyzer/Method	Analyte	Compatible Survey Programs	Page
Hemocue®	Glucose	HCC	66
Roche Reflotron®	Cholesterol	01.04	56-57
	Glucose	C1, C4	56-57
Cholestech LDX®	Total cholesterol		64
	HDL cholesterol	LCW	64
	Triglycerides	LCVV	64
	Glucose		64
Whole blood cholesterol meters	Cholesterol	C1, C4, LCW	56-57, 64
Whole blood glucose meters	Glucose	HCC2, WBGQ	66, 67
Nova StatSensor®/ E-Z-EM EZ Chem™	Creatinine	WBCR	66

Waived Combination HCC, HCC2			
Analyte	Analyte Program Code Challenges/Shipment		
	HCC	HCC2	
Hematocrit			2
Hemoglobin			2
Urinalysis/Urine hCG			2
Whole blood glucose			2 (HCC)/3 (HCC2)

Program Information

- HCC Two 1.0-mL whole blood specimens; two shipments per year
- HCC2 Total of four shipments per year
- Hematocrit, hemoglobin, and urinalysis/urine hCG testing -Two 3.0-mL whole blood specimens and two 10.0-mL urine specimens; two shipments per year: A and C
- Whole blood glucose testing - Three 2.5-mL whole blood specimens; two shipments per year: B and D
- To verify instrument compatibility, refer to the instrument matrix on this page

Whole Blood Creatinine WBCR				
Analyte Program Code Challenges/Shipment				
WBCR				
Creatinine		5		

- Five 4.0-mL whole blood specimens
- For use with the Nova StatSensor®/E-Z-EM EX Chem™
- Three shipments per year

Quality Cross Check—Whole Blood Glucose WBGQ				
Analyte Program Code Challenges/Shipment				
WBGQ				
Glucose	ı	3		

The CAP Accreditation Program requires all accredited laboratories performing waived whole blood glucose testing using glucose meters to perform alternative performance assessment. This program can be used to meet alternative performance assessment requirements.

The Quality Cross Check Program:

- Provides a solution for monitoring performance across multiple instruments, and is in compliance with the CMS directive regarding proficiency testing on multiple instruments.
- Simplifies instrument comparability efforts by providing custom reports with both peer group comparison and instrument comparability statistics.

Improve the reliability of your patient results with CAP Survey Validated Materials

Use the same material that is sent in the Surveys program to:

- Identify and troubleshoot instrument/method problems
- · Correlate results with other laboratories or instruments
- · Document correction of problems identified in Surveys
- · Utilize material with confirmed results as an alternative external quality control
- · Identify potential proficiency testing failures

Each laboratory receives a Survey Participant Summary, which includes readily available results.

Chemistry/TDM—Validated Material

Validated Material	Program Code	Corresponding Survey	Pages
Chemistry/TDM	CZVM	CZ	56-58

Program Information

• Five 5.0-mL liquid serum specimens

Simplify your workflow with CVL, the most comprehensive solution for calibration verification and linearity

- Assure confidence in your linearity results with 5-8 specimen levels covering the analytical measurement range (AMR) for most clinical analyzers.
- Mimic patient testing with human serum-based CVL specimens that are shipped twice per year to match frequency of regulatory requirements.
- Receive thorough feedback with a customized report that contains the most rigorous statistical analysis.
- Expedite linearity evaluations within two business days through our *LN Express*SM Service.

See the Instrumentation Validation Tools section of this catalog to determine programs that best fit your laboratory's CVL needs.

- Three 2.0-mL whole blood specimens
- Report up to 30 instruments
- · Two shipments per year



Urine Chemistry

Analytes/procedures in **bold** type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Urine Chem	nistry—General	J
Analyte	Program Code	Challenges/Shipment
	U	
Amylase	I	3
Calcium		3
Chloride		3
Creatinine	I	3
Glucose		3
Magnesium		3
Nitrogen, total	•	3
Osmolality		3
Phosphorus	I	3
Potassium	I	3
Protein, total		3
Sodium		3
Urea nitrogen	ı	3
Uric acid		3
Urine albumin, quantitative		3
Urine albumin:creatinine ratio	I	3

Program Information

- Six 15.0-mL urine specimens
- One mailing per year will include an additional educational specimen for uric acid testing for a total of seven challenges per year
- Conventional and International System of Units (SI) reporting offered
- · Two shipments per year

Accuracy-Based Urine ABU			
Analyte	Program Code	Challenges/Shipment	
	ABU		
Calcium	ı	3	
Creatinine	I	3	
Urine albumin, quantitative	ı	3	
Urine albumin: creatinine ratio		3	

Target values for albumin are obtained by LC-MS/MS after trypsin digestion, performed by the Renal Testing Laboratory, Mayo Clinic, Rochester, MN, using calibration materials prepared from human serum albumin (>99% pure).

Other analytes will be compared by peer group for harmonization purposes.

- Three 5.0-mL human urine specimens
- Two shipments per year

Kidney Stone Risk Assessment KSA			
Analyte	Program Code	Challenges/Shipment	
	KSA		
Citrate	I	3	
Cystine	I	3	
Oxalate	ı	3	
Sulfate	I	3	

Program Information

- Three 13.5-mL liquid urine specimens
- Two shipments per year

Urine Chemistry—Special N, NX			
Analyte	Program Code	Challenges/Shipment	
	N, NX		
3-methoxytyramines		3	
5-hydroxyindoleacetic acid	ı	3	
17-hydroxycorticosteroids		3	
17-ketosteroids		3	
Aldosterone	I	3	
Coproporphyrins		3	
Cortisol, urinary free		3	
Dopamine		3	
Epinephrine		3	
Homovanillic acid	•	3	
Metanephrine		3	
Norepinephrine	I	3	
Normetanephrine	I	3	
Uroporphyrin	ı	3	
Vanillylmandelic acid	I	3	

Program Information

- N Six 10.0-mL lyophilized urine specimens and three 10.0-mL liquid urine specimens
- NX All lyophilized Survey N specimens in duplicate and three 10.0-mL liquid urine specimens
- Two shipments per year

Myoglobin, Urine MYG			
Analyte	Program Code	Challenges/Shipment	
	MYG		
Myoglobin, urine (qualitative and quantitative)		2	

- Two 1.0-mL urine specimens
- Two shipments per year

Porphobilinogen, Urine UPBG			
Analyte	Program Code	Challenges/Shipment	
	UPBG		
Porphobilinogen	I	3	

Program Information

- Three 5.0-mL urine specimens
- Two shipments per year
- For use with qualitative and quantitative methods

Improve the reliability of your patient results with CAP Survey Validated Materials

Use the same material that is sent in the Surveys program to:

- · Identify and troubleshoot instrument/method problems
- · Correlate results with other laboratories or instruments
- Document correction of problems identified in Surveys
- Utilize material with confirmed results as an alternative external quality control
- · Identify potential proficiency testing failures

Each laboratory receives a Survey Participant Summary, which includes readily available results.

Urine Chemistry—General, Validated Material

Validated Material	Program Code	Corresponding Survey	Page
Urine Chemistry	UVM	U	68

Program Information

• Six 15.0-mL urine specimens

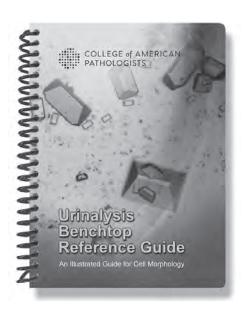
Urinalysis Benchtop Reference Guide (UABRG)

- Thirty-four different cell identifications, including common and rare cells
- Detailed descriptions for each cell morphology
- Eight tabbed sections for easy reference
- A durable and water-resistant format to withstand years of benchtop use—5" by 6½"

Choose code UABRG on your Surveys order form.

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Special Chemistry

Analytes/procedures in **bold** type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

1,5-Anhydroglucitol AG			
Analyte Program Code Challenges/Shipmen			
	AG		
1,5-anhydroglucitol	I	3	

Program Information

- Three 1.0-mL liquid serum specimens
- Two shipments per year

Aldolase ADL			
Analyte	Program Code	Challenges/Shipment	
	ADL		
Aldolase	ı	2	

Program Information

- Two 3.0-mL liquid serum specimens
- Two shipments per year

Angiotensin Converting Enzyme ACE			
Analyte	rte Program Code		
Angiotensin converting enzyme, quantitative		2	

Program Information

- Two 2.0-mL lyophilized serum specimens
- Two shipments per year

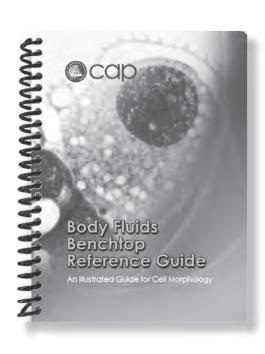
Body Fluids Benchtop Reference Guide (BFBRG)

- Thirty-six color images, including common and rare cells, crystals, and other cell inclusions
- Detailed descriptions of each cell including facts, cell morphology and inclusions
- · Nine tabbed sections for easy reference
 - Erythroid Series
 - Lymphoid Series
 - Myeloid Series
 - Mononuclear Phagocytic Series
 - Lining Cells
 - o Miscellaneous Cells
 - Crystals
 - Microorganisms
 - Miscellaneous Findings
- A durable and water-resistant format to withstand years of benchtop use—5" x 6½"

Choose code BFBRG on your Surveys order form.

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Body Fluid Chemistry FLD			
Analyte	Program Code	Challenges/Shipment	
	FLD		
Albumin	ı	3	
Amylase		3	
CA19-9	ı	1	
CEA	ı	1	
Cholesterol	ı	3	
Creatinine	I	3	
Glucose	ı	3	
Lactate	ı	3	
Lactate dehydrogenase (LD)	I	3	
рН	I	3	
Protein, total	I	3	
Triglycerides	I	3	
Urea nitrogen	ı	1 per year	

- Three 3.0-mL simulated liquid body fluid specimens
- Conventional and International System of Units (SI) reporting offered
- · Two shipments per year

Additional Information

• For second instrument reporting options, see the Quality Cross Check program, FLDQ, on page 73.

Approximately 40% of all CAP proficiency testing (PT) errors are clerical*

Minimize your risk and treat PT more like a patient sample with automated PT reporting from e-LAB Solutions Connect:

- Eliminate errors from manual transcription, reducing PT failures
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 \star Based on data received from the CAP's Laboratory Improvement Programs, 40% of all errors submitted are clerical.



Quality Cross Check—Body Fluid Chemistry FLDQ			
Analyte	Program Code	Challenges/Shipment	
	FLDQ		
Albumin	I	3	
Amylase	I	3	
CA19-9	I	1	
Carcinoembryonic antigen (CEA)	ı	1	
Cholesterol	I	3	
Creatinine	ı	3	
Glucose	ı	3	
Lactate	ı	3	
Lactate dehydrogenase (LD)	ı	3	
рН	ı	3	
Protein, total	ı	3	
Triglycerides	ı	3	
Urea nitrogen	1	1	

This program does not meet regulatory requirements for proficiency testing; see Survey FLD on page 72. For additional information about the CAP Quality Cross Check program, see page 42.

The Quality Cross Check Program:

- Provides a solution for monitoring performance across multiple instruments, and is in compliance with the CMS directive regarding proficiency testing on multiple instruments.
- Simplifies instrument comparability efforts by providing custom reports with both peer group comparison and instrument comparability statistics.

Body Fluid Chemistry 2 FLD2			
Analyte	Program Code	Challenges/Shipment	
	FLD2		
Alkaline phosphatase	•	3	
Bilirubin	ı	3	
Calcium	ı	3	
Chloride	1	3	
Lipase	ı	3	
Potassium	I	3	
Sodium	•	3	
Uric acid	1	3	

Program Information

- Three 3.0-mL specimens in duplicate
- Report up to three instruments
- Two shipments per year

- Three 3.0-mL liquid body fluid specimens
- Conventional and International System of Units (SI) reporting offered
- · Two shipments per year

Cadmium CD			
Analyte	Challenges/Shipment		
	CD		
Beta-2-microglobulin, urine	I	3	
Cadmium, urine		3	
Cadmium, whole blood	ı	3	
Creatinine, urine	ı	3	

This Survey meets the Occupational Safety and Health Administration (OSHA) guidelines for proficiency testing (OSHA standard-29 CFR 1910.1027AppF).

Program Information

- Three 6.0-mL whole blood specimens and three 13.0-mL urine specimens
- Conventional and International System of Units (SI) reporting offered
- · Six shipments per year

Cerebrospinal Fluid Chemistry M, OLI			
Analyte	Prog	ram Code	Challenges/Shipment
	М	OLI	
Albumin, quantitative			3
Electrophoresis (albumin and gamma globulin)	•	•	3
Glucose			3
IgG, quantitative		ı	3
Lactate			3
Lactate dehydrogenase (LD)			3
Protein, total			3
Oligoclonal bands			3

Program Information

- M Three 5.0-mL simulated liquid spinal fluid specimens
- OLI Three 1.0-mL simulated liquid spinal fluid specimens and three paired serum specimens; one educational activity to calculate CSF IgG index and synthesis rate
- Two shipments per year



Cystatin C CYS			
Analyte	Program Code	Challenges/Shipment	
	CYS		
Cystatin C	I	2	

- Two 1.0-mL liquid serum specimens
- Two shipments per year

Fecal Fat FCFS			
Analyte Program Code Challenges/			
	FCFS		
Fecal fat, qualitative	ı	2	

- Two 10.0-g simulated fecal fat specimens
- For microscopic detection of neutral fats (triglycerides) and/or split fats (total free fatty acids)
- Two shipments per year

Fructosamine FT			
Analyte	Program Code	Challenges/Shipment	
	FT		
Fructosamine	I	2	

Program Information

- Two 1.0-mL liquid serum specimens
- Two shipments per year

Glucose-6-Phosphate Dehydrogenase G6PDS				
Analyte Program Code Challenges/Shipment				
G6PDS				
G6PD, qualitative and quantitative	I	2		

Program Information

- Two 0.5-mL lyophilized hemolysate samples
- Two shipments per year

Lipoprotein-Associated Phospholipase A ₂ PLA		
Analyte	Program Code	Challenges/Shipment
	PLA	
Lipoprotein-associated phospholipase (Lp-PLA ₂) concentration	ı	2
Lipoprotein-associated phospholipase (Lp-PLA ₂) activity	ı	2

- Two 1.0-mL liquid specimens
- Two shipments per year

Lipoprotein and Protein Electrophoresis LPE, SPE, UBJP Analyte **Program Code** Challenges/Shipment LPE SPE **UBJP** Lipoprotein electrophoresis 2 IgA, quantitation 2 п IgG, quantitation ı 2 IgM, quantitation ı 2 2 M-protein (Paraprotein) identification П 2 Protein, total 2 Protein electrophoresis Protein electrophoresis pattern Ī 2 interpretation 2 Urine Bence Jones proteins

Program Information

- LPE Two 1.0-mL liquid serum specimens
- SPE Two 1.0-mL lyophilized serum specimens; two educational protein electrophoresis dry challenges per year
- UBJP Two 10.0-mL urine specimens
- Two shipments per year



Lamellar Body Count LBC		
Procedure	Program Code	Challenges/Shipment
	LBC	
Lamellar body count	ı	3

Program Information

- Three 2.0-mL simulated liquid amniotic fluid specimens
- For use with LBC methods performed on all hematology analyzers
- Two shipments per year

Plasma Hemoglobin PHG		
Analyte	Program Code	Challenges/Shipment
	PHG	
Plasma hemoglobin		2

- Two 2.0-mL liquid specimens
- Two shipments per year

Procalcitonin PCT		
Analyte	Program Code	Challenges/Shipment
	PCT	
Procalcitonin	ı	3

Pseudocholinesterase C7			
Analyte	Program Code	Challenges/Shipment	
	C7		

- Three 1.0-mL lyophilized serum specimens
- Two shipments per year

- · One 2.0-mL lyophilized serum specimen
- Three shipments per year



1

Salivary Cortisol SALC		
Analyte	Program Code	Challenges/Shipment
	SALC	
Salivary cortisol	I	3

Accuracy-Based Testosterone, Estradiol ABS **Analyte Program Code** Challenges/Shipment **ABS** Albumin 3 Calcium 3 Cortisol ī 3 3 Estradiol Follicle-stimulating hormone (FSH) NEW 3 3 Luteinizing hormone (LH) NEW ī Sex hormone-binding globulin ī 3 (SHGB) Testosterone 3 3 Testosterone, bioavailable ı Testosterone, free 3 Thyroid-stimulating hormone (TSH) 3

Additional Information

Pseudocholinesterase

• The Centers for Disease Control and Prevention (CDC) will set target values for testosterone and estradiol using the established reference methods.

Program Information

- Three 1.0-mL synthetic oral fluid specimens
- Two shipments per year

- Three 1.0-mL human serum specimens
- Two shipments per year

Total Bile Acids TBLA		
Analyte	Program Code	Challenges/Shipment
	TBLA	
Total bile acids	ı	3

- Three 5.0-mL liquid serum specimens
- Two shipments per year

Trace Metals R		
Analyte	Program Code	Challenges/Shipment
	R	
Aluminum	ı	3
Chromium	ı	3
Copper	ı	3
Manganese	ı	3
Selenium	ı	3
Zinc	ı	3

Program Information

- Three 5.0-mL liquid serum specimens
- Conventional and International System of Units (SI) reporting offered
- Two shipments per year

Trace Metals, Urine TMU		
Analyte	Program Code	Challenges/Shipment
	TMU	
Aluminum	I	2
Arsenic	I	2
Chromium	I	2
Cobalt	I	2
Copper	I	2
Lead	I	2
Manganese	I	2
Mercury	I	2
Selenium	ı	2
Thallium	ı	2
Zinc	1	2

- Two 25.0-mL urine specimens
- Conventional and International System of Units (SI) reporting offered
- For laboratories that monitor trace metals at normal and toxic levels
- Two shipments per year



Trace Metals	, Whole Blood Ti	MWB NEW
Analyte	Program Code	Challenges/Shipment
	TMWB	
Aluminum	I	3
Arsenic, total		3
Chromium	I	3
Cobalt		3
Copper		3
Manganese	ı	3
Mercury	ı	3
Selenium	ı	3
Thallium	I	3
Zinc		3

- Three 6.0-mL whole blood specimens
- Conventional and International System of Units (SI) reporting offered
- Two shipments per year
- Designed for laboratories that monitor trace metals at normal and toxic levels

Sweat Analysis Series SW1, SW2, SW3, SW4			
Analyte Program Code Challenges/Shipmer			
	SW1, SW2, SW3, SW4		
Chloride	•	3	
Conductivity	I	3	

For method compatibility, see chart below.

Sweat Analysis Series Compatibility Matrix

Method/Procedure		Progra	m Code		Materials Included
	SW1	SW2	SW3	SW4	
Orion direct electrode					Precut 2-cm diameter Whatman filter papers
Wescor Macroduct [™] and Nanoduct [®] Systems					22-gauge blunt-tipped needles
CF Indicator System®					Polystyrene boats and chloride-free sponges
All other methodologies					No additional materials provided

- Three 5.0-mL simulated liquid human sweat specimens
- Two shipments per year

Viscosity V				
Analyte	Program Code	Challenges/Shipment		
V				
Viscosity ■ 2				

viscosity v			
Analyte	Program Code	Challenges/Shipment	
	V		
Viscosity		2	

- Two 10.0-mL serum specimens
- Two shipments per year

Soluble Trans	ferrin Receptor	STFR
Analyte	Program Code	Challenges/Shipment
	STFR	
Soluble transferrin receptor (sTfR)	I	3

Program Information

- Three 2.5-mL liquid human serum specimens
- Two shipments per year

Improve the reliability of your patient results with CAP Survey **Validated Materials**

Use the same material that is sent in the Surveys program to:

- · Identify and troubleshoot instrument/method problems
- · Correlate results with other laboratories or instruments
- Document correction of problems identified in Surveys
- · Utilize material with confirmed results as an alternative external quality control
- Identify potential proficiency testing failures

Each laboratory receives a Survey Participant Summary, which includes readily available results.

Cerebrospinal Fluid Validated Material

Validated Material	Program Code	Corresponding Survey	Page
Cerebrospinal Fluid	MVM	М	74

Program Information

• Three 5.0-mL simulated liquid spinal fluid specimens

Endocrinology



Ensure accuracy in your noninvasive prenatal testing with our new NIPT Survey.

- Mimics patient testing with maternal plasma specimens
- Offers compatibility across all major testing methods, including SNP, counting, targeted, and shotgun methods



Endocrinology

Analytes/procedures in **bold** type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Ligand—(General k	K, KK, K2	
Analyte	Program Code		Challenges/Shipment
	K, KK	K2	
Alpha-fetoprotein (AFP)	ı		5
CEA	ı	1	3
Cortisol	ı		5
Ferritin	ı	ı	3
Folate, serum	ı	ı	3
hCG, quantitative	ı		5
Immunoglobulin E (IgE)	ı		5
Prostate-specific antigen (PSA)	ı	ı	2 (K,KK)/3 (K2)
Prostate-specific antigen, complexed (cPSA)	ı		2
Prostate-specific antigen, free	ı		2
Prostatic acid phosphatase (PAP)	ı		3
T3, free (triiodothyronine, free)	ı		5
T3, total (triiodothyronine, total)	ı		5
T3 uptake and related tests	ı		5
T4, free (thyroxine, free)	ı		5
T4, total (thyroxine, total)	ı		5
Thyroid-stimulating hormone (TSH)	I		5
Vitamin B ₁₂	ı	ı	3

- K Five 5.0-mL liquid serum specimens; three shipments per year
- KK Five 5.0-mL liquid serum specimens in duplicate; three shipments per year
- K2 Three 5.0-mL liquid serum specimens; two shipments per year

B-Type Natriuretic Peptides BNP, BNP5			
Analyte Challenges/Shipment			
	Program Code		
	BNP	BNP5	
BNP	2	5	
NT-proBNP	2	5	

Additional Information

- The College of American Pathologists Accreditation Program requires all accredited laboratories performing non-waived testing for BNP and NT-proBNP to complete 15 PT challenges per year.
- For i-STAT®, use Plasma Cardiac Markers programs PCARM or PCARMX.
- For second instrument reporting options, see the Quality Cross Check program, BNPQ, below.

Quality Cross Check—BNP BNPQ				
Analyte Program Code Challenges/Shipment				
BNPQ				
BNP	I	3		
NT-proBNP	3			

This program does not meet regulatory requirements for proficiency testing; see Survey BNP or BNP5 above. For additional information about the CAP Quality Cross Check program, see page 42.

The Quality Cross Check Program:

- Provides a solution for monitoring performance across multiple instruments, and is in compliance with the CMS directive regarding proficiency testing on multiple instruments.
- Simplifies instrument comparability efforts by providing custom reports with both peer group comparison and instrument comparability statistics.

Program Information

- BNP Two 1.0-mL liquid plasma specimens
- Conventional and International System of Units (SI) reporting offered; two shipments per year
- BNP5 Five 1.0-mL liquid plasma specimens
- Conventional and International System of Units (SI) reporting offered; three shipments per year

- Three 1.5-mL liquid specimens
- Report up to three instruments
- Conventional and International System of Units (SI) reporting offered
- · Two shipments per year

Ligand—Special Y, YY, DY			
Analyte	Progra	m Code	Challenges/Shipment
	Y, YY	DY	
11-deoxycortisol			3
17-hydroxyprogesterone			3
Androstenedione			3
DHEA sulfate	•		3
Estradiol	•		3
Estriol, unconjugated (uE3)			3
Follicle-stimulating hormone (FSH)	•		3
Growth hormone (GH)	•		3
IGF-1 (somatomedin C)			3
Luteinizing hormone (LH)	•		3
Progesterone	•		3
Prolactin			3
Testosterone	•		3
Testosterone, bioavailable		1	3
Testosterone, free		•	3
Sex hormone-binding globulin (SHBG)		1	3

- Y-Three 5.0-mL liquid serum specimens in duplicate
- YY Three 5.0-mL liquid serum specimens in triplicate
- · DY Must order in conjunction with Survey Y or
- · Conventional and International System of Units (SI) reporting offered
- Two shipments per year



Antimüllerian Hormone AMH				
Analyte Program Code Challenges/Shipment				
	АМН			
Antimüllerian hormone	I 3			

Antimüllerian Hormone AMH				
Analyte Program Code Challenges/Shipment				
АМН				
Antimüllerian hormone 3				

25-OH Vitamin D, Total VITD Analyte **Program Code** Challenges/Shipment VITD

Program Information

- Three 1.0-mL lyophilized serum specimens
- Two shipments per year

Program Information

3

- Three 1.0-mL liquid serum specimens
- · Conventional and International System of Units (SI) reporting offered
- Two shipments per year

25-OH vitamin D, total

Bone and Growth BGS			
Analyte	Program Code	Challenges/Shipment	
	BGS		
IGF-1 (somatomedin C)	I	3	
Osteocalcin	I	3	

- Three 1.0-mL liquid serum specimens
- Conventional and International System of Units (SI) reporting offered
- Two shipments per year



Accuracy-Based Vitamin D ABVD				
Analyte	Program Code Challenges/Shipment			
	ABVD			
25-OH vitamin D (D2 and D3)	I	3		

Additional Information

- The Centers for Disease Control and Prevention (CDC) will establish reference targets using isotope-dilution LC-MS/MS method.
- Specimens are collected by a modified application of Clinical Laboratory and Standards Institute Guideline CLSI C37-A, Preparation and Validation of Commutable Frozen Human Serum Pools as Secondary Reference Materials for Cholesterol Measurement Procedures; Approved Guideline.

Bone and Mineral Metabolism, Urine BU					
Analyte Program Code Challenges/Shipme					
	BU				
C-telopeptide (CTx)	I	2			
Creatinine	•	2			
Deoxypyridinoline (DPD)	•	2			
N-telopeptide (NTx)	•	2			
Pyridinoline (PYD)	I	2			

Program Information

- Three 1.0-mL liquid human serum specimens
- Serum is from multi-donor endogenous pools
- Conventional and International System of Units (SI) reporting offered
- Two shipments per year

- Two 2.0-mL lyophilized human urine specimens
- Conventional and International System of Units (SI) reporting offered
- Two shipments per year



N-telopeptide

Bone Markers and Vitamins BMV1, BMV2, BMV3, BMV4, BMV5, BMV6 Challenges/ Analyte **Program Code** Shipment BMV1 BMV2 BMV3 BMV4 BMV5 BMV6 1,25 dihydroxy 3 vitamin D Bone-specific 3 alkaline phosphatase ı 3 Vitamin A Vitamin E (alpha 3 tocopherol, gamma tocopherol) C-telopeptide 3

Program Information

- BMV1 through BMV4 Three 5.0-mL liquid serum
 specimens for each program
- BMV5 and BMV6 Three
 1.0-mL liquid serum
 specimens for each program
- Two shipments per year

Erythropoietin EPO			
Analyte	Program Code	Challenges/Shipment	
	EP0		
Erythropoietin	I	2	

Program Information

- Two 1.5-mL serum specimens
- Two shipments per year



3

П

Fetal Fibronectin FF					
Analyte Program Code Challenges/Shipment					
FF					
Fetal fibronectin 2					

Program Information

- Two 1.2-mL liquid specimens
- Two shipments per year

Insulin, Gastrin, C-Peptide, and PTH ING						
Analyte	yte Program Code Challenges/Shipment					
	ING					
C-peptide	I	3				
Gastrin	■ 3					
Insulin	■ 3					
Parathyroid hormone (PTH)	ı	3				

- Three 5.0-mL lyophilized serum specimens
- Conventional and International System of Units (SI) reporting offered
- · Two shipments per year



Second Trimester Maternal Screening FP, FPX						
Analyte	Program Code Challenges/Shipmen					
	FP, FPX					
Alpha-fetoprotein (AFP), amniotic fluid	ı	2				
Alpha-fetoprotein (AFP), serum	1	5				
Dimeric inhibin A (DIA)	ı	5				
Estriol, unconjugated (uE3)	ı	5				
Human chorionic gonadotropin (hCG), quantitative	ı	5				

The CAP designed this Survey for laboratories using AFP and hCG for prenatal screening purposes only. For all other applications, see Survey K or KK on page 82.

Program Information

- FP Five 1.0-mL liquid serum specimens; two 1.0-mL simulated amniotic fluid specimens
- FPX All Survey FP serum specimens in duplicate; two 1.0-mL simulated amniotic fluid specimens
- Three shipments per year

First Trimester Maternal Screening FP1T, FP1B					
Analyte	Progra	Program Code Challenges/Shipment			
	FP1T	FP1T FP1B			
Total hCG			5		
Free beta hCG			5		
PAPP-A	ı		5		

The CAP designed these Surveys for laboratories using hCG for prenatal screening purposes only. For all other applications, see Survey K or KK on page 82.

Program Information

- FP1T Five 1.0-mL serum specimens
- FP1B Five 1.0-mL serum specimens
- Three shipments per year

Noninvasive Pre	enatal Testing	NIPT NEW
Analyte	Program Code	Challenges/Shipment
	NIPT	
Cell-free DNA screening for fetal aneuploidy	ı	3

Noninvasive Pre	enatal Testing	NIPT (NEW)
Analyte	Program Code	Challenges/Shipment
	NIPT	
Cell-free DNA screening for fetal aneuploidy	ı	3

Program Information

- · Three maternal plasma samples
- Two shipments per year

Quality Cross Check—Parathyroid Hormone PTHQ					
Analyte Program Code Challenges/Shipmen					
	PTHQ				
Parathyroid hormone (PTH)		3			

This program does not meet regulatory requirements for proficiency testing; see Survey ING on page 86. For additional information about the CAP Quality Cross Check program, see page 42.

The Quality Cross Check Program:

- · Provides a solution for monitoring performance across multiple instruments, and is in compliance with the CMS directive regarding proficiency testing on multiple instruments.
- Simplifies instrument comparability efforts by providing custom reports with both peer group comparison and instrument comparability statistics.

- Three 5.0-mL lyophilized serum specimens in duplicate
- · Report up to three instruments
- Two shipments per year

Pharmacogenetics PGX, PGX1, PGX2, PGX3					
Analyte/Procedure		Program Code			Challenges/ Shipment
	PGX	PGX1	PGX2	PGX3	
CYP2C19	•				3
CYP2C9					3
CYP2D6					3
CYP3A4					3
CYP3A5					3
SLC01B1 (rs4149056)					3
VKORC1	•				3
IL28B (rs12979860)		•			3
HLA-B*15:02					3
HLA-B*57:01			•		3
DPYD				1	3
TPMT				1	3
UGT1A1				ı	3

Additional Information

- UGT1A1 (PGX3 Survey) tests the laboratory's ability to detect variants in the TATA
 repeat sequence in the UGT1A1 promotor (eg, UGT1A1*28 with seven TA repeats).
 The ability to detect variants in other regions of the UGT1A1 gene is not part of this
 program.
- Survey PGX2 is designed for laboratories that provide *HLA-B*57:01* testing to identify risk of hypersensitivity to abacavir and *HLA-B*15:02* testing to identify risk of hypersensitivity to carbamazepine. The intended response is qualitative (presence/absence of the allele). This Survey is not appropriate for laboratories that perform molecular HLA typing. For HLA typing proficiency testing, please consult the HLA Molecular Typing (DML) Survey.

RBC Folate FOL			
Analyte Program Code Challenges/Shipment			
	FOL		
RBC folate	ı	2	

Program Information

- Three 25.0-µg extracted DNA specimens
- Includes allele detection (genotyping) and/or interpretive challenges
- Two shipments per year

- Two 2.0-mL lyophilized whole blood specimens
- Conventional and International System of Units (SI) reporting offered
- Three shipments per year

Renin and Aldosterone RAP						
Analyte	Program Code	Challenges/Shipment				
RAP						
Aldosterone	1	3				
Renin 3						

- Three 2.0-mL lyophilized plasma specimens
- Conventional and International System of Units (SI) reporting offered
- Two shipments per year



Tumor Markers TM, TMX				
Analyte	Program Code	Challenges/Shipment		
	TM,TMX			
Adrenocorticotropic hormone (ACTH)	ı	3		
Beta-2-microglobulin	ı	3		
CA 15-3	ı	3		
CA 19-9	ı	3		
CA 27.29	ı	3		
CA 72-4	ı	3		
CA 125	ı	3		
Calcitonin	ı	3		
Thyroglobulin	ı	3		

Program Information

- TM Three 2.0-mL liquid serum specimens
- TMX All Survey TM specimens in duplicate
- Two shipments per year

Human Epididymis Protein 4 HUEP						
Analyte	Program Code	Challenges/Shipment				
HUEP						
Human epididymis protein 4 ■ 3						

- Three 1.0-mL lyophilized serum specimens
- Two shipments per year

Improve the reliability of your patient results with CAP Survey Validated Materials

Use the same material that is sent in the Surveys program to:

- Identify and troubleshoot instrument/method problems
- Correlate results with other laboratories or instruments
- Document correction of problems identified in Surveys
- · Utilize material with confirmed results as an alternative external quality control
- · Identify potential proficiency testing failures

Each laboratory receives a Survey Participant Summary, which includes readily available results.

Endocrinology—Validated Materials

Validated Material	Program Code	Corresponding Survey	Page
Ligand—General	KVM	K	82
Ligand—Special	YVM	Υ	84

Program Information

- KVM Five 5.0-mL liquid serum specimens; three shipments per year
- YVM Six 5.0-mL liquid serum specimens in duplicate; two shipments per year

We are here to help. Fast Focus on Compliance—the inspector's quick guide

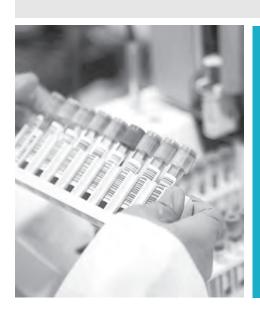
Multiply inspector confidence in just 10 minutes with the CAP's Fast Focus on Compliance mini-training vignettes. Using real world examples, inspectors will arm themselves with practical approaches to handle new and perplexing topics:

- Inspecting Personnel Records
- 12 Inspector Tools to Make Your Inspection Go More Smoothly
- Competency Assessment
- Identifying Systemic Issues—Critical Role of the Inspection Team Leader
- · Documenting Your Inspection Findings
- IQCP—What It Means to the Inspector

Access these concentrated topics online by searching Inspector Training at cap.org.

8

Blood Gas, Critical Care, and Oximetry



Our programs closely mimic patient testing to ensure accuracy in the laboratory.

- Test specimen levels that reflect clinical decision points.
- Keep current with the latest laboratory best practices with educational content supplied in our participant summary reports.
- Gain confidence in your results by comparing your performance against the largest peer groups.

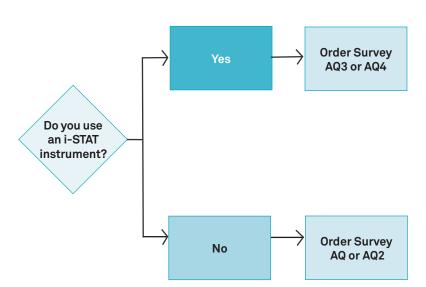
Blood Gas, Critical Care, and Oximetry

Analytes/procedures in bold type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Critical Care Blood Gas AQ, AQ2, AQ3, AQ4					
Analyte		Progra	m Code		Challenges/Shipment
	AQ	AQ2	AQ3	AQ4	
Calcium, ionized	•	ı			2
Chloride			1		5
Hematocrit		•			5
Hemoglobin, estimated	•				5
Lactate					2
Magnesium, ionized		•			2
PCO ₂		•		•	5
рН					5
PO ₂					5
Potassium	•				5
Sodium					5
tCO ₂		•			5
Creatinine		•			5
Glucose		•			5
Urea nitrogen (BUN)		ı		ı	5

For second instrument reporting options, see the Quality Cross Check programs, AQQ, AQ2Q, AQ3Q, and AQ4Q, on page 93.

- Five 2.5-mL aqueous specimens in duplicate and five 2.5-mL specimens for hematocrit testing in duplicate; appropriate for all methods except i-STAT®
- AQ3, AQ4 Five 2.5-mL specimens in duplicate for i-STAT methods only
- Conventional and International System of Units (SI) reporting offered
- Three shipments per year



Quality Cross Check—Blood Gas AQQ, AQ2Q, AQ3Q, AQ4Q					
Analyte		Progra	m Code		Challenges/Shipment
	AQQ	AQ2Q	AQ3Q	AQ4Q	
Calcium, ionized					3
Chloride	•				3
Hematocrit					3
Hemoglobin, estimated	•				3
Lactate					3
Magnesium, ionized					3
PCO ₂					3
рН					3
PO ₂					3
Potassium	•				3
Sodium					3
tCO ₂					3
Creatinine					3
Glucose					3
Urea nitrogen (BUN)					3

These programs do not meet regulatory requirements for proficiency testing; see Surveys AQ and AQ2-AQ4 on page 92. For additional information about the CAP Quality Cross Check program, see page 42.

The Quality Cross Check Program:

- Provides a solution for monitoring performance across multiple instruments, and is in compliance with the CMS directive regarding proficiency testing on multiple instruments.
- Simplifies instrument comparability efforts by providing custom reports with both peer group comparison and instrument comparability statistics.

- AQQ, AQ2Q Three 2.5-mL specimens in triplicate and three 2.5-mL specimens for hematocrit testing in triplicate; appropriate for all methods except i-STAT®
- AQ3Q, AQ4Q Three
 1.7-mL specimens in triplicate for i-STAT methods only
- Report up to three instruments
- Conventional and International System of Units (SI) reporting offered
- Two shipments per year

Blood Oximetry SO					
Analyte	Program Code	Challenges/Shipment			
	S0				
Carboxyhemoglobin	I	5			
Hematocrit, estimated	I	5			
Hemoglobin, total	I	5			
Methemoglobin	I	5			
Oxyhemoglobin	I	5			

Additional Information

- This Survey is not compatible with Oxicom-2000, -2100, or -3000 whole blood oximeters.
- For second instrument reporting options, see the Quality Cross Check program, SOQ, below.

Program Information

- Five 1.8-mL stabilized human hemoglobin solution specimens
- Conventional and International System of Units (SI) reporting offered
- Three shipments per year

Quality Cross Check—Blood Oximetry SOQ					
Analyte	Program Code	Challenges/Shipment			
	SOQ				
Carboxyhemoglobin		3			
Hematocrit, estimated	I	3			
Hemoglobin, total	I	3			
Methemoglobin		3			
Oxyhemoglobin 3					

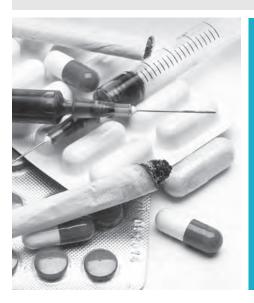
This program does not meet regulatory requirements for proficency testing; see Survey SO above. For additional information about the CAP Quality Cross Check program, see page 42.

The Quality Cross Check Program:

- Provides a solution for monitoring performance across multiple instruments, and is in compliance with the CMS directive regarding proficiency testing on multiple instruments.
- Simplifies instrument comparability efforts by providing custom reports with both peer group comparison and instrument comparability statistics.

- Three 1.2-mL liquid specimens in triplicate
- Report up to three instruments
- Conventional and International System of Units (SI) reporting offered
- Two shipments per year

9 Toxicology



Toxicology testing is changing at a rapid pace—so is our proficiency testing.

 The CAP's Toxicology Resource Committee annually reviews toxicology Survey compounds and modifies them in accordance with the appearance and prevalence of new drugs to stay contemporary.

New Programs NEW

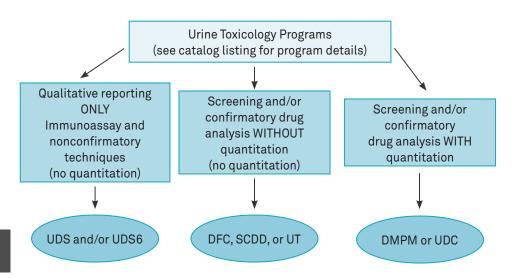
Trace Metals, Whole Blood (TMWB)	103
Toxicology Quality Program (TQP) Delayed until 2019	108
New Analyte/Drug Additions NEW	
Drug Monitoring for Pain Management (DMPM)	106
Oral Fluid for Drugs of Abuse (OFD)	100
Toxicology (T)	

Urine Toxicology (UT).......97

Toxicology

Analytes/procedures in **bold** type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Use this flowchart as a guide for ordering appropriate toxicology Surveys for your laboratory's testing menu.



-		
	Mixed and Other Matrices Toxicology Programs	
	(see catalog listing for program details)	
	\	
	Screening and/or confirmatory drug analysis with quantitation	
	•	
	T, FTC, OFD	

Toxicology T				
Analyte	Program Code	Challenges/Shipment		
	Т			
See drug listing on next page	ı	5		

Program Information

- A total of five specimens consisting of 20.0-mL liquid serum and 50.0-mL liquid urine specimens
- For laboratories performing qualitative and quantitative drug analysis on serum and qualitative analysis on urine specimens
- Three shipments per year

Urine Toxicology UT				
Analyte	Program Code	Challenges/Shipment		
	UT			
See drug listing on next page	I	5		

- Five 50.0-mL liquid urine specimens
- For laboratories performing qualitative drug analysis with qualitative confirmatory testing
- Three shipments per year

T and UT Programs Drug Listing

Challenges will include a mix of drugs from the list below.

6-acetylmorphine (6-AM)	Delta-9-THC (serum only)	Mephedrone	O-desmethyltramadol
7-aminoclonazepam	Delta-9-THC-COOH	Meprobamate	Olanzapine
7-aminoflunitrazepam	Desalkylflurazepam	Methadone	Opiate group
Acetaminophen	Desipramine	Methadone	Oxazepam
Alpha-hydroxyalprazolam	Desmethylclomipramine	metabolite (EDDP)	Oxycodone
Alprazolam	Desmethylcyclobenzaprine*	Methamphetamine	Oxymorphone
Amitriptyline	Desmethylsertraline	Methylenedioxy-	Paroxetine
Amphetamine	Dextromethorphan	amphetamine (MDA)	Pentobarbital
Amphetamine group	Diazepam	Methylenedioxy- methamphetamine	Phencyclidine
Aripiprazole NEW	Dihydrocodeine	(MDMA)	Phenethylamine
Atenolol	Diltiazem	Methylenedioxy-	Pheniramine
Atropine	Diphenhydramine	pyrovalerone (MDPV)	Phenobarbital
Barbiturate group	Doxepin	Methylphenidate	Phentermine
Benzodiazepine	Doxylamine	Metoprolol	Phenylephrine
group	Duloxetine	Mirtazapine	Phenytoin
Benzoylecgonine	Ecgonine ethyl ester	Morphine	Pregabalin NEW
Brompheniramine	Ecgonine methyl ester	N-desmethyltramadol	Propoxyphene
Buprenorphine	Ephedrine	Naproxen	Propranolol
Bupropion	Fentanyl	Nicotine	Pseudoephedrine
Butalbital	Flunitrazepam	Norbuprenorphine	Quetiapine
Cannabinoids	Fluoxetine	Norchlordiazepoxide	Quinidine
Carbamazepine	Flurazepam	Norclomipramine	Quinine
Carbamazepine-10,	Gabapentin NEW	Norcodeine	Ranitidine
11-epoxide	Hydrocodone	Norcyclobenzaprine*	Salicylates
Carisoprodol	Hydromorphone	Nordiazepam	Sertraline
Chlordiazepoxide	Hydroxyzine	Nordoxepin	Strychnine
Chlorpheniramine	Ibuprofen	Norfentanyl	Temazepam
Citalopram	Imipramine	Norfluoxetine	Topiramate NEW
Clomipramine	Ketamine	Norketamine	Tramadol
Clonazepam	Lamotrigine	Normeperidine	Trazodone
Clozapine	Levetiracetam	Noroxycodone	Tricyclic group
Cocaethylene	Lidocaine	Norpropoxyphene	Trimipramine
Cocaine	Lorazepam	Norsertraline	Valproic acid
Codeine	Lysergic acid diethylamide	Nortrimipramine	Venlafaxine
Cotinine	(LSD)	Nortriptyline	Verapamil
Cyclobenzaprine	Meperidine	Norverapamil	Zolpidem

^{*}Same compound

CAP/AACC Urine Drug Testing, Screening UDS, UDS6 **Analyte Program Code** Challenges/Shipment UDS6 UDS Limited Acetaminophen 5 3 5 Amphetamine 5 3 Amphetamine/methamphetamine group Barbiturate group 5 3 Benzodiazepine group 5 3 Benzoylecgonine/cocaine metabolites 5 3 5 3 Buprenorphine and metabolites Delta-9-THC-COOH 5 3 5 3 Ethanol Fentanyl 5 3 5 3 Lysergic acid diethylamide (LSD) 3 5 Methadone 5 3 Methadone metabolite (EDDP) Methamphetamine 5 3 5 3 Methaqualone 5 3 Methylenedioxymethamphetamine (MDMA) 5 3 Opiate group Oxycodone 5 3 5 3 Phencyclidine Propoxyphene 5 3 5 Tricyclic group 3

Program Information

- UDS Five 10.0-mL liquid urine specimens; three shipments per year
- UDS6 Three 10.0-mL liquid urine specimens; two shipments per year
- For laboratories performing drugs of abuse testing on urine specimens using immunoassay or other screening techniques only
- Participants will have access to the AACC quarterly newsletter, Clinical & Forensic Toxicology News



Urine Drug Adulterant/Integrity DAI		
Analyte	Program Code	Challenges/Shipment
	DAI	
Creatinine		3
Glutaraldehyde		3
Nitrite		3
Oxidants		3
рН		3
Specific gravity		3

- Three 25.0-mL urine specimens
- Two shipments per year

CAP/AACC Forensic Urine Drug Testing,

Confirmato	ory UDC	
Analyte	Program Code	Challenges/Shipment
	UDC	
6-acetylmorphine (6-AM)		10
Alpha-hydroxyalprazolam		10
Amphetamine		10
Benzoylecgonine		10
Buprenorphine		10
Butalbital		10
Codeine		10
Delta-9-THC-COOH		10
Hydrocodone		10
Hydromorphone		10
Lorazepam		10
Methadone		10
Methadone metabolite (EDDP)		10
Methamphetamine		10
Methaqualone		10
Methylenedioxyamphetamine (MDA)		10
Methylenedioxyethylamphetamine (MDEA)		10
Methylenedioxymethamphetamine (MDMA)		10
Morphine		10
Norbuprenorphine		10
Nordiazepam		10
Norpropoxyphene		10
Oxazepam		10
Oxycodone		10
Oxymorphone		10
Phencyclidine		10
Phenobarbital		10
Propoxyphene		10
Secobarbital		10
Temazepam		10
Adulterant/Integrity Indicator		
Creatinine		10
рН		10
Specific gravity		10

- Ten 50.0-mL liquid urine specimens
- For laboratories that perform both screening and confirmatory testing, including quantitation, for drugs of abuse in urine specimens; laboratories are asked to report creatinine, pH, and specific gravity for each specimen to ensure specimen adulteration has not occurred
- Participants will have access to the AACC quarterly newsletter, Clinical & Forensic Toxicology News
- Four shipments per year





Oral Fluid for Drugs of Abuse OFD		
Analyte	Program Code	Challenges/Shipment
	OFD	
Amphetamine Group		5
Amphetamine		5
Methamphetamine	I	5
Methylenedioxyamphetamine (MDA)		5
Methylenedioxyethylamphetamine (MDEA)	I	5
Methylenedioxymethamphetamine (MDMA)		5
Benzodiazepine Group		5
Alprazolam NEW		5
Diazepam		5
Nordiazepam		5
Oxazepam NEW		5
Temazepam		5
Buprenorphine NEW	ı	5
Buprenorphine and norbuprenorphine NEW		5
Cocaine and/or metabolite	I	5
Benzoylecgonine		5
Cocaine		5
Cannabinoids		5
Delta-9-THC		5
Delta-9-THC-COOH NEW		5
Methadone		5
Opiate Group		5
6-acetylmorphine (6-AM)	I	5
Codeine	•	5
Hydrocodone	I	5
Hydromorphone		5
Morphine		5
Oxycodone		5
Oxymorphone	I	5
Phencyclidine (PCP)		5

- Five 2.0-mL oral fluid specimens
- For laboratories performing drug screening, confirmation, and quantitation
- Four shipments per year



Vitreous Fluid, Postmortem VF		
Analyte	Program Code	Challenges/Shipment
	VF	
Acetone		3
Chloride		3
Creatinine		3
Ethanol		3
Glucose		3
Potassium		3
Sodium		3
Vitreous urea nitrogen		3

- Three 5.0-mL synthetic vitreous fluid specimens
- For forensic and other toxicology laboratories that perform quantitative analysis of vitreous fluid
- Conventional and International System of Units (SI) reporting offered
- Two shipments per year

Serum Drug Screening SDS			
Analyte	Program Code	Challenges/Shipment	
	SDS		
Acetaminophen, quantitative	•	3	
Acetone, semiquantitative and qualitative	•	3	
Barbiturate group, qualitative	1	3	
Benzodiazepine group, qualitative	1	3	
Salicylate, quantitative		3	
Total tricyclic antidepressants, qualitative		3	

Program Information

- Three 2.0-mL serum specimens
- For laboratories that perform serum drug screening using immunoassay or other screening techniques
- Two shipments per year



CAP/AACC Alcohol/Volatiles AL1,* AL2 **Program Code Analyte** Challenges/Shipment AL1* AL2 Whole Blood Serum 5 Acetone, quantitative П 5 Ethanol, quantitative П Ethylene glycol, qualitative and 5 quantitative Isopropanol, quantitative П 5 Methanol, quantitative 5

Program Information

 AL1 - Five 5.0-mL liquid whole blood specimens; conventional reporting



- AL2 Five 2.0-mL liquid serum specimens; conventional and International System of Units (SI) reporting offered
- Three shipments per year



^{*}The American Society of Crime Laboratory Directors/Laboratory Accreditation Board Proficiency Review Committee (ASCLD/LAB PRC) has approved Survey AL1.

Ethanol Biomarkers ETB		
Analyte	Program Code	Challenges/Shipment
	ETB	
Ethyl glucuronide (EtG), qualitative and quantitative	ı	3
Ethyl sulfate (EtS), quantitative	I	3

- Three 10.0-mL synthetic urine specimens
- Two shipments per year



CAP/AACC Blood Lead BL			
Analyte	Program Code	Challenges/Shipment	
	BL		
Lead	I	5	

This Survey meets the Occupational Safety and Health Administration (OSHA) requirements for proficiency testing [OSHA lead standards-29 CFR 1910.1025(j)(2)(iii)].

Program Information

- Five 6.0-mL liquid nonhuman whole blood specimens
- Conventional and International System of Units (SI) reporting offered
- Three shipments per year

 Three 6.0-mL whole blood specimens and three 13.0-mL urine specimens

International System of Units



Program Information

· Conventional and

(SI) reporting offeredSix shipments per year

Cadmium CD			
Analyte	Program Code	Challenges/Shipment	
	CD		
Beta-2-microglobulin, urine	I	3	
Cadmium, urine	ı	3	
Cadmium, whole blood	ı	3	
Creatinine, urine	I	3	

This Survey meets the Occupational Safety and Health Administration (OSHA) guidelines for proficiency testing (OSHA standard-29 CFR 1910.1027AppF).

- Three 25.0-mL urine specimens
- Designed for laboratories that qualitatively and/or quantitatively test for anabasine, cotinine, and/or nicotine in urine
- Two shipments per year

Nicotine and Tobacco Alkaloids NTA			
Analyte	Program Code	Challenges/Shipment	
	NTA		
Anabasine	I	3	
Cotinine	ı	3	
Nicotine		3	

Trace Metals, Urine TMU		
Analyte	Program Code	Challenges/Shipment
	TMU	
Aluminum	ı	2
Arsenic	ı	2
Chromium	ı	2
Cobalt	ı	2
Copper	ı	2
Lead	ı	2
Manganese	ı	2
Mercury	ı	2
Selenium	ı	2
Thallium	ı	2
Zinc	ı	2

- Two 25.0-mL urine specimens
- Conventional and International System of Units (SI) reporting offered
- For laboratories that monitor trace metals at normal and toxic levels
- Two shipments per year



Trace Me	tals, Whole Blood T	MWB NEW
Analyte	Program Code	Challenges/Shipment
	TMWB	
Aluminum	ı	3
Arsenic, total	ı	3
Chromium	ı	3
Cobalt	ı	3
Copper	I	3
Manganese	ı	3
Mercury	ı	3
Selenium	ı	3
Thallium	ı	3
Zinc	ı	3

- Three 6.0-mL whole blood specimens
- Conventional and International System of Units (SI) reporting offered
- Two shipments per year
- Designed for laboratories that monitor trace metals at normal and toxic levels

Forensic Toxicology, Criminalistics FTC		
Analyte	Program Code	Challenges/Shipment
	FTC	
See drug listing below	I	4

The American Society of Crime Laboratory Directors/Laboratory Accreditation Board Proficiency Review Committee (ASCLD/LAB PRC) has approved Survey FTC.

Program Information

- Three 20.0-mL whole blood specimens and one 20.0-mL synthetic urine specimen
- For crime and hospital laboratories that have forensic toxicology divisions performing qualitative and quantitative analysis of drugs in whole blood specimens along with a urine qualitative challenge
- Two shipments per year

FTC Program Drug Listing

Challenges will include a mix of drugs from the list below.

6-acetylmorphine (6-AM)	Ecgonine ethyl ester	Norfluoxetine
7-aminoclonazepam	Ecgonine methyl ester	Norketamine
7-aminoflunitrazepam	Ephedrine	Norpropoxyphene
Acetaminophen	Fentanyl*	Norsertraline
Alpha-hydroxyalprazolam	Fluoxetine	Nortriptyline
Alprazolam	Flurazepam*	Oxazepam
Amitriptyline	Gamma-hydroxybutyrate (GHB)	Oxycodone
Amphetamine	Hydrocodone	Oxymorphone
Benzoylecgonine	Hydromorphone	Paroxetine
Brompheniramine	Imipramine	Phencyclidine
Butalbital	Ketamine	Phenethylamine
Carisoprodol	Lorazepam	Phenobarbital
Chlorpheniramine	Lysergic acid diethylamide (LSD)	Phentermine
Clonazepam	Meperidine*	Phenytoin
Cocaethylene	Meprobamate	Propoxyphene
Cocaine	Methadone	Pseudoephedrine
Codeine	Methadone metabolite (EDDP)	Salicylate
Cyclobenzaprine*	Methamphetamine	Secobarbital
Delta-9-THC	Methylenedioxyamphetamine (MDA)	Sertraline
Delta-9-THC-COOH	Methylenedioxymethamphetamine	Temazepam
Desipramine	(MDMA)	Tramadol*
Desmethylcyclobenzaprine	Morphine*	Trazodone
Dextromethorphan	N-desmethyltramadol	Zolpidem
Diazepam	Nordiazepam	
Diphenhydramine	Nordoxepin	
Doxepin	Norfentanyl	*and/or metabolite(s)

Synthetic Cannabinoid/Designer Drugs SCDD		
Analyte	Program Code	Challenges/Shipment
	SCDD	
Synthetic cannabinoid/designer drugs		3

Additional Information

Synthetic cannabinoids and designer drug stimulants are widespread and constantly changing in respect to the available chemical moieties. In order to stay contemporary, the CAP has decided to modify the compounds in this program in accordance with the appearance and prevalence of new compounds.

Program Information

- Three 20.0-mL urine specimens
- For laboratories that perform screening and confirmatory testing for the compounds found in this program
- Two shipments per year



SCDD Program Drug Listing

Challenges will include a mix of drugs.

For the most current list of drugs, please go to cap.org. Click on Catalog and Shipping Information.

The list is located under the PT Order Supplements header.

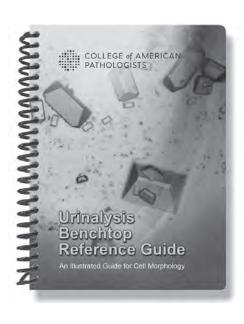
Urinalysis Benchtop Reference Guide (UABRG)

- Thirty-four different cell identifications, including common and rare cells
- Detailed descriptions for each cell morphology
- Eight tabbed sections for easy reference
- A durable and water-resistant format to withstand years of benchtop use—5" by 6½"

Choose code UABRG on your Surveys order form.

Or, view sample pages and order online:

- printed books at estore.cap.org
- · ebooks at ebooks.cap.org



Drug Monitoring for Pain Management DMPM		
Analyte	Program Code	Challenges/Shipment
	DMPM	
See drug listing below		3

- Three 40.0-mL urine specimens
- For laboratories offering qualitative, confirmatory, and/or quantitative urine drug analysis for pain management
- · Includes clinical cases and questions along with detailed descriptions of how to interpret test results
- Two shipments per year



DMPM Program Drug Listing

Challenges will include a mix of drugs from the list below.

Amphetamine group	Delta-9-THC-COOH	N-desmethyltramadol
6-acetylmorphine (6-AM)	Diazepam	Norbuprenorphine
7-aminoclonazepam	Fentanyl	Nordiazepam
Alpha-hydroxyalprazolam	Fentanyl and/or metabolites	Norfentanyl
Alprazolam	Gabapentin NEW	Normeperidine
Amphetamine	Hydrocodone	Noroxycodone
Barbiturate group	Hydromorphone	Noroxymorphone
Benzodiazepine group	Lorazepam	Norpropoxyphene
Benzoylecgonine	Lorazepam glucuronide	O-desmethyltramadol
Buprenorphine	Meperidine	Opiate group
Buprenorphine and/or metabolites	Meperidine and/or metabolites	Oxazepam
Butalbital	Meprobamate	Oxycodone
Cannabinoids	Methadone	Oxymorphone
Carisoprodol	Methadone metabolite (EDDP)	Pregabalin NEW
Carisoprodol and/or metabolites	Methamphetamine	Propoxyphene
Clonazepam	Methylenedioxyamphetamine (MDA)	Propoxyphene and/or metabolites
Cocaine	Methylenedioxymethamphetamine	Temazepam
Cocaine and/or metabolites	(MDMA)	Tramadol
Codeine	Morphine	Tramadol and/or metabolites

Drug-Facilitated Crime DFC		
Analyte	Program Code	Challenges/Shipment
	DFC	
See drug listing below		3

Program Information

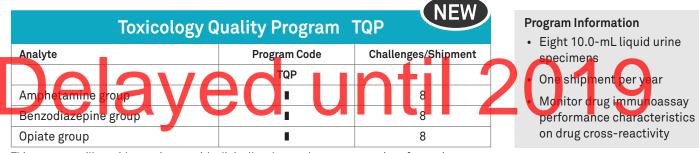
- Three 25.0-mL urine specimens
- For laboratories performing qualitative urine drug analysis with confirmation testing
- Designed for laboratories performing testing for drugs associated with drugfacilitated crimes, which target drugs at much lower concentrations than in other toxicology Surveys
- Two shipments per year



DFC Program Drug Listing

Challenges will include a mix of drugs from the list below.

/ hd	Fantand	Name and the line
4-hydroxytriazolam	Fentanyl	Norsertraline
7-aminoclonazepam	Fluoxetine	Nortriptyline
7-aminoflunitazepam	Gamma hydroxybutyrate (GHB)	Oxazepam
alpha-hydroxyalprazolam	Hydrocodone	Oxycodone
Amitriptyline	Hydromorphone	Oxymorphone
Amobarbital	Imipramine	Paroxetine
Amphetamine	Ketamine	Pentobarbital
Benzoylecgonine	Lorazepam	Phencyclidine (PCP)
Brompheniramine	Meperidine	Phenobarbital
Butalbital	Meprobamate	Phenytoin
Carisoprodol	Methadone	Propoxyphene
Chlorpheniramine	Methadone metabolite (EDDP)	Scopolamine
Citalopram/escitalopram	Methamphetamine	Secobarbital
Clonidine	Methylenedioxyamphetamine (MDA)	Sertraline
Codeine	Methylenedioxymethamphetamine	Temazepam
Cyclobenzaprine	(MDMA)	Tetrahydrozoline
Delta-9-THC-COOH	Morphine	Tramadol
Desipramine	Nordoxepin	Valproic Acid
Dextromethorphan	Norfluoxetine	Zaleplon
Diphenhydramine	Norketamine	Ziprasidone
Doxepin	Normeperidine	Zolpidem
Doxylamine	Norpropoxyphene	Zopiclone/Eszopiclone



This program will provide specimens with clinically relevant drug concentrations for testing across various assay manufacturers' kits, including Abbott, Beckman Coulter, Roche, Siemens, Ortho Clinical Diagnostics, and Thermo Fisher Scientific.

Improve the reliability of your patient results with CAP Survey Validated Materials

Use the same material that is sent in the Surveys program. Each laboratory receives a Survey Participant Summary, which includes readily available results.

Toxicology, Validated Material

Validated Material	Program Code	Corresponding Survey	Page
Urine Drug Testing, Screen	UDSM	UDS	98

Program Information

- Five 10.0-mL liquid urine specimens
- Three shipments per year

Find a practical guide to toxicology laboratory operations with this resource

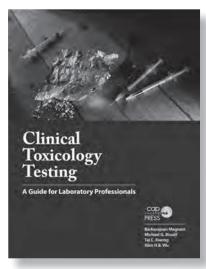
Clinical Toxicology Testing A Guide for Laboratory Professionals

Complex issues face the laboratory director or pathologist who offers toxicology services. This thorough reference book will guide both experienced physicians and those in training through the pharmacological principles, testing menus, and methodologies for toxicology testing.

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Item number: PUB220 Softcover; 304 pages; 2012



Be confident of the accuracy of your results.

- Specimens in the Accuracy-Based Surveys are matrix-effect free and have target values that are traceable to certified reference materials.
- Grading is independent of instrument peer groups, allowing health care systems to compare performance of multiple instrument systems across sites.

Accuracy-Based Programs

Accuracy-Based ProgramsValidated Materials	
New Analyte/Drug Additions NEW	
Accuracy-Based Testosterone, Estradiol (ABS)	111
Creatinine Accuracy CVL (LN24)	112

Accuracy-Based Programs

Analytes/procedures in **bold** type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Accuracy-Based Lipids ABL		
Analyte	Program Code	Challenges/Shipment
	ABL	
Apolipoprotein A1*		3
Apolipoprotein B*		3
Cholesterol*	I	3
HDL cholesterol*		3
Non-HDL cholesterol		3
LDL cholesterol		3
Lipoprotein (a)	I	3
Triglycerides*		3

^{*}This analyte will be evaluated against the reference method.

Program Information

- Three 1.0-mL human serum specimens
- Two shipments per year

Accuracy-Based Vitamin D ABVD		
Analyte	Program Code	Challenges/Shipment
	ABVD	
25-OH vitamin D (D2 and D3)		3

Additional Information

- The Centers for Disease Control and Prevention (CDC) will establish reference targets using isotope-dilution LC-MS/MS method.
- Specimens are collected by a modified application of Clinical Laboratory and Standards Institute Guideline CLSI C37-A, Preparation and Validation of Commutable Frozen Human Serum Pools as Secondary Reference Materials for Cholesterol Measurement Procedures; Approved Guideline.

- Three 1.0-mL liquid human serum specimens
- Serum is from multi-donor endogenous pools
- Conventional and International System of Units (SI) reporting offered
- Two shipments per year

Accuracy-Based Testosterone, Estradiol ABS			
Analyte	Program Code	Challenges/Shipment	
	ABS		
Albumin	ı	3	
Calcium	ı	3	
Cortisol	ı	3	
Estradiol	ı	3	
Follicle-stimulating hormone (FSH) NEW	ı	3	
Luteinizing hormone (LH) NEW	1	3	
Sex hormone-binding globulin (SHGB)	ı	3	
Testosterone	ı	3	
Testosterone, bioavailable		3	
Testosterone, free	ı	3	
Thyroid-stimulating hormone (TSH)	1	3	

Program Information

- Three 1.0-mL human serum specimens
- Two shipments per year

Additional Information

• The Centers for Disease Control and Prevention (CDC) will set target values for testosterone and estradiol using the established reference methods.

Accuracy-Based Urine ABU			
Analyte	Program Code	Challenges/Shipment	
	ABU		
Calcium	•	3	
Creatinine	I	3	
Urine albumin, quantitative	I	3	
Urine albumin: creatinine ratio	ı	3	

Target values for albumin are obtained by LC-MS/MS after trypsin digestion, performed by the Renal Testing Laboratory, Mayo Clinic, Rochester, MN, using calibration materials prepared from human serum albumin (>99% pure).

Other analytes will be compared by peer group for harmonization purposes.

- Three 5.0-mL human urine specimens
- Two shipments per year

Creatinine Accuracy Calibration Verification/Linearity LN24		
Analyte	Program Code	
	LN24	LN24 Target Range
Creatinine	ı	0.6-4.0 mg/dL
Estimated glomerular filtration rate (eGFR) NEW	ı	

LN Express service is available.

Additional Information

The College of American Pathologists (CAP) and the National Kidney Disease Education Program (NKDEP) have an initiative to harmonize clinically reported creatinine values. This initiative is analogous to what the federal health agencies and the clinical laboratory community did to improve the accuracy of cholesterol and glycohemoglobin testing.

Harmonized Thyroid ABTH Analyte **Program Code** Challenges/Shipment **ABTH** 3 T3, free (triiodothyronine, free) 3 T3, total (triiodothyronine, total) 3 Ī T4, free (thyroxine, free) 3 T4, total (thyroxine, total) П 3 Thyroid-stimulating hormone (TSH)

Additional Information

- Analytes will be evaluated using harmonization.
- Specimens are collected by a modified application of Clinical Laboratory and Standards Institute Guideline CLSI C37-A, Preparation and Validation of Commutable Frozen Human Serum Pools as Secondary Reference Materials for Cholesterol Measurement Procedures; Approved Guideline.

Hemoglobin A _{1c} Accuracy Calibration Verification/Linearity LN15		
Analyte	Program Code	
	LN15	LN15 Target Range
Hemoglobin A _{1c}		5%-12%

CAP-assigned target values derived from Hemoglobin A_{1c} measurements assayed by National Glycohemoglobin Standardization Program (NGSP) secondary reference laboratories.

LN Express service is available.

Program Information

- Six 1.0-mL human serum specimens
- Two shipments per year

Program Information

- Three 1.0-mL frozen human specimens
- Two shipments per year

- Six 0.8-mL liquid human whole blood specimens
- Two shipments per year

Hemoglobin A _{1c} GH2, GH5			
Analyte	Challenges/Shipment		
	Program Code		
	GH2	GH5	
Hemoglobin A _{1c}	3	5	

Additional Information

- These Surveys will be evaluated against the National Glycohemoglobin Standardization Program (NGSP) reference method.
- The College of American Pathologists Accreditation Program requires all accredited laboratories performing non-waived testing for Hemoglobin A_{1c} to complete 15 PT challenges per year.
- For second instrument reporting options, see the Quality Cross Check program, GHQ, on page 63.

Program Information

- GH2 Three 0.8-mL liquid human whole blood specimens; two shipments per year
- GH5 Five 0.8-mL liquid human whole blood specimens; three shipments per year

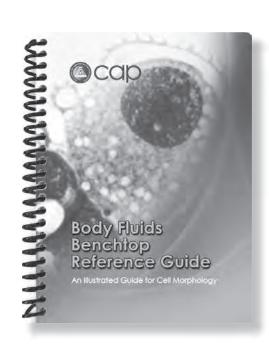
Body Fluids Benchtop Reference Guide (BFBRG)

- Thirty-six color images, including common and rare cells, crystals, and other cell inclusions
- Detailed descriptions of each cell including facts, cell morphology and inclusions
- Nine tabbed sections for easy reference
 - Erythroid Series
 - Lymphoid Series
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- A durable and water-resistant format to withstand years of benchtop use—5" x 6½"

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10

Validated Materials

Improve the reliability of your patient results with CAP Survey Validated Materials

Use the same material that is sent in the Surveys program to:

- · Identify and troubleshoot instrument/method problems
- Correlate results with other laboratories or instruments
- Document correction of problems identified in Surveys
- · Utilize material with confirmed results as an alternative external quality control
- Identify potential proficiency testing failures

Each laboratory receives a Survey Participant Summary, which includes readily available results.

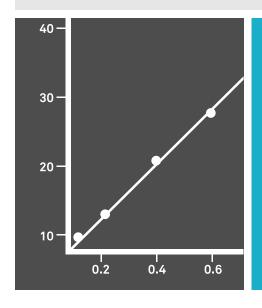
Chemistry, Validated Materials			
Validated Material	Validated Material Code	Corresponding Survey	Page
Chemistry and Therapeutic Drugs	CZVM	CZ	56-58
Cerebrospinal Fluid	MVM	M	74
Urine Chemistry—General	UVM	U	68

Coagulation—Limited, Validated Material				
Validated Material Code Corresponding Survey				
Coagulation—Limited CGM CGL				

Endocrinology—Validated Materials				
Validated Material Code Corresponding Survey Page				
Ligand Assay—General KVM K				
Ligand—Special YVM Y				

Toxicology, Validated Material				
Validated Material Code Corresponding Survey				
Urine Drug Testing, Screen	UDSM	UDS	98	

Instrumentation Validation Tools



Simplify your workflow with the most comprehensive solution for calibration verification and linearity.

- Ensure confidence in your linearity results with 5–8 specimen levels included with each CVL program.
- Receive a customized report with the most rigorous statistical analysis.
- Expedite linearity evaluations through our *LN Express*SM Service.

Instrumentation Validation Tools

Calibration Verification/Linearity	
New Analyte/Drug Additions NEW	
Creatinine Accuracy CVL (LN24) Estimated glomerular filtration rate (eGFR)	125
Program Changes	
High-Sensitivity C-Reactive Protein CVL (LN21)	124

Discontinued Programs

New target range

Whole Blood Ethanol Calibration Verification/Linearity (LN14)
Calibration Verification/Linearity, Validated Materials (LN2VM, LN2VM1, LDM, LLM, LUM)

11

Calibration Verification/Linearity

The CAP CVL program

Our program will help you meet CLIA regulations and CAP Laboratory Accreditation Program requirements for calibration verification and analytical measurement range (AMR) verification under 42 CFR493.1255(b)(3) for most analytes. In addition, you will receive a linearity assessment to help identify instrument/method performance issues before they can affect your patient results.

With your enrollment in the CAP CVL program you will receive:

· Testing Kit

- Kit instructions—Contain important information to help you complete testing and accurately report your results
- Result form
- Specimens—The majority of CAP CVL programs offer human-based materials to closely mimic your patient

Customized Report Package

- Executive Summary—A quick overview of both your calibration verification and linearity results for all reported analytes
- Calibration Verification Evaluation
- Linearity Evaluation
 - Receive your linearity evaluations through LN Express[™], our expedited delivery service, within two business days for select CVL programs by logging in to e-LAB Solutions Suite
- Linearity Troubleshooting Report
- Participant Summary—A summary of laboratory performance that includes peer group statistics and enhanced diagnostic information for early insight into potential problems

Additional Tools

- o Calibration Verification/Linearity Program User's Guide—Get assistance in interpreting your evaluations and reports as well as helpful troubleshooting information with suggested actions. Also available online by logging in to e-LAB Solutions Suite
- Calibration Verification Troubleshooting Guide—The guide provides suggested actions if you receive a calibration verification result of Different, or if your evaluation result is Verified over a range that does not include all of your reported results
- o Calibration Verification/Linearity Surveys Investigation Checklist for Problematic Results—Interpretative checklists are included to help with troubleshooting and documentation

CVL Program	Page No.	Corresponding Proficiency Testing Survey	Page No.
LN2 - Chemistry, Lipid, Enzyme CVL	118	C1, C3/C3X, C4,	
LN2BV - Chemistry, Lipid, Enzyme all Beckman (except AU), Vitros CVL	118	CZ/CZX/CZ2X	56-58
LN3 - Therapeutic Drug Monitoring CVL	119	CZ/CZX/CZ2X/Z	56-58
LN5 - Ligand CVL	119-120		
LN5S - Ligand all Siemens ADVIA (Centaur, CP, and XP) CVL	119-120	K/KK	82
LN6 - Urine Chemistry CVL	120	U	68
LN7 - Immunology CVL	121	IG/IGX	202
LN8 - Reproductive Endocrinology CVL	121	Y/YY	84
LN9 - Hematology CVL	121	FH series, HE series	135, 134
LN11 - Serum Ethanol CVL	122	AL2	101
LN12 - C-Reactive Protein CVL	122	CDD	202
LN12E - C-Reactive Protein, Extended CVL	122	CRP	202
LN13, LN13C - Blood Gas/Critical Care CVL	122-123	AQ, AQ2, AQ3, AQ4	92
LN15 - Hemoglobin A _{1c} Accuracy CVL	123	GH2, GH5	63
LN16 - Homocysteine CVL	123	HMS	64
LN17 - Whole Blood Glucose CVL	123		
LN18, LN19 - Reticulocyte CVL	124	RT, RT2, RT3, RT4	139
LN20 - Urine Albumin CVL	124	U	68
LN21 - High-Sensitivity C-Reactive Protein CVL	124	HSCRP	64
LN22 - Flow Cytometry CVL	124	FL	209
LN23 - Prostate-Specific Antigen CVL	125	K/KK	82
LN24 - Creatinine Accuracy CVL	125	C1, C3/C3X, C4, CZ/CZX/CZ2X	56-58
LN25, LN27 - Troponin I and T CVL	125	CRT, CRTI, TNT	62
LN30 - B-Type Natriuretic Peptides CVL	125	BNP	61
LN31 - Immunosuppressive Drugs CVL	126	CS	59
LN32 - Ammonia CVL	126	C1, C3/C3X, CZ/CZX/CZ2X	56-58
LN33 - Serum Myoglobin CVL	126	CRT, CRTI	62
LN34 - Tumor Markers CVL	126	TM/TMX	89
LN35 - Thrombophilia CVL	127	CGS2	160
LN36 - Heparin CVL	127	CGS4	160
LN37 - von Willebrand Factor Antigen CVL	127	CGS3	160
LN38 - CMV Viral Load CVL	127	VLS, VLS2	193
LN39 - HIV Viral Load CVL	127	HIVG, HV2	193
LN40 - Vitamin D CVL	127	VITD	84
LN41 - Procalcitonin CVL	128	PCT	77
LN42 - D-Dimer CVL	128	CGL, CGDF	158
LN43 - Lamellar Body Count CVL	128	LBC	149
LN44 - Fibrinogen CVL	128	CGL	158
LN45 - HCV Viral Load CVL	127	HCV2	193

All CVL Surveys provide individual evaluation reports by analytes, an Executive Summary, and graphical plots for linearity and calibration verification.

Chemistry, Lipid, Enzyme Calibration Verification/Linearity LN2, LN2BV					
Analyte	Program Code	LN2	LN2 LN2BV		Units
	LN2, LN2BV	(All Instruments)	All Beckman (except AU)	Vitros	
Albumin			1.5-9.0		g/dL
Calcium			4.0-18.0		
Chloride			60-180		mmol/L
CO ₂			7–40		mmol/L
Creatinine			0.3-32.0		mg/dL
Glucose			20-780		mg/dL
Iron			10-950		µg/dL
Magnesium			0.3-10.0		mg/dL
Osmolality			200-600		m0sm/kg H ₂ 0
Phosphorus			0.5-20.0		mg/dL
Potassium			1.5-13.0		mmol/L
Protein			1.5-10.0		g/dL
Sodium			90-215		mmol/L
Urea nitrogen/Urea			3-190		mg/dL
Uric acid			1-25		mg/dL
Alkaline phosphatase		25-1,800	25-1,000	25-1,100	U/L
ALT (SGPT)		10-900	10-650	30-700	U/L
Amylase		30-1,800	30-900	30-800	U/L
AST (SGOT)		10-900	10-500	10-700	U/L
Creatine kinase		25-2,000	25-1,200	25-700	U/L
CK-2 (MB) Mass		1–250	1-300	1–200	ng/mL
Gamma glutamyl transferase	•	10-1,400	10-900	10-1,100	U/L
Lactate dehydrogenase		50-1,800	50-700	185-3,000	U/L
Lipase		20-1,400	20-190	150-2,500	U/L
Bilirubin, direct		0.1–10.0			mg/dL
Bilirubin, total		0.2-25.0			mg/dL
Cholesterol		35-625			mg/dL
HDL			7–120		mg/dL
Triglycerides			20-700		mg/dL

• Seven 5.0-mL liquid serum specimens for basic chemistry, six 3.0-mL liquid serum specimens for direct and total bilirubin, seven 2.0-mL liquid serum specimens for lipids, and seven 5.0-mL liquid serum specimens for enzymes

Program Information

- LN2 Appropriate for most major instruments
- LN2BV Appropriate for Beckman (except AU) and Vitros instruments only
- Two shipments per year

LN Express service is available.

Please note that the ranges listed are an estimate of the values recovered. Some instruments may recover lower or higher values than the ranges listed.

Therapeutic Drug Monitoring Calibration Verification/Linearity LN3			
Analyte	Program Code		
	LN3	LN3 Target Ranges	
Acetaminophen	1	20-450 μg/mL	
Amikacin	ı	2-45 μg/mL	
Carbamazepine	ı	2–18 μg/mL	
Digoxin	ı	0.5-4.4 ng/mL	
Gentamicin	I	1–11 μg/mL	
Lidocaine	I	1–10 μg/mL	
Lithium	I	0.3-4.0 mmol/L	
N-acetylprocainamide (NAPA)	I	2-25 μg/mL	
Phenobarbital	I	8-70 μg/mL	
Phenytoin	I	5–35 μg/mL	
Primidone	I	1–22 μg/mL	
Procainamide	I	2–18 μg/mL	
Quinidine	I	0.4-7.0 μg/mL	
Salicylate	I	7-90 mg/dL	
Theophylline	ı	5–35 μg/mL	
Tobramycin	ı	1–12 μg/mL	
Valproic acid	ı	15-140 μg/mL	
Vancomycin	ı	7–90 μg/mL	

LN Express service is available.

Ligand Calibration Verification/Linearity LN5, LN5S **Target Ranges** Analyte **Program Code** LN5, LN5S* **LN5 Target Ranges LN5S Target Ranges** AFP 1.0-900.0 ng/mL CEA П 0.5-750.0 ng/mL 0.6-90.0 ng/mL Cortisol $1-65 \mu g/dL$ П Ferritin 2-1,100 ng/mL Folate П 1.3-20 ng/mL Human chorionic 5-14,000 mIU/mL gonadotropin (hCG) T3, total (triidothyronine) П 0.5 - 7.0 ng/mLT4, total (thyroxine) $1-80 \mu g/dL$ Continued on the next page

Program Information

- Six 4.0-mL liquid serum specimens
- A seventh 4.0-mL liquid serum specimen for acetaminophen and vancomycin
- Two shipments per year

Program Information

- LN5 Eight 4.0-mL liquid serum specimens; appropriate for most major instruments except Siemens ADVIA Centaur, XP, and CP users
- LN5S Thirteen 4.0-mL liquid serum specimens; appropriate for Siemens ADVIA Centaur, XP, and CP users
- Two shipments per year

Please note that the ranges listed are an estimate of the values recovered. Some instruments may recover lower or higher values than the ranges listed.

Ligand Calibration Verification/Linearity LN5, LN5S continued					
Analyte	Program Code Target Ranges				
	LN5, LN5S*	LN5, LN5S* LN5 Target Ranges LN5S Target Ranges			
Thyroid-stimulating hormone (TSH)		0.01–100 μIU/mL			
Vitamin B ₁₂	■ 100−2,200 pg/mL				

^{*}The LN5S CVL will allow Siemens ADVIA Centaur users to report other major instruments for analytes other than CEA, if needed.

Program Information

- LN5 Eight 4.0-mL liquid serum specimens; appropriate for most major instruments except Siemens ADVIA Centaur, XP, and CP users
- LN5S Thirteen 4.0-mL liquid serum specimens; appropriate for Siemens ADVIA Centaur, XP, and CP users
- Two shipments per year

Urine Chemistry Calibration Verification/Linearity LN6

Analyte	Program Code	
	LN6	LN6 Target Ranges
Amylase	I	40-1,500 U/L
Calcium	I	5–30 mg/dL
Chloride	I	20-330 mmol/L
Creatinine	I	20-460 mg/dL
Glucose	I	25-640 mg/dL
Osmolality	I	30 –1,800 m0sm/kg H ₂ 0
Phosphorus	I	15-200 mg/dL
Potassium	I	7–225 mmol/L
Protein, total	I	10-235 mg/dL
Sodium	I	20-340 mmol/L
Urea nitrogen/Urea	I	20-2,000 mg/dL
Uric acid	I	6-150 mg/dL

LN Express service is available.

Program Information

- Eighteen 4.0-mL liquid urine specimens
- Two shipments per year

Please note that the ranges listed are an estimate of the values recovered. Some instruments may recover lower or higher values than the ranges listed.

LN Express service is available.

Immunology Calibration Verification/Linearity LN7			
Analyte	Program Code		
	LN7	LN7 Target Ranges	
Alpha-1-antitrypsin	1	25-616 mg/dL	
Complement C3	ı	21-420 mg/dL	
Complement C4	1	5-100 mg/dL	
IgA	ı	32-650 mg/dL	
IgG	ı	150-3,000 mg/dL	
IgM	I	25-450 mg/dL	
Transferrin	I	38-950 mg/dL	

LN Express service is available.

Reproductive Endocrinology Calibration Verification/Linearity LN8 **Analyte Program Code** LN8 **LN8 Target Ranges** Estradiol 25-4,500 pg/mL Follicle-stimulating hormone (FSH) 3-190 mIU/mL Human chorionic gonadotropin (hCG) 5-8,000 mIU/mL Luteinizing hormone (LH) 2-190 mIU/mL Progesterone 1-50 ng/mL Prolactin 3-315 ng/mL Testosterone 20-1,500 ng/dL

Hematology Calibration Verification/Linearity LN9 Analyte **Program Code** LN9 **LN9 Target Ranges** Hemoglobin 1.5-24.0 g/dL Platelet count $10-2,500 \times 10^9/L$ **RBC** count $0.5 - 8.00 \times 10^{12}/L$ WBC count 0.5-350.0 x 109/L

Program Information

- Six 2.0-mL liquid serum specimens
- Two shipments per year

Program Information

- Seven 4.0-mL liquid serum specimens
- Two shipments per year

Program Information

- Twenty 3.0-mL liquid specimens
- Two shipments per year

Please note that the ranges listed are an estimate of the values recovered. Some instruments may recover lower or higher values than the ranges listed.

LN Express service is available.

LN Express service is available.

Serum Ethanol Calibration Verification/Linearity LN11				
Analyte Program Code				
LN11 LN11 Target Range				
Serum ethanol	ı	15-550 mg/dL		

LN Express service is available.

Program Information

- Seven 3.0-mL liquid serum specimens
- Two shipments per year

C-Reactive Protein; C-Reactive Protein, Extended Calibration Verification/Linearity LN12, LN12E

Analyte	Program Code		Program Code	
	LN12	LN12 Target Range	LN12E	LN12E Target Range
C-reactive protein		5-110 mg/L		6-320 mg/L

LN Express service is available.

Not appropriate for reporting high-sensitivity C-reactive protein (hsCRP). For reporting hsCRP, use LN21 on page 124.

Program Information

- LN12 Five 1.0-mL liquid serum specimens; appropriate for Beckman Immage, Siemens Dimension, and Vitros instruments
- LN12E Six 1.0-mL liquid serum specimens; appropriate for Abbott Architect, Beckman (except Immage), Roche, and Siemens (except Dimension) instruments
- · Select program based on appropriate target range for assay used
- · Two shipments per year

Blood Gas/Critical Care Calibration Verification/Linearity LN13, LN13C

Analyte	Program Code		Program Code	
	LN13	LN13 Target Ranges	LN13C	LN13C Target Ranges
PCO ₂		12-91 mm Hg		12–91 mm Hg
рН		6.83-7.82		6.83-7.82
PO ₂		18-490 mm Hg		18-490 mm Hg
Calcium, ionized				0.15-3.3 mmol/L
Chloride				62–148 mmol/L
Glucose				10-465 mg/dL
Lactate				0.2–18 mmol/L
Continued on the next pa	age		'	

Program Information

- Ten 2.5-mL ampules of aqueous specimens
- · Two shipments per year

Please note that the ranges listed are an estimate of the values recovered. Some instruments may recover lower or higher values than the ranges listed.

Blood Gas/Critical Care Calibration Verification/Linearity LN13, LN13C continued				
Analyte	Program Code		Program Code	
	LN13	LN13 Target Ranges	LN13C	LN13C Target Ranges
Potassium				0.5-10.7 mmol/L
Sodium				83–172 mmol/L

Program Information

- Ten 2.5-mL ampules of aqueous specimens
- · Two shipments per year

Hemoglobin A _{1c} Accuracy Calibration Verification/Linearity LN15				
Analyte	Program Code			
LN15 LN15 Target Range				
Hemoglobin A _{1c}	ı	5%-12%		

CAP-assigned target values derived from Hemoglobin A_{1c} measurements assayed by National Glycohemoglobin Standardization Program (NGSP) secondary reference laboratories.

LN Express service is available.

Program Information

Program Information
Six 0.8-mL liquid human whole blood specimens
Two shipments per year

- Six 1.0-mL liquid serum specimens
- Two shipments per year

Homocysteine Calibration Verification/Linearity LN16			
Analyte	Program Code		
	LN16	LN16 Target Range	
Homocysteine		5–65 μmol/L	

LN Express service is available.

Whole Blood Glucose Calibration Verification/Linearity LN17 Analyte Program Code LN17 LN17 Target Range Whole blood glucose ■ 50-400 mg/dL

LN Express service is available.

Program Information

- Five 2.0-mL liquid whole blood specimens
- Report up to 10 different ancillary testing sites or instruments
- Two shipments per year

Please note that the ranges listed are an estimate of the values recovered. Some instruments may recover lower or higher values than the ranges listed.

Instrumentation Validation Tools

Reticulocyte Calibration Verification/Linearity LN18, LN19				
Instrument/Method	Program Code		Program Code	
	LN18	LN18 Target Range	LN19	LN19 Target Range
Coulter Gen∙S™, LH 500, LH 700 series, and UniCel DxH				0.3%-27.0%
All other instruments		0.3%-24.0%		

LN Express service is available.

Program Information

- LN18 Five 2.5-mL liquid whole blood specimens with pierceable caps
- LN19 Five 3.0-mL liquid whole blood cell specimens with pierceable caps
- Two shipments per year

Urine Albumin Calibration Verification/Linearity LN20

Analyte	Program Code	
	LN20	LN20 Target Ranges
Urine albumin	I	10-350 mg/L
Urine creatinine	ı	20-500 mg/dL

Program Information

- Six 5.0-mL urine specimens
- Two shipments per year

High-Sensitivity C-Reactive Protein Calibration Verification/Linearity LN21

Analyte	Program Code	
	LN21	LN21 Target Range
High-sensitivity C-reactive protein	ı	0.5-18.0 mg/L

LN Express service is available.

Program Information

- Six 1.0-mL liquid serum specimens
- For high-sensitivity methods
- Two shipments per year

Flow Cytometry Calibration Verification/Linearity LN22

Analyte	Program Code	
	LN22	LN22 Target Ranges
CD3+	ı	50%-70% positive
CD3+ T lymphocytes absolute	ı	350–4,000 cells/μL
CD3+/CD4+	ı	1%-40% positive
CD3+/CD4+ T lymphocytes absolute	1	6–2,000 cells/μL
CD3+/CD8+	ı	25%-40% positive
CD3+/CD8+ T lymphocytes absolute	ı	250–1,600 cells/μL

Program Information

- · Seven 1.0-mL liquid whole blood specimens
- Two shipments per year

Please note that the ranges listed are an estimate of the values recovered. Some instruments may recover lower or higher values than the ranges listed.

Prostate-Specific Antigen Calibration Verification/Linearity LN23			
Analyte	Program Code		
LN23 LN23 Target Rang			
Prostate-specific antigen		0.1-90.0 ng/mL	

Creatinine Accuracy Calibration Verification/Linearity LN24			
Analyte	Program Code		
	LN24	LN24 Target Range	
Creatinine		0.6-4.0 mg/dL	
Estimated glomerular filtration rate (eGFR) NEW			

LN Express service is available.

Additional Information

The College of American Pathologists (CAP) and the National Kidney Disease Education Program (NKDEP) have an initiative to harmonize clinically reported creatinine values. This initiative is analogous to what the federal health agencies and the clinical laboratory community did to improve the accuracy of cholesterol and glycohemoglobin testing.

Troponin Calibration Verification/Linearity LN25, LN27				
Analyte	Program Code Program Code			
	LN25	LN25 Target Range	LN27	LN27 Target Range
Troponin I	•	0.05-60.00 ng/mL		
Troponin T				0.1-27.00 ng/mL

B-Type Natriuretic Peptides Calibration Verification/Linearity LN30			
Analyte	Program Code		
	LN30	LN30 Target Ranges	
BNP	•	30-6,500 pg/mL	
NT-proBNP	I	50-50,000 pg/mL	

LN Express service is available.

Program Information

- Twelve 1.0-mL liquid serum specimens
- Two shipments per year

Program Information

- Six 1.0-mL human serum specimens
- Two shipments per year

Program Information

- LN25 Seven 2.0-mL liquid serum specimens
- LN27 Six 2.0-mL liquid serum specimens
- Two shipments per year

Program Information

- Seven 1.0-mL liquid plasma specimens
- · Two shipments per year

Please note that the ranges listed are an estimate of the values recovered. Some instruments may recover lower or higher values than the ranges listed.

11

Instrumentation Validation Tools

Immunosuppressive Drugs Calibration Verification/Linearity LN31					
Analyte Program Code					
LN31 LN31 Target Ranges					
Cyclosporine	60-1,200 ng/mL				
Tacrolimus	I	1.5-30.0 ng/mL			

Program Information

- Seven 2.0-mL liquid whole blood hemolysate specimens
- Two shipments per year

Ammonia Calibration Verification/Linearity LN32						
Analyte Program Code						
LN32 LN32 Target Range						
mmonia ■ 13-900 μmol/L						

LN Express service is available.

Program Information

- Seven 2.0-mL liquid serum specimens
- · Two shipments per year

Serum Myoglobin Calibration Verification/Linearity **LN33** Analyte **Program Code** LN33 **LN33 Target Range** Myoglobin Ī 25-900 ng/mL

LN Express service is available.

Program Information

- Seven 1.0-mL liquid serum specimens
- Two shipments per year

Tumor Markers Calibration Verification/Linearity LN34					
Analyte Program Code					
	LN34	LN34 Target Ranges			
CA 125	I	1–1,000 U/mL 2–190 U/mL			
CA 15-3	I				
CA 19-9	■ 10-900 U/mL				

LN Express service is available.

Program Information

- Seven 3.0-mL liquid serum specimens
- Two shipments per year

Please note that the ranges listed are an estimate of the values recovered. Some instruments may recover lower or higher values than the ranges listed.

Coagulation Calibration Verification/Linearity LN35, LN36, LN37					
Analyte	Pr	ogram Co	de		
	LN35	LN35 LN36 LN37		Target Ranges	
Antithrombin activity	•			10%-130%	
Protein C activity	•			10%-100%	
Heparin, low molecular weight				0.1-2.0 U/mL	
Heparin, unfractionated				0.1-1.3 U/mL	
von Willebrand factor antigen				5%-140%	

The LN35, LN36, and LN37 CVL programs meet the CAP Accreditation requirements HEM.38009, 38010, and 38011.

LN Express service is available.

Program Information

- LN35, LN37 Six 1.0-mL frozen plasma specimens per mailing
- LN36 Twelve 1.0-mL frozen plasma specimens per mailing, which include six for low molecular weight heparin and six for unfractionated heparin
- Two shipments per year; ships on dry ice

Viral Load Calibration Verification/Linearity LN38, LN39, LN45

Analyte	Program Code		de			
	LN38* LN39 LN45		LN45	Target Ranges		
CMV viral load				316-1.0M IU/mL		
HIV viral load				50-5.0M IU/mL		
HCV viral load				50-280M IU/mL		

^{*}The biohazard warning applies to Survey LN38.

LN Express service is available.

Program Information

- LN38 Six 1.5-mL frozen plasma specimens
- Two shipments per year; ships on dry ice



- LN39 Six 2.5-mL plasma specimens
- LN45 Seven 2.5-mL frozen plasma specimens
- Two shipments per year; ships on dry ice (dry ice does not apply to LN39)

Vitamin D Calibration Verification/Linearity LN40

Analyte	Program Code	
	LN40	Target Range
25-OH vitamin D, total		4-120 ng/mL

LN Express service is available.

Program Information

- Six 1.0-mL serum specimens
- Two shipments per year



Refer to the Ordering Information provided for information regarding additional dangerous goods and related fees.

Please note that the ranges listed are an estimate of the values recovered. Some instruments may recover lower or higher values than the ranges listed.

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Procalcitonin Calibration Verification/Linearity LN41						
Analyte	Analyte Program Code					
LN41 Target Range						
Procalcitonin ■ 0.3-190 ng/mL						

LN Express service is available.

Program Information

- Six 1.0-mL frozen serum specimens
- Two shipments per year; ships on dry ice

D-Dimer Calibration Verification/Linearity LN42						
Analyte	Analyte Program Code					
LN42 Target Range						
D-dimer ■ 220–5,500 ng/mL FEU						

LN Express service is available.

Program Information

- Six 1.0-mL plasma specimens
- Two shipments per year



Lamellar Body Count Calibration Verification/Linearity LN43					
Analyte Program Code					
LN43 Target Range					
Lamellar body count ■ 5–200 particles x 10 ⁹ /L					

LN Express service is available.

Program Information

- Six 2.0-mL simulated liquid amniotic fluid specimens
- For use with lamellar body count methods performed on hematology analyzers
- Two shipments per year

Fibrinogen Calibration Verification/Linearity LN44					
Analyte	Program Code				
LN44		Target Range			
Fibrinogen ■ 80-900 mg/dL					
	I				

LN Express service is available.

Program Information

- Six 1.0-mL frozen plasma specimens
- Two shipments per year; ships on dry ice

Please note that the ranges listed are an estimate of the values recovered. Some instruments may recover lower or higher values than the ranges listed.

Instrumentation Quality Management Programs

Instrumentation I					
Challenges	Program Code				
	I				
	A Shipment	B Shipment	C Shipment		
Adjustable micropipette calibration verification/linearity	•				
Analytical balance check			ı		
Gravimetric pipette calibration	•				
Microtiter plate linearity					
Refractometer calibration					
Spectrophotometer (stray light check)	•				
Absorbance check – UV wavelength					
Fluorescent intensity check – fluorescent microscopes		ı			
Ocular micrometer calibration		•			
Osmometer study		•			
Peak absorbance measurement					
pH meter check					
Photometric calibration – visible wavelength		•			

WARNING: The Instrumentation (I) Survey specimens may contain corrosive or toxic substances, environmental hazards, or irritants.

- Designed to assess instruments not routinely challenged during the proficiency testing process
- Includes appropriate materials to assess important functional parameters, including accuracy and linearity
- Three shipments per year

Interfering Substance IFS					
Analyte	Program Code				
	IFS				
	Bilirubin Interferent	Hemoglobin Interferent	Lipid Interferent		
Alanine aminotransferase (ALT/SGPT)	ı		ı		
Albumin	ı	•	ı		
Alkaline phosphatase	ı	I	ı		
Amylase	ı		ı		
Aspartate aminotransferase (AST/SGOT)	ı	•	ı		
Calcium	ı	•	ı		
Chloride	ı	•	ı		
CK2 (MB) mass	ı		ı		
Creatine kinase (CK)	ı		ı		
Creatinine	ı		ı		
Gamma glutamyl transferase (GGT)	ı		ı		
Glucose	ı	•	ı		
Iron	ı	•	ı		
Lactate dehydrogenase (LD)	ı		ı		
Lipase	ı	•	ı		
Magnesium	ı	•	ı		
Osmolality	ı		ı		
Phosphorus	ı	•	ı		
Potassium	•		ı		
Protein, total	ı	I	ı		
Sodium	ı	I	ı		
Urea nitrogen (BUN)	•		ı		
Uric acid			ı		

The material expires December 1, 2018.

- Eighteen 10.0-mL liquid serum specimens
- Designed for verifiying manufacturing interference specifications and investigating discrepant results caused by interfering substances
- Submit results any time prior to the material's expiration date
- One shipment per year

Serum Carryover SCO				
Analyte Program Code				
	SC0			
Creatinine	ı			
hCG				
Lactate dehydrogenase (LD)				
Phenytoin	ı			

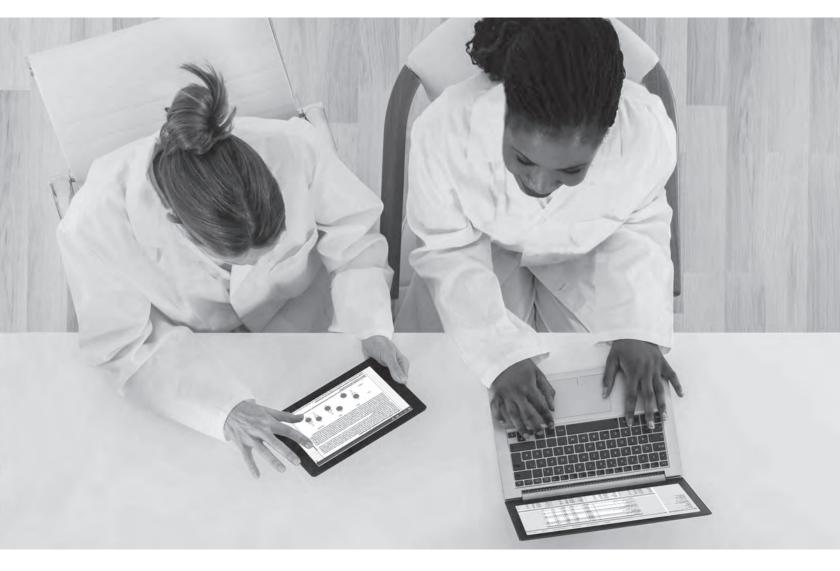
Urine Toxicology Carryover UTCO					
Analyte	Program Code				
	UTCO				
Benzoylecgonine	ı				
Delta-9-THC-COOH	ı				
Opiates	ı				
Amphetamine					

Program Information

- One 10.0-mL liquid serum specimen (low level) and one 5.0-mL liquid serum specimen (high level)
- Designed to screen for instrument sample probe carryover
- One shipment per year

- Two 40.0-mL urine specimens (low and high levels)
- Designed to screen for instrument sample probe carryover
- One shipment per year

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12 Hematology and Clinical Microscopy



Our programs are supported by 500 experts in laboratory medicine.

These experts spend countless hours monitoring testing trends to:

- Determine specimen specifications to challenge participants.
- Keep our offerings contemporary with new analytes and programs.
- Provide peer-reviewed CME, SAM, and CE education.

Hematology and Clinical Microscopy

Hematology
New Programs NEW
Quality Cross Check—Reticulocyte Series (RTQ, RT2Q, RT3Q, RT4Q)140
New Analyte Additions NEW
Clinical Microscopy Miscellaneous Photopage (CMMP)149 Spermatozoa

Urinalysis and Clinical Microscopy (CMP2, CMP3)

Discontinued Programs

Hematology

Analytes/procedures in **bold** type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Hematology—Basic HE, HEP						
Analyte/Procedure	Progra	Program Code Challenge				
	HE	HEP				
Blood cell identification			10			
Hematocrit	1 1		5			
Hemoglobin			5			
MCV, MCH, and MCHC	1 1		5			
MPV	1 1		5			
Platelet count	I I		5			
RDW	1 1		5			
Red blood cell count			5			
White blood cell count			5			

Program Information

- Five 3.0-mL whole blood specimens
- HEP Ten images, each available as photographs and online images
- Three shipments per year



Blood Cell Identification, Photographs BCP, BCP2					
Procedure	Progra	m Code	Challenges/Shipment		
	BCP BCP2				
Blood cell identification		ı	5		
Educational challenge(s)			5 (BCP)/1 (BCP2)		

Program Information

- BCP Ten images, each available as photographs and online images
- BCP2 Six images, each available as photographs and online images
- · Three shipments per year



Erythrocyte Sedimentation Rate ESR, ESR1, ESR2, ESR3 Challenges/ **Procedure Program Code** Shipment **ESR** ESR1 ESR2 ESR3 All methods except the ALCOR, Alifax®, Sedimat 15®, and Sedimat 3 15 Plus Sedimat 15, Sedimat 15 Plus 3 3 Alifax ALCOR iSED 3

- ESR, ESR1 Three 6.0-mL whole blood specimens
- ESR2 Three 3.0-mL simulated whole blood specimens
- ESR3 Three 4.5-mL whole blood specimens
- · Two shipments per year

Hematology Automated Differential Series FH1–FH13, FH1P–FH13P					eries
Analyte/Procedure		Program Code Challenges Shipment			Challenges/ Shipment
	FH1- FH10	FH1P- FH10P	FH13	FH13P	
Blood cell identification				•	10
Hematocrit					5
Hemoglobin					5
Immature granulocyte parameter					5 (FH9 only)
Large unstained cell (LUC)					5 (FH4 only)
MCV, MCH, and MCHC					5
MPV				•	5
Nucleated red blood cell count (nRBC)	•	•		•	5 (FH3, FH9, and FH13)
Platelet count					5
RDW		•		•	5
Red blood cell count				•	5
White blood cell count					5
WBC differential					5

For second instrument reporting options, see the Quality Cross Check programs, FH3Q, FH4Q, FH6Q, and FH9Q, on page 136.

Centrifugal Hematology FH15Analyte/ProcedureProgram CodeChallenges/ShipmentFH15FH15HematocritI5HemoglobinI5Platelet countI5WBC countI5WBC differential (2-part)I5

Program Information

- FH1-FH10 and FH1P-FH10P

 Five 2.5-mL whole blood
 specimens with pierceable caps
- FH13 and FH13P Five 2.0-mL whole blood specimens with pierceable caps
- FHP series Ten images, each available as photographs and online images
- For method compatibility, see instrument matrix on page 137
- Three shipments per year



- Five 0.6-mL whole blood specimens
- For use with QBC instruments; not intended for spun hematocrit methods
- Three shipments per year

12

Quality Cross Check—Hematology Series FH3Q, FH4Q, FH6Q, FH9Q					
Analyte/Procedure	Program Code Challenges/ Shipment				
	FH3Q	FH4Q	FH6Q	FH9Q	
Hematocrit	•	•	ı		3
Hemoglobin			ı		3
Immature granulocyte parameter					3
Large unstained cells (LUC)		ı			3
MCV, MCH, MCHC	•	ı	•		3
MPV			ı		3
Nucleated red blood cell count (nRBC)				•	3
Platelet count			ı		3
RDW			ı		3
Red blood cell count	•	1	•	ı	3
White blood cell count		ı			3
WBC differential	I	ı	I		3

These programs do not meet regulatory requirements for proficiency testing; see the FH Series on page 135. For additional information about the CAP Quality Cross Check program, see page 42.

The Quality Cross Check Program:

- · Provides a solution for monitoring performance across multiple instruments, and is in compliance with the CMS directive regarding proficiency testing on multiple instruments.
- Simplifies instrument comparability efforts by providing custom reports with both peer group comparison and instrument comparability statistics.

Program Information

- Three 2.5-mL whole blood specimens with pierceable
- · Report up to three instruments
- For method compatibility, see instrument matrix on page 137
- Two shipments per year

Have you created or updated your CAP Profile?

Each laboratory staff member should have their own profile. Your profile is transferrable when you leave your current position. Use it to maintain information about yourself, including:

- Business affiliations
- Certifications
- Contact preferences
- Inspector-related information
- Personal contact information
- Specialties and skills
- Addresses

To create or update your profile, visit cap.org, log in, and click on UPDATE MY PROFILE.



Hematology Automated Differential Series, Instrument Matrix

Instrument					FH and F	HQ Series	3			
	FH1	FH2	FH3/ FH3Q	FH4/ FH4Q	FH6	FH6Q	FH9/ FH9Q	FH10	FH13	FH15
Abbott Cell-Dyn® 1200, 1400, 1600, 1700, 1800, Emerald™	•									
Horiba ABX 9000+, 9018+, 9020+										
Sysmex K-series, KCP-1, KX-21/21N, pocH-100i, XP-series	•									
CDS/Medonic M-Series		ı								
Coulter® AcT™, MD, ONYX™, S880, S-plus V, ST, STKR, T-series		•								
Drew Scientific DC-18, Drew3, Excell 10/16/18, I-1800,		•								
Horiba ABX Micros		ı								
Mindray BC - 2800, 3000/3200 series		ı								
Siemens ADVIA® 360										
Abbott Cell-Dyn 3000, 3200, 3500, 3700, 4000, Ruby™, Sapphire™			1							
Cell-DYN Emerald 22			ı							
Coulter DxH 500			1							
Drew Scientific Excell 22, 2280			ı							
Orphee Mythic 22 AL, Orphee Mythic 22 OT										
Siemens ADVIA 560			ı							
Siemens ADVIA 120, 120 w/SP1, 2120				ı						
Coulter Gen-S™, HmX, LH500, MAXM™, MAXM A/L, STKS, VCS™					•	•				
Sysmex XE-2100, XE-2100C, XE-2100D, XE-2100DC, XE-2100L, XE-5000, XN-Series, XS-500i, XS-800i, XS-1000i, XS-1000iAL, XS-1000iC, XT-1800i, XT-2000i, XT-4000i, XE-2100D/L (Blood Center)							•			
Coulter AcT 5 diff (AL, CP, OV)								ı		
DIRUI BF Series										
Horiba ABX Pentra 60, 80, 120										
Coulter LH750, LH755, LH780, LH785, Unicel DxH						•			•	
QBC										•

12

Blood Parasite BP					
Procedure	Program Code Challenges/Shipmen				
	ВР				
Thin/thick blood film sets*	ı	5			

^{*}This Survey will include corresponding thick films when available.

Program Information

- · Five Giemsa-stained blood film sets, photographs, and/or online images
- · A variety of blood parasites, including Plasmodium, Babesia, Trypanosoma, and filarial worms
- Three shipments per year

Bone Marrow Cell Differential BMD					
Procedure	Challenges/Shipment				
	BMD				
Bone marrow differential, including myeloid:erythroid ratio		1			
Bone marrow cell identification	I	5			

Additional Information

- Examine an online whole slide image that includes a manual 500 bone marrow differential count and annotated cells for identification.
- · Recognize and integrate problem-solving skills through the use of interpretive questions found throughout the discussion.
- Evaluate cell morphology and identify specific cells in bone marrow.
- · See system requirements on page 13.

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Program Information

- One online bone marrow aspirate whole slide image that includes five annotated cells for identification
- Powered by DigitalScope[®] technology
- Two online activities per year; your CAP shipping contact will be notified via email when the activity is available



Fetal Red Cell Detection HBF **Procedure Program Code** Challenges/Shipment **HBF** 2 Kleihauer-Betke and flow cytometry 2 Rosette fetal screen 1 Acid elution whole slide image П

- Two 1.2-mL liquid whole blood specimens
- · Not designed for F cell quantitation
- Two online whole slide images per year with optional grids for cell counting
- Powered by DigitalScope[®] technology
- · Two shipments per year

Hemoglobinopathy HG			
Procedure	Program Code	Challenges/Shipment	
	HG		
Hemoglobin identification and quantification	ı	4	
"Dry lab" educational challenges	I	2	
Hemoglobin A ₂ quantitation	ı	4	
Hemoglobin F quantitation	I	1	
Sickling test, qualitative	ı	4	

Program Information

- Four 0.5-mL stabilized red blood cell specimens
- Two "dry lab" educational challenges (case histories, electrophoresis patterns, and clinical interpretation questions)
- Two shipments per year

Reticulocyte Series RT, RT2, RT3, RT4					
Instrument/Method	Program Code		•	Challenges/Shipment	
	RT	RT2	RT3	RT4	
Abbott Cell-Dyn 4000, Sapphire, Siemens ADVIA 120/2120, and all other automated and manual methods					3
Abbott Cell-Dyn 3200, 3500, 3700, Ruby					3
Coulter GenS, HmX, LH500, LH700 series, MAXM, STKS, Unicel DxH					3
Sysmex XE-2100, XE-2100C, XE-5000, XN Series, XT-2000i, XT-4000i				•	3
Pierceable caps					3

- RT, RT2 Three 1.0-mL stabilized red blood cell specimens
- RT3, RT4 Three 2.0-mL stabilized red blood cell specimens
- Includes percentage and absolute result reporting
- Two shipments per year

Quality Cross Check—Reticulocyte RTQ, RT2Q, RT3Q, RT4Q Challenges/ Instrument/Method **Program Code** Shipment RTQ RT2Q RT3Q RT4Q Abbott Cell-Dyn 4000, Sapphire, 3 Siemens ADVIA 120/2120, and all other automated and manual methods Abbott Cell-Dyn 3200, 3500, 3700, 3 Ruby Coulter GenS, HmX, LH500, LH700 3 series, MAXM, STKS, Unicel DxH Sysmex XE-2100, XE-2100C, XE-5000, 3 XN Series, XT-2000i, XT-4000i

These programs do not meet regulatory requirements for proficiency testing; see the RT Series on page 139. For additional information about the CAP Quality Cross Check program, see page 42.

The Quality Cross Check Program:

- Provides a solution for monitoring performance across multiple instruments, and is in compliance with the CMS directive regarding proficiency testing on multiple instruments.
- Simplifies instrument comparability efforts by providing custom reports with both peer group comparison and instrument comparability statistics.

Program Information

- RTQ, RT2Q Three 1.0-mL stabilized red blood cell specimens
- RT3Q Three 3.0-mL stabilized red blood cell specimens
- RT4Q Three 2.0-mL stabilized red blood cell specimens
- Includes percentage and absolute result reporting
- Report up to three instruments
- Two shipments per year

Sickle Cell Screening SCS			
Procedure	Program Code	Challenges/Shipment	
	scs		
Sickling test, qualitative	I	3	

Transfusion-Related Cell Count TRC			
Procedure	Procedure Program Code		
	TRC		
Platelet count (platelet-rich plasma)	ı	5	
WBC count	I	4	
Dry challenge	I	2	

WBC counts must be performed using a Nageotte chamber, fluorescence microscopy, or by flow cytometry.

Program Information

- Three 1.0-mL stabilized human erythrocyte specimens
- Two shipments per year

- Five 1.2-mL suspensions of platelet-rich plasma
- Two 1.0-mL vials leukocytereduced platelet material
- Two 1.0-mL vials leukocytereduced red blood cells
- Three shipments per year

Waived Combination HCC, HCC2			
Analyte	Program Code Challenges/Shipmen		
	HCC	HCC2	
Hematocrit			2
Hemoglobin			2
Urinalysis/Urine hCG			2
Whole blood glucose	I	I	2 (HCC)/3 (HCC2)

Program Information

- HCC Two 1.0-mL whole blood specimens; two shipments per year
- HCC2 Total of four shipments per year
- Hematocrit, hemoglobin, and urinalysis/urine hCG testing -Two 3.0-mL whole blood specimens and two 10.0-mL urine specimens; two shipments per year: A and C
- Whole blood glucose testing - Three 2.5-mL whole blood specimens; two shipments per year: B and D
- To verify instrument compatibility, refer to the instrument matrix on page 66

Rapid Total White Blood Cell Count RWBC			
Procedure	Program Code	Challenges/Shipment	
	RWBC		
Rapid total white blood cell count	ı	5	

Program Information

- Five 2.0-mL whole blood specimens
- For use with the HemoCue WBC instrument
- Three shipments per year

Virtual Peripheral Blood Smear VPBS			
Procedure	Program Code	Challenges/Shipment	
	VPBS		
WBC differential		3	
Platelet estimate		3	
RBC morphology	1	3	
Blood cell identification	I	15	

Additional Information

- Examine online whole slide images that include a manual 100 WBC differential count and annotated cells for identification.
- Evaluate and identify red blood cell (RBC) morphology and identify specific white blood cells (WBC) in peripheral blood.
- Recognize and integrate problem-solving skills through the use of interpretive questions found throughout the discussion.
- · See system requirements on page 13.

- Three online peripheral blood whole slide images that include 15 annotated cells for identification
- Powered by DigitalScope technology
- Two online activities per year; your CAP shipping contact will be notified <u>via email</u> when the activity is available



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Expanded Virtual Peripheral Blood Smear EHE1			
Procedure	Program Code	Challenges/Shipment	
	EHE1		
WBC differential	1	2	
Platelet estimate	I	2	
RBC morphology	I	2	
WBC morphology	I	2	
Blood cell identification	ı	10	

Additional Information

- More challenging and/or complex testing.
- Examine online whole slide images that include a manual 100 WBC/differential count and annotated cells for identification.
- · Comprehensive case studies.
- Ability to recognize and integrate problem-solving skills through the use of interpretive questions found throughout discussion.
- Evaluate and identify red blood cell (RBC) morphology and identify specific white blood cells (WBC) in peripheral blood.
- See system requirements on page 13.

- Two online peripheral blood whole slide images that include 10 annotated cells for identification
- Powered by DigitalScope technology
- Two online activities per year; your CAP shipping contact will be notified <u>via email</u> when the activity is available

Hematopathology Online Education HPATH/HPATH1					
Program Code Challenges/Shipment					
HPATH/HPATH1					
Hematopathology online case eview					

Additional Information

HPATH educates pathologists, hematolopathologists, and hematologists with an interest in hematopathology to assess and improve their diagnostic skills in hematopathology.

- All cases have been specially selected to highlight important changes in the 2016 revision of the WHO Classification.
- · Clinical history and relevant laboratory data.
- At least one online whole slide image of peripheral blood, bone marrow, spleen, lymph node, or other tissue.
- Results of ancillary studies such as immunohistochemistry, flow cytometry, FISH, karyotyping, and molecular studies, where appropriate.
- · Case discussion and discussion of differential diagnoses.
- · Five SAM questions per case.
- See system requirements on page 13.

- HPATH Five diagnostic challenges/online whole slide images with clinical history; reporting with CME/SAM credit is available for one pathologist/ hematologist. For additional pathologist/hematologist, order HPATH1
- HPATH1 Reporting option with CME/SAM credit for each additional pathologist and hematologist (within the same institution); must order in conjunction with Survey HPATH
- Earn a maximum of 12.5 CME/SAM credits (AMA PRA Category 1 Credits™) per pathologist and a maximum of 12.5 CE credits per hematologist for completion of an entire year
- This activity meets the ABP MOC Part IV Practice Performance Assessment requirements
- Powered by DigitalScope technology
- Two online activities per year; your CAP shipping contact will be notified <u>via email</u> when the activity is available





12

Clinical Microscopy

Analytes/procedures in bold type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Urinalysis and Clini	CMP, CMP1		
Analyte/Procedure	Progra	ım Code	Challenges/Shipment
	СМР	CMP1	
Bilirubin			3
Blood or hemoglobin			3
Body fluid photographs			3
Glucose			3
hCG urine, qualitative			3
Ketones		ı	3
Leukocyte esterase		ı	3
Nitrite			3
Osmolality		ı	3
рН		ı	3
Protein, qualitative	•	ı	3
Reducing substances		ı	3
Specific gravity	ı	ı	3
Urine sediment photographs	•	ı	3
Urobilinogen	ı		3

Program Information

- CMP Three 10.0-mL liquid urine specimens; for use with all instruments except iCHEM; six images, each available as photographs and online images
- CMP1 Three 12.0-mL liquid urine specimens; for use with iCHEM instruments; six images, each available as photographs and online images
- · Two shipments per year



Additional Information

For second instrument reporting options, see the Quality Cross Check program, CMQ, on page 145.

Test your diagnostic skills as a pathologist with CPIP

Online, hands-on and interactive, the Clinical Pathology Improvement Program (CPIP) enables pathologists to sharpen their diagnostic skills in real time by working through an actual case. Each month, you will receive a new scenario, including slide images and clinical background. As the case unfolds, more information is revealed, just as in the laboratory. Participants who successfully complete the posttest may apply their earned credits to their MOC SAM requirements. Enjoy a full year of CPIP and earn up to 15 CME/SAM credits.

Choose code CPIP/CPIP1 on your Surveys order form.

Quality Cross Check—Urinalysis CMQ				
Analyte	Program Code	Challenges/Shipment		
	CMQ			
Bilirubin	ı	3		
Blood or hemoglobin	ı	3		
Glucose	I	3		
hCG urine, qualitative	I	3		
Ketones		3		
Leukocyte esterase		3		
Nitrite		3		
Osmolality		3		
рН		3		
Protein, qualitative		3		
Reducing substances	ı	3		
Specific gravity	I	3		
Urobilinogen	1	3		

This program does not meet regulatory requirements for proficiency testing; see Surveys CMP and CMP1 on page 144. For additional information about the CAP Quality Cross Check program, see page 42.

The Quality Cross Check Program:

- Provides a solution for monitoring performance across multiple instruments, and is in compliance with the CMS directive regarding proficiency testing on multiple instruments.
- Simplifies instrument comparability efforts by providing custom reports with both peer group comparison and instrument comparability statistics.

Clinical Microscopy Miscellaneous Photopage CMMP				
Procedure	Program Code Challenges/Shipn			
	СММР			
Fern test (vaginal)	ı	1		
KOH preparation (skin or vaginal)	ı	1		
Nasal smear	ı	1		
Pinworm preparation	ı	1		
Spermatozoa NEW	ı	1		
Stool for leukocytes	ı	1		
Urine sediment photographs	ı	3		
Vaginal wet preparation photographs (for clue cells, epithelial cells, trichomonas, and yeast)	ı	1		

Program Information

- Three 10.0-mL liquid urine specimens for use with all instruments
- Report up to three instruments
- Two shipments per year

- Ten images, each available as photographs and online images
- · Two shipments per year

Virtual Body Fluid VBF				
Procedure	Program Code	Challenges/Shipment		
	VBF			
Total nucleated cells differential	I	2		
Body fluid cell identification				

Additional Information

- Examine online whole slide images that include a manual differential count and annotated cells for identification.
- Evaluate cell morphology and identify specific cells in a body fluid.
- See system requirements on page 13.

Amniotic Fluid Leakage AFL					
Procedure	edure Program Code Challenges/Shipment				
AFL					
pH interpretation 3					

Program Information

- Two online whole slide body fluid images that include 10 annotated cells for identification
- Powered by DigitalScope technology
- Two online activities per year; your CAP shipping contact will be notified <u>via email</u> when the activity is available

Program Information

- Three 2.0-mL liquid specimens
- For use with nitrazine paper and the Amniotest™
- Two shipments per year

Automated Body Fluid Series ABF1, ABF2, ABF3				
Procedure	Program Code Challenges/Shipment			
	ABF1	ABF2	ABF3	
Red blood cell fluid count				2
White blood cell fluid count			ı	2

For method compatibility, see instrument matrix below.

Automated Body Fluid, Instrument Matrix

Instrument	ABF Series		
	ABF1	ABF2	ABF3
Advanced Instruments GloCyte, Siemens ADVIA 120/2120 series	1		
Coulter LH 700 Series, Unicel DxH		•	
Sysmex XE-2100, XE-5000, XN- series, XT-1800i, XT-2000i, XT- 4000i			
IRIS iQ [®] 200			

- Two 3.0-mL simulated body fluid specimens
- Two shipments per year

Automated Urine Microscopy UAA, UAA1				
Analyte	Progra	Program Code Challenges/Shipment		
	UAA UAA1			
Casts, semiquantitative			2	
Crystals, semiquantitative			2	
Epithelial cells, semiquantitative			2	
Red blood cells, quantitative	•		2	
White blood cells, quantitative	ı		2	

- UAA Two 10.0-mL liquid urine specimens for use with IRIS and Roche instruments
- UAA1 Two 12.0-mL liquid urine specimens for use with Sysmex instruments
- Two shipments per year

Automated Urinalysis, Instrument Matrix

Instrument	UAA,	UAA1
	UAA	UAA1
DIRUI FUS	X	
IRIS Iq200	X	
Roche cobas u701	X	
ARKRAY Auction Hybrid		X
77 Elektronika		X
Sysmex UF 50, 100, 500i, 1000i, 5000		X
Sysmex UX 2000		Х

Crystals BCR, BFC, URC					
Procedure	Program Code Challenges/Shipment				
	BCR BFC URC				
Bile crystal identification				2	
Body fluid crystal identification	1 2			2	
Urine crystal identification	1 2				

D	ipstick Confirmatory D	sc
Analyte	Program Code	Challenges/Shipment
	DSC	
Riliruhin		2

Sulfosalicylic acid (SSA)

Program Information

- BFC Two 1.5-mL simulated body fluid specimens (eg, synovial fluid)
- URC Two 1.5-mL urine specimens
- BCR Two photographs
- Two shipments per year

- Two 12.0-mL liquid urine specimens
- For use with methods to confirm positive bilirubin and protein dipstick results
- Two shipments per year

Fecal Fat FCFS		
Analyte	Program Code	Challenges/Shipment
	FCFS	
Fecal fat, qualitative		2

- Two 10.0-g simulated fecal fat specimens
- For microscopic detection of neutral fats (triglycerides) and/or split fats (total free fatty acids)
- Two shipments per year

Fetal Hemoglobin APT		
Analyte	Program Code	Challenges/Shipment
	APT	
Fetal hemoglobin (gastric fluid)		2

Program Information

- Two 1.2-mL simulated gastric fluid specimens
- Two shipments per year

Gastric Occult Blood GOCB		
Analyte	Program Code	Challenges/Shipment
	GOCB	
Gastric occult blood	I	3
Gastric pH	ı	3

Program Information

- Three 2.0-mL simulated gastric specimens
- Two shipments per year

Glucose-6-Phosphate Dehydrogenase G6PDS				
Analyte	alyte Program Code Challenges/Shipment			
	G6PDS			
G6PD, qualitative and quantitative	I	2		

Program Information

- Two 0.5-mL lyophilized hemolysate samples
- Two shipments per year

Hemocytometer Fluid Count HFC		
Procedure	Program Code	Challenges/Shipment
	HFC	
Cytopreparation differential		3
Red blood cell fluid count		3
White blood cell fluid count		3

- Three 1.0-mL simulated body fluid specimens
- Two shipments per year

Hemocytometer Fluid Count, International HFCI		
Procedure	Program Code	Challenges/Shipment
	HFCI	
Red blood cell fluid count	I	3
White blood cell fluid count	I	3
Body fluid differential	ı	2

This program meets the CAP's Accreditation Program requirements.

Additional Information

- Examine online whole slide images that include a manual differential count.
- See system requirements on page 13.

Lamellar Body Count LBC		
Procedure	Program Code	Challenges/Shipment
	LBC	
Lamellar body count		3

Occu	lt Blood OCB	
Analyte	Program Code	Challenges/Shipment
	OCB	
Occult blood		3

Additional Information

For second instrument reporting options, see the Quality Cross Check program, OCBQ, on page 150.

Program Information

- Three 2.0-mL simulated body fluid specimens; two online whole slide images for 2- and 5-part differential
- Designed for international laboratories that have experienced significant shipping and receiving issues and need longer program stability
- Powered by DigitalScope technology
- Two shipments per year

Program Information

- Three 2.0-mL simulated liquid amniotic fluid specimens
- For use with LBC methods performed on all hematology analyzers
- Two shipments per year

- Three 2.0-mL simulated fecal specimens
- Two shipments per year

Quality Cross Check—Occult Blood OCBQ		
Analyte	Program Code	Challenges/Shipment
	OCBQ	
Occult blood	ı	3

This program does not meet regulatory requirements for proficiency testing; see Survey OCB on page 149. For additional information about the CAP Quality Cross Check program, see page 42.

The Quality Cross Check Program:

- Provides a solution for monitoring performance across multiple instruments, and is in compliance with the CMS directive regarding proficiency testing on multiple instruments.
- Simplifies instrument comparability efforts by providing custom reports with both peer group comparison and instrument comparability statistics.

Program Information

- Three 2.0-mL simulated fecal specimens
- Report up to three instruments
- Two shipments per year

Rupture of Fetal Membranes Testing ROM1			
Procedure Program Code Challenges/Shipmen			
	ROM1		
Rupture of fetal membranes	ı	3	

Program Information

- Three 0.5-mL simulated vaginal specimens for methods such as Amnisure, Actim PROM, and Clinical Innovations
- Two shipments per year

Special Clinical Microscopy SCM1, SCM2			
Analyte/Procedure	Progra	m Code	Challenges/Shipment
	SCM1	SCM2	
Urine hemosiderin, Prussian blue	ı		3
Urine eosinophils, Wright stain		ı	3

Program Information
• Three images, each available
as photographs and online
images
Two shipments per year

Ticks, Mites, and Other Arthropods TMO								
Procedure Program Code Challenges/Shipmen								
	ТМО							
Tick, mite, and arthropod identification	ı	3						

•	Three images, each available
	as photographs and online
	images

• Two shipments per year

Program Information

Urine hCG UHCG						
Procedure	Program Code	Challenges/Shipment				
	UHCG					
Urine hCG, qualitative	•	5				

- Five 1.0-mL urine specimens
- · Three shipments per year

Urine Albumin and Creatinine, Semiquant UMC							
Analyte/Procedure	Program Code	Challenges/Shipment					
	UMC						
Creatinine	I	2					
Urine albumin (microalbumin): creatinine ratio	ı	2					
Urine albumin (microalbumin), semiquantitative	ı	2					

For quantitative reporting, refer to Survey U, page 68.

Program Information

- Two 3.0-mL liquid urine specimens
- For use with dipstick and semiquantitative methods only
- Two shipments per year

Worm Identification WID						
Procedure	Program Code Challenges/Shipm					
	WID					
Worm identification		3				

Program Information

- Three images, each available as photographs and online images
- Two shipments per year

World-class recognition deserves to be displayed.



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Proudly display the mark. It distinguishes you as one of almost 8,000 laboratories worldwide that have attained CAP accreditation, the most respected and recognized laboratory accreditation in the world.

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In just seconds, the CAP's Performance Analytics Dashboard provides valuable insights into your laboratory's performance, letting you proactively focus energy on areas that need immediate attention while filtering out distractions. Updated daily, this complimentary Surveys and CAP accreditation performance monitoring tool reduces the stress of managing today's laboratory by giving you fast access to a single laboratory's or an expansive network's performance.

To view a demo, search Performance Analytics Dashboard at cap.org.

13

13 Reproductive Medicine



Access unique, integrated resources.

Reproductive medicine's integrated laboratory improvement program offers:

- Unique accreditation checklist specific for the subspecialty of reproductive medicine
- Comprehensive proficiency testing and educational programs for andrology and embryology

Reproductive Medicine

Andrology and Embryology......154

13

Andrology and Embryology

Analytes/procedures in **bold** type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Semen Analysis SC, SC1, PV, SM, SV, ASA									
Procedure		Program Code Ch							
	SC	SC1	PV	SM	SV	ASA			
Sperm count and presence/absence (manual methods and CASA systems)	•						2		
Sperm count and presence/absence (automated methods)							2		
Postvasectomy sperm count and presence/absence							2		
Sperm morphology							2		
Sperm viability							2		
Antisperm antibody IgG							2		

Program Information

- SC Two 0.3-mL stabilized sperm specimens
- SC1 Two 1.0-mL stabilized sperm specimens
- PV Two 0.3-mL stabilized sperm specimens with counts appropriate for postvasectomy testing
- SM Two prepared slides for staining
- SV Two eosin-nigrosinstained slides
- ASA Two 0.3-mL serum specimens
- Two shipments per year



Sperm Motility, Morphology, and Viability SMCD, SM1CD, SM2CD **Procedure Program Code** Challenges/Shipment **SMCD** SM1CD SM2CD Sperm count П 2 2 Sperm motility/forward progression П Sperm morphology 2 2 Sperm viability

- SMCD One CD-ROM with video clips
- SM1CD Two challenges, each available as images on CD-ROM and online whole slide images powered by DigitalScope® technology
- SM2CD Two challenges, each available as images on CD-ROM and online whole slide images powered by DigitalScope technology
- Two shipments per year



Embryology EMB							
Procedure	Program Code	Challenges/Shipment					
	EMB						
Embryo transfer and quality assessment (three- and five-day-old embryos)	ı	4					

- One CD-ROM with video clips
- Two shipments per year

Ligand—Special Y, YY, DY								
Analyte	Progra	ım Code	Challenges/Shipment					
	Y, YY	DY						
11-deoxycortisol			3					
17-hydroxyprogesterone			3					
Androstenedione			3					
DHEA sulfate			3					
Estradiol			3					
Estriol, unconjugated (uE3)			3					
Follicle-stimulating hormone (FSH)			3					
Growth hormone (GH)			3					
IGF-1 (somatomedin C)			3					
Luteinizing hormone (LH)			3					
Progesterone			3					
Prolactin			3					
Testosterone			3					
Testosterone, bioavailable			3					
Testosterone, free			3					
Sex hormone-binding globulin (SHBG)			3					

Program Information

- Y Three 5.0-mL liquid serum specimens in duplicate
- YY Three 5.0-mL liquid serum specimens in triplicate
- DY Must order in conjunction with Survey Y or YY
- Conventional and International System of Units (SI) reporting offered
- Two shipments per year



Antimüllerian Hormone AMH Analyte Program Code Challenges/Shipment AMH Antimüllerian hormone 3

- Three 1.0-mL lyophilized serum specimens
- Two shipments per year

Are you ready for your CAP inspection?



The CAP Accreditation Readiness Assessment (CARA®) is an on-site evaluation and education program for laboratories just beginning their pursuit of CAP accreditation.

CARA focuses on:

- Facilitating an in-depth understanding of CAP requirements as they apply to your laboratory
- Helping you manage the time and resources necessary for compliance with CAP accreditation requirements and preparation for your initial inspection
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Email us at readiness-assessment@cap.org to accelerate your quality journey.

14

14 Coagulation



Meet requirements for calibration verification and linearity for coagulation testing.

- · Hemostasis test methods that are calibrated and directly measure the concentration of an analyte require calibration verification/linearity (CVL).
- Coagulation programs available include Heparin CVL (LN36), von Willebrand Factor Antigen CVL (LN37), D-Dimer CVL (LN42), Thrombophilia CVL (LN35), and Fibrinogen CVL (LN44).

New Programs NEW



Apixaban Anticoagulant Monitoring (APXBN)161

Discontinued Programs

Whole Blood D-Dimer (WBDD)

D-dimer

Coagulation

Analytes/procedures in bold type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Coagulation—Limited CGB, CGL, CGDF								
Analyte	Program Code Challenges, Shipment							
	CGB	CGL	CGDF					
Activated partial thromboplastin time	•			5				
Fibrinogen	1		5					
International normalized ratio (INR)*	1 1			5				
Prothrombin time	1 1			5				
D-dimer**				2 per year				
Fibrin(ogen) degradation products, plasma**	1 1		2 per year					
Fibrin(ogen) degradation products, serum**		ı		2 per year				

^{*}Participants reporting INR results will receive a special evaluation to assess the INR calculation.
**D-dimer and FDP are shipped with the CGL A and C mailings.

Additional Information

For second instrument reporting options, see the Quality Cross Check program, CGLQ, below.

Quality Cross Check—Coagulation CGLQ Analyte Program Code Challenges/Shipment CGLQ CGLQ Activated partial thromboplastin time Image: CGLQ and CG

1

1

This program does not meet regulatory requirements for proficiency testing; see Survey CGL above. For additional information about the CAP Quality Cross Check program, see page 42.

The Quality Cross Check Program:

Fibrin(ogen) degradation products,

Fibrin(ogen) degradation products, serum

- Provides a solution for monitoring performance across multiple instruments, and is in compliance with the CMS directive regarding proficiency testing on multiple instruments.
- Simplifies instrument comparability efforts by providing custom reports with both peer group comparison and instrument comparability statistics.

Program Information

- CGB Five 1.0-mL lyophilized plasma specimens; three shipments per year
- CGL Five 1.0-mL lyophilized plasma specimens; three shipments per year; one 1.0-mL plasma specimen and one 1.0-mL serum specimen; three shipments per year
- CGDF One 1.0-mL serum specimen; one 1.0-mL lyophilized plasma specimen; two shipments per year



- Three 1.0-mL lyophilized plasma specimens in triplicate, one 1.0-mL plasma specimen, and one 2.0-mL serum specimen
- Report up to three instruments
- · Two shipments per year

Coagulation—E	xtended CGE, CGE	(
Analyte	Program Code	Challenges/ Shipment
	CGE, CGEX	
See analyte listing below		2

- CGE Two 1.0-mL lyophilized plasma specimens in triplicate
- CGEX Two 1.0-mL lyophilized plasma specimens (five vials each)
- Two shipments per year

Coagulation Analyte Listing

50:50 mixing study, PT and aPTT Activated partial thromboplastin time

Activated protein C resistance

Alpha-2-antiplasmin

Anti-beta-2-glycoprotein (IgG and IgM)

Antiphospholipid antibody (IgG, IgM, and IgA)

Antithrombin activity/antigen

Dilute prothrombin time

Dilute Russell's viper venom time

Euglobulin test

Factors II, V, VII, VIII, IX, X, XI,

XII, and XIII

Factor VIII assay

Fibrin monomer Fibrinogen activity

Fibrinogen antigen

Heparin-induced thrombocytopenia (HIT)

High molecular weight kininogen

Kaolin-activated aPTT

Kaolin clotting time

Lupus anticoagulant

Plasminogen activator inhibitor

Plasminogen activity/antigen

Prekallikrein

Protein C

Protein S

Prothrombin fragment 1.2

Prothrombin time

Reptilase time

Thrombin-antithrombin

Thrombin time

Tissue plasminogen activator

von Willebrand factor activity:

- Collagen binding
- Glycoprotein I_b binding
- Ristocetin cofactor

von Willebrand factor antigen

von Willebrand multimer analysis

Coagulation Special Testing Series CGS1, CGS2, CGS3, CGS4, CGS5, CGS6, CGS7, CGS8

000 - , 000	0,0		ouc	7,0	400			
Module/Analyte			Cha	llenges	/Shipm	nent		
				Progra	m Code			
	CGS1	CGS2	CGS3	CGS4	CGS5	CGS6	CGS7	CGS8
Activated partial thromboplastin time*	2		2	3				
International normalized ratio (INR)	2			3				
Prothrombin time*	2			3				
Lupus Anticoagulant and Mixing St	udies	Module	Э					
Dilute Russell's viper venom time	2							
Lupus anticoagulant (confirmation and screen)	2							
50:50 mixing studies, PT and aPTT	2							
Thrombophilia Module								
Activated protein C resistance		2						
Antithrombin (activity, antigen)		2						
Protein C (activity, antigen)		2						
Protein S (activity, free antigen, total antigen)		2						
von Willebrand Factor Antigen Mod	lule							
Factor VIII assay			2					
von Willebrand factor (antigen, activity, multimers)			2					
Heparin Module								
Heparin activities using methodologies including Anti-Xa (unfractionated, low molecular weight, and hybrid curve)				3				
Thrombin time				3				
Heparin-Induced Thrombocytopen	ia Mod	ule						
Appropriate with methods such as Gen-Probe Lifecodes PF4 IgG and Gen-Probe Lifecodes PF4 Enhanced® assays					2			
Appropriate with the Akers Biosciences, Inc. PIFA® Heparin/ Platelet Factor 4 Rapid Assay						3		
Continued on the next page								

^{*}Not appropriate for meeting regulatory requirements, see page 158.

- CGS1, CGS2, CGS3 A total of two 2.0-mL of lyophilized plasma
- CGS4 Three 1.0-mL lyophilized plasma specimens
- CGS5 Two 60.0-µL serum specimens
- CGS6, CGS8 Three 400.0-µL frozen serum specimens
- CGS7 Three 1.0-mL lyophilized plasma specimens in duplicate
- Two shipments per year

Coagulation Special Te CGS4, CGS5, C		_					, CGS	3,
Module/Analyte Challenges/Shipment								
		Program Code						
	CGS1	CGS2	CGS3	CGS4	CGS5	CGS6	CGS7	CGS8
Heparin-Induced Thrombocytopen	ia Mod	ule co	ntinue	d				
Appropriate with the Akers Biosciences, Inc. PIFA PlussPF4™ Heparin/Platelet Factor 4 Rapid Assay								3
ADAMTS13 Module								
ADAMTS13 (activity, inhibitor							3	

^{*}Not appropriate for meeting regulatory requirements, see page 158.

screen, and titer)

Program Information

- CGS1, CGS2, CGS3 A total of 2.0-mL of lyophilized plasma
- CGS4 Three 1.0-mL lyophilized plasma specimens
- CGS5 Two 60.0-µL serum specimens
- CGS6, CGS8 Three 400.0-μL frozen serum specimens
- CGS7 Three 1.0-mL lyophilized plasma specimens in duplicate
- Two shipments per year

Apixaban, Dabigatran, Fondaparinux, **Rivaroxaban Anticoagulant Monitoring** APXBN, DBGN, FNPX, RVBN

Analyte	Pr	Program Code			
	APXBN NEW	DBGN	FNPX	RVBN	
Activated partial thromboplastin time*	•				3
Prothrombin time*					3
Thrombin time					3
Apixaban					3
Dabigatran					3
Fondaparinux					3
Rivaroxaban					3

^{*}Not appropriate for meeting regulatory requirements, see page 158.

Program Information

3

- Three 1.0-mL lyophilized specimens
- Two shipments per year

1	4
tion	

Activated Clotting Time Series CT, CT1, CT2, CT3, CT5						
Instrument/Cartridge		Prog	gram (Code		Challenges/Shipment
	СТ	CT1	CT2	СТЗ	СТ5	
Helena Actalyke®	•					3
Helena Cascade POC						3
IL Gem® PCL ACT						3
IL Gem PCL ACT-LR						3
IL GEM PCL Plus ACT				ı		3
IL GEM PCL Plus ACT-LR						3
ITC Hemochron® CA510/FTCA510	ı					3
ITC Hemochron FTK-ACT	ı					3
ITC Hemochron Jr. Signature/ACT+				ı		3
ITC Hemochron Jr. Signature/ACT-LR						3
ITC Hemochron P214/P215	ı					3
i-STAT® Celite® and Kaolin ACT						3
Medtronic HemoTec ACT/ACTII/ACT Plus HR-ACT		•				3
Medtronic HemoTec ACT/ACTII/ACT Plus LR-ACT		•				3
Medtronic HemoTec ACT/ACTII/ACT Plus R-ACT		•				3
Medtronic Hepcon HMS, HMS Plus		•				3
Sienco Sonoclot®	1					3

Additional Information

For second instrument reporting options, see the Quality Cross Check programs CTQ-CT3Q and CT5Q, on page 163.

- CT Three 3.0-mL lyophilized whole blood specimens with corresponding diluents
- CT1 Three 1.7-mL lyophilized whole blood specimens with corresponding diluents
- CT2 Three 0.5-mL lyophilized whole blood/ diluent ampules
- CT3 Three 0.5-mL lyophilized whole blood/ diluent ampules
- CT5 Three 1.7-mL lyophilized whole blood specimens with corresponding diluents
- Two shipments per year

Quality Cross Check— Activated Clotting Time Series CTQ, CT1Q, CT2Q, CT3Q, CT5Q

Instrument/Cartridge	Program Code					Challenges/ Shipment
	CTQ	CT1Q	CT2Q	CT3Q	CT5Q	
Helena Actalyke®						3
Helena Cascade POC						3
IL Gem® PCL ACT						3
IL Gem PCL ACT-LR						3
IL GEM PCL Plus ACT						3
IL GEM PCL Plus ACT-LR						3
ITC Hemochron® CA510/FTCA510						3
ITC Hemochron FTK-ACT						3
ITC Hemochron Jr. Signature/ACT+						3
ITC Hemochron Jr. Signature/ACT-LR						3
ITC Hemochron P214/P215						3
i-STAT® Celite® and Kaolin ACT						3
Medtronic HemoTec ACT/ACTII/ACT Plus HR-ACT						3
Medtronic HemoTec ACT/ACTII/ACT Plus LR-ACT						3
Medtronic HemoTec ACT/ACTII/ACT Plus R-ACT						3
Medtronic Hepcon HMS, HMS Plus						3
Sienco Sonoclot®						3

These programs do not meet regulatory requirements for proficiency testing; see Surveys CT-CT3 and CT5 on page 162. For additional information about the CAP Quality Cross Check program, see page 42.

The Quality Cross Check Program:

- Provides a solution for monitoring performance across multiple instruments, and is in compliance with the CMS directive regarding proficiency testing on multiple instruments.
- Simplifies instrument comparability efforts by providing custom reports with both peer group comparison and instrument comparability statistics.

- CTQ Three 3.0-mL lyophilized whole blood specimens in triplicate with corresponding diluents
- CT1Q Three 1.7-mL lyophilized whole blood specimens in triplicate with corresponding diluents
- CT2Q Three 0.5-mL lyophilized whole blood/ diluent ampules in triplicate
- CT3Q Three 0.5-mL lyophilized whole blood/ diluent ampules in triplicate
- CT5Q Three 1.7-mL lyophilized whole blood specimens in triplicate with corresponding diluents
- Report up to three instruments
- Two shipments per year

Platelet Function* PF, PF1						
Instrument/Method Program Code Challenges/Shipment						
	PF	PF1				
Platelet aggregation			2			
PFA-100			2			
Helena Plateletworks®			2			

^{*}This Survey requires the draw of a normal donor sample.

Viscoelastometry TEG						
Instrument/Method	Program Code	Challenges/Shipment				
	TEG					
Viscoelastometry	ı	2				

- PF Four 3.2% sodium citrate vacuum tubes; two 10.0-mL plastic tubes
- PF1 Four 3.2% sodium citrate vacuum tubes; two 10.0-mL plastic tubes
- Two shipments per year

Program Information

- Two 1.0-mL lyophilized whole blood specimens with diluents
- For use with the Haemonetics[™]
 Thromboelastograph[®], including TEG5000 and TEG6s, ROTEM[®] delta hemostasis analyzers
- Two shipments per year

Coagulation Calibration Verification/Linearity LN35, LN36, LN37 **Analyte Program Code** LN35 LN36 LN37 **Target Ranges** Antithrombin activity 10%-130% Protein C activity 10%-100% Heparin, low molecular weight 0.1-2.0 U/mL Heparin, unfractionated 0.1-1.3 U/mL 5%-140% von Willebrand factor antigen

The LN35, LN36, and LN37 CVL programs meet the CAP Accreditation requirements HEM.38009, 38010, and 38011.

LN Express service is available.

- LN35, LN37 Six 1.0-mL frozen plasma specimens per mailing
- LN36 Twelve 1.0-mL frozen plasma specimens per mailing, which include six for low molecular weight heparin and six for unfractionated heparin
- Two shipments per year; ships on dry ice

D-Dimer Calibration Verification/Linearity LN42							
Analyte	Analyte Program Code						
	LN42	Target Range					
D-dimer	•	220-5,500 ng/mL FEU					

LN Express service is available.

Fibrinogen Calibration Verification/Linearity LN44						
Analyte	Analyte Program Code					
LN44 Target Range						
Fibrinogen		80-900 mg/dL				

LN Express service is available.

Drug-Specific Platelet Aggregation PIA, PIAX Procedure **Program Code** Challenges/Shipment PIA PIAX Aspirin assay 3 3 PRU test 3 llb/llla assay

Program Information

- Six 1.0-mL plasma specimens
- Two shipments per year



Program Information

- Six 1.0-mL frozen plasma specimens
- Two shipments per year; ships on dry ice

- PIA Three lyophilized specimens with diluents
- PIAX All Survey PIA specimens in duplicate
- For use with the Accumetrics VerifyNow® System
- · Kit includes sufficient material to perform one assay; multiple assay reporting requires the purchase of PIAX
- Two shipments per year

Whole Blood Coagulation WP3, WP4, WP6, WP9, WP10						
Analyte Challenges/Shipment						
	Program Code					
	WP3 WP4 WP6 WP9 WP10					
International normalized ratio (INR)	5 5 5 5 3					
Prothrombin time	5	5	5	5	-	

For method compatibility, see instrument matrix below.

Program Information

- WP3 Five 1.0-mL lyophilized plasma specimens with corresponding diluents
- WP4, WP6 Five 0.5-mL unitized lyophilized blood specimens
- WP9 Five 0.3-mL lyophilized plasma specimens
- Three shipments per year
- WP10 Three 0.3-mL lyophilized plasma specimens with corresponding diluents; two shipments per year

Whole Blood Coagulation, Instrument Matrix

Instrument		Pro	gram C	ode	
	WP3	WP4	WP6	WP9	WP10
Abbott CoaguSense™					
Helena Cascade POC – Citrated					
Helena Cascade POC – Noncitrated					
IL GEM PCL, PCL Plus – Citrated					
IL GEM PCL, PCL Plus – Noncitrated					
ITC Hemochron Jr. Signature/Signature +, Signature Elite and Jr. II – Citrated cuvette					
ITC Hemochron Jr. Signature/Signature +, Signature Elite and Jr. II – Noncitrated cuvette					
i-STAT					
Roche CoaguChek XS Plus and XS Pro					
Roche CoaguChek XS System					

11-dehydrothromboxane B2 TBX						
Analyte	Program Code	Challenges/Shipment				
	TBX					
11-dehydrothromboxane B2	ı	3				

- Three 0.5-mL lyophilized urine specimens
- For use with Aspirin Works
- Two shipments per year

Platelet Mapping* PLTM						
Analyte Program Code Challenges/Shipment						
PLTM						
AA % aggregation/inhibition	I	2				
ADP % aggregation/inhibition		2				

^{*}This Survey requires the draw of a normal donor sample.

Improve the reliability of your patient results with CAP Survey Validated Materials

Use the same material that is sent in the Surveys program to:

- · Identify and troubleshoot instrument/method problems
- · Correlate results with other laboratories or instruments
- Document correction of problems identified in Surveys
- · Utilize material with confirmed results as an alternative external quality control
- · Identify potential proficiency testing failures

Each laboratory receives a Survey Participant Summary, which includes readily available results.

Coagulation—Limited, Validated Material

Validated Material	Program Code	Corresponding Survey	Page
Coagulation	CGM	CGL	158

Program Information

- One 3.2% sodium citrate and two heparin vacuum tubes; two 3.5-mL plastic tubes; one vial of 0.2M CaCl₂
- For use with the Haemonetics Platelet Mapping[®] assay
- · Two shipments per year

Program Information

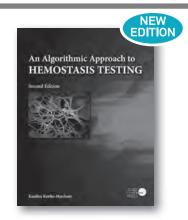
 Five 1.0-mL lyophilized plasma specimens; three shipments per year; one 1.0-mL lyophilized plasma specimen and one 1.0-mL serum specimen; two shipments per year

Give better consultations for hemostasis diagnosis

- New chapters on emergency assessment, consultation, antifibrinolytic and thrombolytic agents, and more
- Insightful case studies
- · Detailed algorithms to assist in diagnosis

View sample pages and order online:

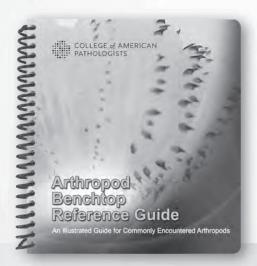
- · printed books at estore.cap.org
- · ebooks at ebooks.cap.org

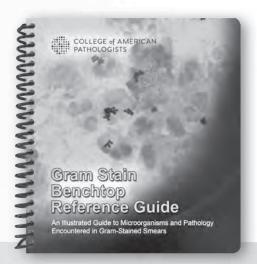


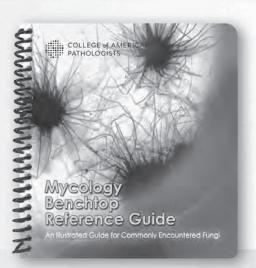
Item number: PUB223 Hardcover; 480 pages; 175+ figures, tables, and

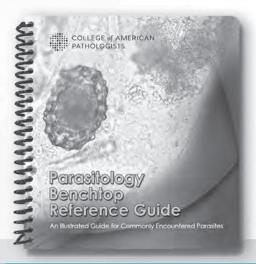
algorithms; 2016

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ABRG	Arthropod Benchtop Reference Guide
GSBRG	Gram Stain Benchtop Reference Guide
MBRG	Mycology Benchtop Reference Guide
PBRG	Parasitology Benchtop Reference Guide

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15 Microbiology



Ensure accuracy of Zika testing.

Our new Vector-Borne Disease Molecular (VBDM) Survey is intended for laboratories performing Zika testing using nucleic acid amplification methodologies.

- Add a level of quality assurance to ensure you deliver accurate test results.
- Provide an additional opportunity for your laboratory staff to fulfill competency requirements.

Microbiology

Bacteriology	170
Mycobacteriology	183
Mycology	184
Parasitology	187
Virology	191
Molecular Microbiology	
Infectious Disease Serology	
New Programs Expanded Parasitology (PEX)	189
MRSA Screen, Molecular, 2 Challenge (MRS2M)	179
MRSA Screen, Molecular, 5 Challenge (MRS5M)	179
Vector-Borne Disease—Molecular (VBDM)	
New Analyte/Drug Additions NEW	
Gastrointestinal Panel (GIP)	199

Adenovirus

Astrovirus

Cyclospora cayetanensis

Enteroaggregative E coli (EAEC)

Enteropathogenic *E coli* (EPEC)

Plesiomonas shigelloides

Sapovirus

Vibrio cholerae

Yersinia enterocolitica

Bacteriology

Analytes/procedures in **bold** type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Guide for Ordering Appropriate Bacteriology Surveys

Procedure	Program Code									
	D	D4	MC1	MC2	MC5	D2	D7	D3	MC4	D1
Bacterial identification	ı	1	ı							
Gram stain	ı	ı	ı					ı		
Antimicrobial susceptibility testing	I	ı	ı							
Bacterial antigen detection	I	ı	ı						I	

Participants must report five specimens for each mailing to meet CLIA requirements for the subspecialty of bacteriology. See the following pages for more detailed information about each Survey.

Bacteriology D				
Procedure	Program Code	Challenges/Shipment		
	D			
Antimicrobial susceptibility testing	ı	1 graded, 1 ungraded		
Bacterial antigen detection	ı	2		
Bacterial identification	ı	5		
Gram stain	I	1		

Additional Information

Antigen detection challenges will be included in the following shipments:

- Shipment A: C. difficile antigen/toxin and spinal fluid meningitis panel
- · Shipment B: Spinal fluid meningitis panel and Group A Streptococcus
- Shipment C: C. difficile antigen/toxin and Group A Streptococcus

Program Information

- Five swab specimens with diluents in duplicate for culture
- Culture sources may include wounds, blood, respiratory, urines, stools, and anaerobes on a rotational basis
- Two specimens for bacterial antigen detection from the following:

One swab for Group A Streptococcus

One 1.0-mL lyophilized specimen for spinal fluid meningitis testing

One 0.5-mL lyophilized specimen for *Clostridium* difficile, for use with rapid or molecular testing methods

· Three shipments per year







Expanded Bacteriology DEX				
Analyte	Program Code	Challenges/Shipment		
	DEX			
Live organisms	I	2		

Additional Information

Expanded Bacteriology (DEX) is an educational opportunity that provides:

- Culture and susceptibility testing challenges for microbiology laboratories that perform complete identification and susceptibility of bacterial isolates including less common or problematic bacteria
- More exposure to emerging bacterial pathogens and novel resistance mechanisms
- Ability to recognize and identify organisms that exhibit multiple drug-resistance patterns
- Recovery and identification of mixed pathogens such as yeast, aerobic, and anaerobic bacteria in cultures containing multiple organisms

Program Information

- Two swab specimens in duplicate with diluents to perform bacterial identification and susceptibility (when directed)
- · Three shipments per year



Microbiology Bench Tools Competency MBT				
Procedure Program Code Challenges/Shipme				
	MBT			
Bacterial identification	•	6		
Antimicrobial susceptibility testing	I	2		

Additional Information

Microbiology Bench Tools Competency (MBT) is a supplemental module for competency assessment and an educational resource for microbiology laboratories. The module:

- Provides organisms that challenge the basic elements of testing at the microbiology bench, including direct observation, monitoring, recording, and reporting of test results
- Can be used for both competency and educational purposes, including teaching and training pathology residents, new employees, and medical and MT/MLT students
- · Provides identification and susceptibility results for supervisor use

This is not a proficiency testing program and participants will not return results to the CAP.

Program Information

- Six swab specimens with diluents for bacterial identification and susceptibility
- Culture sources will vary with each shipment
- Results will be provided with the kit to assess personnel competency
- · Two shipments per year





GC, T	hroat, and	l Urine Cu	Itures D1	, D2, D3, E	7	
Procedure		Program Code				
	D1	D2	D3	D7		
Antimicrobial susceptibility testing		ı			1	
Bacterial identification				ı	5	
Gram stain					1	
Culture source:	Throat	Urine	Cervical	Throat/Urine		
Microbiologic level:	Presence or absence of Group A Streptococcus determination	Organisms identified to the extent of your laboratory's protocol	Presence or absence of Neisseria gonorrhoeae determination	Combination of two throat and three urine culture specimens		

- D1- Five swab specimens with diluents in duplicate
- D2 Five loop specimens with diluents in duplicate, with one susceptibility challenge, and one Gram stain challenge
- D3 Five loop specimens with diluents in duplicate, and one Gram stain challenge
- D7 Two swab specimens with diluents in duplicate, three loop specimens with diluents in duplicate, one susceptibility challenge, and one Gram stain challenge
- Throat swabs compatible with molecular- and culture-based methods
- Three shipments per year





Identify microorganisms quickly and confidently

Gram Stain Benchtop Reference Guide is an illustrated guide to gram-positive and gram-negative organisms. Its rugged construction is well suited for students and medical technologists for heavy use at the workbench.

Features include:

- Theory and application of the Gram stain
- Detailed descriptions of microbial morphology, quantitation, and indicators of pathology
- Examples of more than 35 gram-positive and gram-negative organisms found in blood, body fluids, CSF, urine, and the genital and respiratory tracts
- Seven tabbed sections for easy reference

This sturdy, spiral-bound, laminated guide is conveniently sized at 6½" x 7".

Choose code GSBRG on your Surveys order form.



Bacteriology—Limited D4				
Procedure	Program Code	Challenges/Shipment		
	D4			
Antimicrobial susceptibility testing		1		
Rapid group A Streptococcus antigen detection*	ı	1		
Bacterial identification	I	5		
Gram stain	I	1		
Culture source:	Microbiologic level:			
GC culture	Presence or absence of Neisseria gonorrhoeae determination			
Throat culture Presence or absence of Group A Streptococcus determination				
Urine culture	Organisms identified to the extent of your laboratory's protocol			

^{*}If you are using a waived method for *Streptococcus* testing, these results will not count toward the required five challenges for the subspecialty of microbiology.

- Five loop/swab specimens with diluents in duplicate, and one swab specimen
- Two throat culture specimens, two urine culture specimens, one GC culture specimen, and one bacterial antigen detection specimen
- Throat swabs compatible with molecular- and culture-based methods
- · Three shipments per year





Microbiology—Combination w/GC MC1, MC2			
Procedure	Challenges/Shipment		
	Progra	m Code	
	MC1	MC2	
Antimicrobial susceptibility	1	1	
GC culture	5		
Gram stain	2	1	
Rapid group A Streptococcus antigen detection*	1	1	
Throat culture	2	3	
Urine culture	3	5	

^{*}If you are using a waived method for *Streptococcus* testing, these results will not count toward the required five challenges for the subspecialty of microbiology.

Program Information

- MC1 Eight loop specimens with diluents in duplicate, two swab specimens with diluents in duplicate, and one swab specimen for antigen detection
- MC2 Five loop specimens with diluents in duplicate, three swab specimens with diluents in duplicate, and one swab specimen for antigen dectection
- Urine cultures will only have one susceptibility challenge
- Throat swabs compatible with molecular- and culturebased methods
- Three shipments per year







Urine Colony Count	MC3, MC4	
Procedure	Challenges	S/Shipment
	Progra	m Code
	мсз	MC4
Urine colony count/urine culture identification	2	5
Rapid group A Streptococcus antigen detection*		3
Throat culture		3

^{*}If you are using a waived method for *Streptococcus* testing, these results will not count toward the required five challenges for the subspecialty of microbiology.

- MC3 Two 2.0-mL urine specimens with diluents
- MC4 Five 2.0-mL urine specimens with diluents, three swab specimens (in duplicate) with diluents, and three swab specimens for antigen detection
- Throat swabs compatible with molecular- and culturebased methods
- Three shipments per year





Throat & Urine Culture/Rapid Strep A Antigen Detection MC5				
Procedure	Program Code	Challenges/Shipment		
	MC5			
Antimicrobial susceptibility	I	1		
Gram stain		1		
Rapid group A Streptococcus antigen detection*		2		
Throat culture		3		
Urine culture		3		

^{*}If you are using a waived method for *Streptococcus* testing, these results will not count toward the required five challenges for the subspecialty of microbiology.

Program Information

- Three loop specimens with diluents in duplicate, three swab specimens with diluents in duplicate, and two swab specimens for antigen detection
- Urine cultures will only have one susceptibility challenge
- Throat swabs compatible with molecular- and culturebased methods
- · Three shipments per year





Gram Stain D5		
Procedure Program Code Challenges/Shipment		
	D5	
Gram stain	I	5

Program Information

- Five air-dried, methanolfixed unstained glass slides
- · Three shipments per year





Virtual Gram Stain Competency VGS1, VGS2			
Procedure	Program Code Challenges/Shipment		
	VGS1	VGS2	
Virtual gram stain basic			3
Virtual gram stain advanced			3

Additional Information

- Virtual Gram Stain Basic Competency (VGS1) is for general and new laboratory technologists/technicians. Participants will assess the quality of specimens and stains and will report artifacts and detailed gram-positive and gram-negative morphology. Challenges will include specimens such as CSF, body fluids, and positive blood cultures.
- Virtual Gram Stain Advanced Competency (VGS2) is for experienced laboratory technologists/technicians and microbiologists. Participants will receive challenging images of sputum, body fluids, and other specimens to assess the quality, quantity, and typical morphology of both gram-positive and gram-negative organisms appropriate for the site.
- See system requirements on page 13.

Program Information

- Three online whole slide images
- Results included in the kit to assess personnel competency
- Powered by DigitalScope® technology
- Two shipments per year

Rapid Group Strep A Antigen Detection D6				
Procedure Program Code Challenges/Shipment				
	D6			
Group A Streptococcus antigen detection*	ı	5		

^{*}If you are using a waived method for *Streptococcus* testing, these results will not count toward the required five challenges for the subspecialty of microbiology.

Program Information

- · Five swab specimens
- Not compatible with molecular- and culturebased methods
- Three shipments per year



Rapid Strep A Antigen Detection, Waived D9			
Procedure Program Code Challenges/Shipment			
	D9		
Group A Streptococcus antigen detection	ı	2	

- Two swab specimens
- Not compatible with molecular- and culturebased methods
- Two shipments per year

15

Group B Strep Detection D8			
Procedure Program Code Challenges/Shipment			
	D8		
Group B Streptococcus detection	I	5	

Program Information

- Five swab specimens with diluents
- Compatible with molecularand culture-based methods
- · Three shipments per year



Bacterial Antigen D	etection	ı LBAS,	SBAS
Procedure	Progra	m Code	Challenges/Shipment
	LBAS	SBAS	
Legionella pneumophila antigen detection			2
Streptococcus pneumoniae antigen detection			2

Program Information

- Two 0.5-mL liquid simulated clinical specimens
- Two shipments per year

Blood C	ulture BCS	
Procedure Program Code Challenges/Shipment		
	BCS	
Blood culture bacterial detection and identification	ı	2

Program Information

- Two specimens with diluents for inoculation of blood culture bottles
- · Two shipments per year



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Blood Culture, Staphylococcus aureus BCS1			
Analyte Program Code Challenges/Shipment			
	BCS1		
Staphylococcus aureus/MRSA/MSSA	I	3	

- Three specimens with diluents for inoculation of blood culture bottles
- Compatible with molecular methods for rapid detection of S. aureus/MRSA/MSSA from positive blood culture bottles
- Two shipments per year



PNA FISH	PNA1,	PNA2	
Analyte	Progra	m Code	Challenges/Shipment
	PNA1	PNA2	
Staphylococcus			3
Yeast			3

Program Information

- Three specimens with diluents for inoculation of blood culture bottles
- Two shipments per year



Bordetella pertussis/parapertussis, Molecular BOR				
Analyte Program Code Challenges/Shipment				
	BOR			
Bordetella pertussis	ı	3		
Bordetella parapertussis	I	3		

Program Information

- Three swab specimens
- Designed for molecular techniques
- Two shipments per year

Campylol	pacter CAMP	
Analyte	Program Code	Challenges/Shipment
	CAMP	
Campylobacter	I	2

Program Information

- Two swabs with diluents in duplicate
- For use with rapid antigen, culture-based testing, and molecular methods
- Two shipments per year





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Clostridium difficile, 2 Challenge CDF2			
Procedure	Program Code	Challenges/Shipment	
	CDF2		
Clostridium difficile	ı	2	

Clostridium difficile, 5 Challenge CDF5			
Procedure	Program Code	Challenges/Shipment	
	CDF5		
Clostridium difficile	ı	5	

Chlamydia Antigen Detection HC1, HC3				
Procedure	Program Code		Challenges/Shipment	
	HC1	HC3		
Antigen detection (DFA)			5	
Antigen detection (EIA)			5	

Fecal Lactoferrin FLAC				
Analyte	Program Code	Challenges/Shipment		
	FLAC			
Fecal lactoferrin		3		

Helicobacter pylori Antigen, Stool HPS			
Procedure	Program Code	Challenges/Shipment	
	HPS		
Helicobacter pylori antigen detection		2	

- Two 0.5-mL lyophilized specimens, for use with rapid or molecular testing methods
- Two shipments per year

Program Information

- Five 0.5-mL lyophilized specimens, for use with rapid or molecular testing methods
- · Three shipments per year

Program Information

- HC1 Five 5-well slide specimens; for the detection of chlamydial elementary bodies by DFA
- HC3 Five 2.0-mL liquid specimens for Chlamydia antigen testing by EIA
- Three shipments per year

Program Information

- Three 0.5-mL simulated stool specimens
- For use with rapid methods
- Two shipments per year

Program Information

- Two 0.5-mL fecal suspensions
- Two shipments per year





Methicillin Resistant S. aureus, 2 Challenge MRS			
Procedure Program Code Challenges/Shipmen			
	MRS		
MRSA/MSSA detection	ı	2	

- Two swab specimens with diluents
- For use with culture-based testing
- Two shipments per year



MRSA Screen, Molecular,	2 Challenge Mi	RS2M NEW
Procedure	Program Code	Challenges/Shipment
	MRS2M	
MRSA/MSSA/SA detection	I	2

Program Information

- Two swab specimens (in duplicate)
- For use with molecular methods only
- Two shipments per year

Methicillin-Resistant Staphylococcus aureus Screen MRS5		
Procedure	Program Code	Challenges/Shipment
	MRS5	
MRSA/MSSA detection	ı	5

Program Information

- Five swab specimens with diluents
- For use with culture-based testing
- Three shipments per year



MRSA Screen, Molecular,	5 Challenge M	RS5M (NEW)
Procedure	Program Code	Challenges/Shipment
	MRS5M	
MRSA/MSSA/SA detection	•	5

Program Information

- Five swab specimens (in duplicate)
- For use with molecular methods only
- Three shipments per year



Laboratory Preparedness Exercise LPX		
Analyte	Program Code Challenges/Shipment	
	LPX	
Live organisms		3

Additional Information

The Laboratory Preparedness Exercise (LPX) was developed as a collaborative effort between the College of American Pathologists, the Centers for Disease Control and Prevention (CDC), and the Association of Public Health Laboratories (APHL). Laboratories will be sent live organisms that either exhibit characteristics of bioterrorism agents or demonstrate epidemiologic importance and will be expected to respond following Laboratory Response Network Sentinel Laboratory Guidelines if a bioterrorism agent is suspected. All agents provided are excluded from the CDC's select agent list. These may include strains of *Bacillus anthracis*, *Yersinia pestis*, *Francisella tularensis*, and *Brucella abortus* that have been modified and are safe for testing in a laboratory that contains a certified Class II Biological Safety Cabinet and is capable of handling Category A and B agents.

Rapid U	Irease RUR	
Analyte	Program Code	Challenges/Shipment
	RUR	
Urease		3

Stool Pathogen SP, SPN, SP1				
Analyte	Pro	ogram Co	de	Challenges/Shipment
	SP	SPN	SP1	
Adenovirus 40/41				2
C. difficile				2
Rotavirus				2
Shiga toxin				2
Norovirus				1

Program Information

- Three swab specimens with diluents
- Not available to international customers due to United States export law restrictions
- · Two shipments per year







Program Information

- Three simulated gastric biopsy specimens
- For use with methods such as CLOTEST®
- Two shipments per year

Program Information

- SP Two 1.0-mL liquid specimens; for use with rapid or molecular testing methods; not available to international customers due to United States export law restrictions
- SPN Two 1.0-mL liquid specimens; for use with rapid or molecular testing methods; intended for international laboratories
- SP1 One 1.0-mL liquid specimen compatible with molecular methods only
- · Two shipments per year



Shiga Toxin ST		
Analyte	Program Code Challenges	
	ST	
Shiga toxin		2

- Two 0.5-mL liquid specimens
- For use with direct shiga toxin testing only; not compatible with culture methods, cytotoxicity assays, or PCR
- Not available to international customers due to United States export law restrictions
- Two shipments per year

Bacterial Vaginosis BV		
Procedure	Program Code	Challenges/Shipment
	BV	
Bacterial vaginosis detection	I	3

Program Information

- Three 1.0-mL liquid specimens
- For OSOM® BVBlue users
- Two shipments per year

Vaginitis S	creen \	/S, VS1	
Analyte	Progra	m Code	Challenges/Shipment
	VS*	VS1**	
Candida sp.			5
Gardnerella vaginalis			5
Trichomonas vaginalis			5

^{*}The biohazard warning applies to Survey VS.

Program Information

 VS - Five swabs for DNA probe technology; BD Affirm™ VP III probe detection method; three shipments per year



 VS1 - Five swabs for methods such as Sekisui OSOM Trichomonas Rapid Test, Trichomonas vaginalis methods; two shipments per year



^{**}Molecular users are encouraged to use *Trichomonas vaginalis*, Molecular TVAG on page 189.

Vaginitis Screen, Virtual Gram Stain VS2		
Procedure	Program Code	Challenges/Shipment
	VS2	
Interpretation of Gram-stained vaginal smears	ı	3

Additional Information

• See system requirements on page 13.

•	Three online
	!

whole slide images

Program Information

- · Powered by DigitalScope technology
- Two activities per year; your CAP shipping contact will be notified via email when the activity is available

Vancomycin-Resistant <i>Enterococcus</i> VRE		
Procedure	Program Code	Challenges/Shipment
	VRE	
Vancomycin-resistant Enterococcus (VRE) detection		2

Program Information

- Two swabs with diluents
- For use with molecular methods and culture-based testing
- Two shipments per year





Mycobacteriology

Analytes/procedures in **bold** type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Mycobacteriology E		
Procedure	Program Code	Challenges/Shipment
	E	
Acid-fast smear	I	1
Antimycobacterial susceptibility testing	I	1 graded, 1 ungraded
Mycobacterial identification*		5

^{*}This procedure requires identification of Mycobacterium tuberculosis.

Program Information

- Five simulated clinical isolates with diluents and one specimen for performing an acid-fast bacillus smear
- Identification may be performed by culture or molecular methods
- · Two shipments per year



Mycobacteriology—Limited E1		
Procedure	Program Code	Challenges/Shipment
	E1	
Acid-fast smear	ı	5
Mycobacterial culture	ı	5

Program Information

- Five simulated specimens for acid-fast smears and/or for the determination of the presence or absence of acid-fast bacillus by culture
- · Two shipments per year



Molecular MTB Detection and Resistance MTBR		
Procedure Program Code Challenges/Shipment		
	MTBR	
Mycobacterium tuberculosis detection	ı	3
Rifampin resistance	ı	3

Program Information

- Three 1.25-mL simulated sputum specimens for use with molecular methods
- · Not suitable for culture
- · Two shipments per year



Mycology

Analytes/procedures in **bold** type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Mycology and Aerobic Actinomycetes F		
Procedure	Program Code	Challenges/Shipment
	F	
Antifungal susceptibility testing	ı	1
Cryptococcal antigen detection	ı	2 per year
Mold and yeast identification	ı	5

Program Information

- Five loops for culture with diluents in duplicate and one 1.0-mL simulated cerebrospinal fluid specimen (A and B shipments only)
- Identification of yeasts, molds, and aerobic actinomycetes may be performed by molecular- and culture-based methods
- · Three shipments per year





Yeast F1		
Procedure	Program Code	Challenges/Shipment
	F1	
Antifungal susceptibility testing	I	1
Cryptococcal antigen detection		2 per year
Yeast identification	ı	5

Program Information

- Five loops for culture with diluents in duplicate and one 1.0-mL simulated cerebrospinal fluid specimen (A and B shipments only)
- Identification of yeast may be performed by molecularand culture-based methods
- · Three shipments per year





Candida Culture F3		
Procedure	Program Code	Challenges/Shipment
	F3	
Yeast identification	ı	5

- Five loops for culture with diluents in duplicate
- Identification of Candida species may be performed by culture, molecular, and rapid methods
- Three shipments per year



Cryptococcal Antigen Detection CRYP		
Analyte	Program Code	Challenges/Shipment
	CRYP	
Cryptococcal antigen detection	ı	5

Program Information

- Five 1.0-mL simulated cerebral spinal fluids
- Three shipments per year

Galactomannan FGAL		
Analyte	Program Code	Challenges/Shipment
	FGAL	
Galactomannan - Aspergillus		3

Program Information

- Three liquid specimens
- For use with methods such as Bio-Rad Platelia™
- Two shipments per year

Fungal Serology FSER		
Procedure	Program Code	Challenges/Shipment
	FSER	
Serological detection of specific fungal antibodies	ı	3

Program Information

- Three serum specimens
- For use with immunodiffusion methods
- Designed for the detection of antibodies to Aspergillus, Blastomyces, Coccidiodes, and Histoplasma
- Two shipments per year



15

Fungal Smear FSM		
Procedure	Program Code	Challenges/Shipment
	FSM	
KOH preparation/calcofluor white	ı	3

India Ink IND		
Procedure	Program Code	Challenges/Shipment
	IND	
India ink	I	2

Pneumocystis PCP1, PCP2, PCP4				
Procedure	Program Code Challenges/Shipmen			
	PCP1	PCP2	PCP4	
PCP – Calcofluor white stain				3
PCP – DFA stain				3
PCP – GMS stain			ı	3

Program Information

- · Three slides
- Two shipments per year

Program Information

- Two liquid specimens
- Two shipments per year

Program Information

- Three images, each available as photographs and online images for Pneumocystis jirovecii
- Two shipments per year

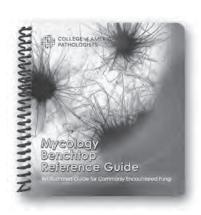
Find fungi fast

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- Portable (6½" x 7")
- Durable, water-resistant laminated format to withstand years of benchtop use

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Item number: MBRG Spiral bound; 92 pages; 70+ images; 2013

Parasitology

Analytes/procedures in **bold** type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Parasitology P, P3	, P4, P	5		
Procedure		Prograi	m Code	
	Р	Р3	P4	P5
Fecal suspension (wet mount)				
Fecal suspension (Giardia and/or Cryptosporidium immunoassay and modified acid-fast stain)	ı	ı		•
Giemsa-stained blood smear				
Preserved slide (for permanent stain)			I	

Additional Information

- The proficiency testing materials used for the Parasitology Surveys contain formalin as a preservative.
- Modified acid-fast stain results do not meet CLIA requirements for parasite identification.

- P Five specimens consisting of thin and thick films for blood and tissue parasite identification; preserved slides for permanent stain; 0.75-mL fecal suspensions for direct wet mount examination, photographs, and/or online images; two 0.75-mL fecal suspensions for Giardia and/or Cryptosporidium immunoassay testing and modified acid-fast stain
- P3 Five 0.75-mL fecal suspensions for direct wet mount examination, photographs, and/or online images; one 0.75-mL fecal suspension for Giardia and/or Cryptosporidium immunoassay testing and modified acid-fast stain
- P4 Five specimens
 consisting of 0.75-mL fecal
 suspensions for direct wet
 mount examination, preserved
 slides for permanent stain,
 photographs, and/or online
 images; one 0.75-mL fecal
 suspension for Giardia
 and/or Cryptosporidium
 immunoassay testing and
 modified acid-fast stain
- P5 Five 0.75-mL fecal suspensions for Giardia and/or Cryptosporidium immunoassay testing and modified acid-fast stain
- Three shipments per year



15

Blood Parasite BP				
Procedure	Program Code	Challenges/Shipment		
	ВР			
Thin/thick blood film sets*	ı	5		

^{*}This Survey will include corresponding thick films when available.

Program Information

- Five Giemsa-stained blood film sets, photographs, and/or online images
- A variety of blood parasites, including Plasmodium, Babesia, Trypanosoma, and filarial worms
- Three shipments per year

Rapid Ma	alaria RMAL	
Procedure	Program Code	Challenges/Shipment
	RMAL	
Rapid malaria detection	I	3

^{*}Detects *Plasmodium falciparum* specific histidine-rich protein 2 (HRP2). May not be compatible with methods that use pLDH enzyme detection for mixed malaria infections.

Program Information

- Three 0.5-mL antigen specimens
- Two shipments per year

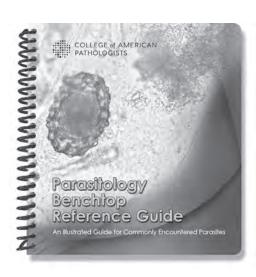
Parasitology Benchtop Reference Guide (PBRG)

- More than 70 identifications for parasites commonly encountered in the clinical laboratory
- Detailed descriptions of the parasite morphology, ecology, and clinical significance
- Color images of microscopic morphologies using routine parasitology stains and preparations
- Color images of macroscopic worms routinely submitted to the clinical laboratory
- Five tabbed sections for easy reference
 - Blood Parasites
 - o Intestinal Protozoa
 - o Intestinal Helminths
 - o Miscellaneous Specimens
 - Macroscopic Worms
- A durable and water-resistant format to withstand years of benchtop use—6½" x 7"

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Expanded Pa	arasitology PE	NEW
Procedure	Program Code	Challenges/Shipment
	PEX	
Parasite identification	I	3

This program provides an educational opportunity to challenge laboratory professionals' competency in the identification of parasites utilizing photo images.

Program Information

- Three images, each available as photographs and online images
- Two shipments per year

Ticks, Mites, and Other Arthropods TMO					
Procedure Program Code Challenges/Shipment					
ТМО					
Tick, mite, and arthropod identification	I	3			

Program Information

- Three images, each available as photographs and online images
- Two shipments per year

Trichomonas vaginalis, Molecular TVAG			
Analyte	Challenges/Shipment		
Trichomonas vaginalis	ı	3	

Program Information

- · Three liquid specimens
- Designed for molecular techniques
- Two shipments per year

Worm Iden	tification WID	
Procedure	Program Code	Challenges/Shipment
	WID	
Worm identification	I	3

Program Information

- Three images, each available as photographs and online images
- Two shipments per year

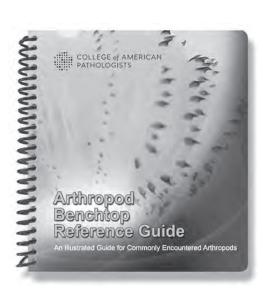
Arthropod Benchtop Reference Guide (ABRG)

- Numerous identifications of ectoparasites commonly encountered in the clinical laboratory
- Detailed descriptions of the most significant morphologic elements, ecology, and clinical significance
- A durable and water-resistant format to withstand years of benchtop use—6½" x 7"

Choose code ABRG on your Surveys order form.

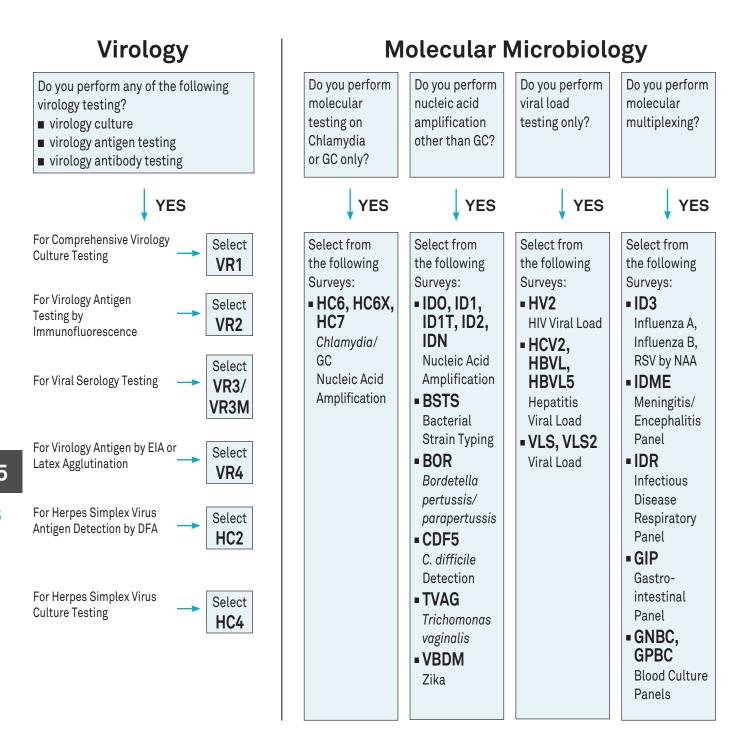
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Virology and Molecular Microbiology Testing

Use this flowchart as a guide for ordering the appropriate Virology and Molecular Microbiology Surveys for your laboratory's testing menu. For the subspecialty of virology, you must test five specimens per mailing. If you have any questions, please call the Customer Contact Center at 800-323-4040 or 847-832-7000 option 1.



Virology

Analytes/procedures in **bold** type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Virology Culture VR1				
Procedure	Program Code	Challenges/Shipment		
	VR1			
Chlamydia trachomatis culture	ı	1		
Viral isolation/identification	ı	5		
Educational challenge	ı	1 per year		

Program Information

- Five 0.5-mL specimens for viral culture and one 0.5-mL specimen for *Chlamydia* trachomatis culture
- · Three shipments per year



Virology Antigen Detection (DFA) VR2				
Analyte/Procedure	Program Code	Chall	enges/Shi _l	pment
	VR2	Α	В	С
Adenovirus antigen	ı	1	1	
Cytomegalovirus antigen	•	1	1	
Herpes simplex virus (HSV) antigen	I		1	1
Influenza A antigen	ı	1		1
Influenza B antigen	•		1	
Parainfluenza antigen	•	1		1
Respiratory syncytial virus (RSV) antigen	•	1		1
Varicella-zoster antigen	•		1	1
Educational challenge		1		

Program Information

- Five 5-well slide specimens
- Three shipments per year



15

Virology Antigen Detection (Non-DFA) VR4				
Analyte	Program Code	Challenges/Shipment		
	VR4			
Adenovirus (Not 40/41) antigen	I	5		
Influenza A antigen	I	5		
Influenza B antigen	ı	5		
Respiratory syncytial virus (RSV) antigen	ı	5		
Rotavirus antigen	ı	5		

Program Information

- Five 1.5-mL specimens
- For use with enzyme immunoassay and/or latex agglutination methods
- Three shipments per year

Herpes Simplex Virus HC2, HC4				
Procedure	Program Code Challenges/Shipment			
	HC2	HC4*		
Antigen detection (DFA)			5	
Culture			5	

^{*}The biohazard warning applies to Survey HC4.

Program Information

- HC2 Five 5-well slide specimens
- HC4 Five 0.5-mL lyophilized specimens
- Three shipments per year



Test your diagnostic skills as a pathologist with CPIP

Online, hands-on and interactive, the Clinical Pathology Improvement Program (CPIP) enables pathologists to sharpen their diagnostic skills in real time by working through an actual case. Each month, you will receive a new scenario, including slide images and clinical background. As the case unfolds, more information is revealed, just as in the laboratory. Participants who successfully complete the posttest may apply their earned credits to their MOC SAM requirements. Enjoy a full year of CPIP and earn up to 15 CME/SAM credits.

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Molecular Microbiology

Analytes/procedures in **bold** type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Hepatitis Viral Load	HCV2, HB	VL, HBVL5	5
Procedure Challenges/Shipment			nt
	Program Code		
	HCV2	HBVL	HBVL5
HCV genotyping	1		
HCV, qualitative	1		
HCV viral load	5		
HBV viral load		3	5

Program Information

- HCV2 Five 1.5-mL liquid plasma specimens; three shipments per year
- HBVL Three 1.25-mL plasma specimens; two shipments per year
- HBVL5 Five 2.0-mL plasma specimens; three shipments per year

HIV Viral Load HV2, HIVG			
Procedure	Progra	m Code	Challenges/Shipment
	HV2	HIVG	
HIV-RNA viral load			5
HIV genotyping			1

Program Information

- HV2 Five 2.5-mL plasma specimens
- HIVG One 1.0-mL plasma specimen
- Three shipments per year

Viral Load	VLS,	/LS2	
Procedure	Progra	m Code	Challenges/Shipment
	VLS	VLS2	
BK viral load			2
CMV viral load			2
EBV viral load			2
Adenovirus viral load			2
HHV6 viral load			2

- VLS Six 1.0-mL EDTA plasma specimens; two shipments per year
- VLS2 Ten 2.0-mL EDTA plasma specimens; three shipments per year

Viral Load Calibration Verification/Linearity LN38, LN39, LN45				
Analyte	Р	rogram Coo		
	LN38*	LN39	LN45	Target Ranges
CMV viral load	•			316-1.0M IU/mL
HIV viral load				50-5.0M IU/mL
HCV viral load				50-280M IU/mL
*The biohazard warning applies to Survey LN38.				
LN Express service is available.				

- LN38 Six 1.5-mL frozen plasma specimens
- Two shipments per year; ships on dry ice



- LN39 Six 2.5-mL plasma specimens
- LN45 Seven 2.5-mL frozen plasma specimens
- · Two shipments per year; ships on dry ice (dry ice does not apply to LN39)

C. trachomatis/GC by NAA HC6, HC6X, HC7			
Procedure	Program Code Challenges/Shipment		
	HC6,* HC6X*	HC7	
Nucleic acid amplification (NAA)			5
Nucleic acid amplification (NAA/DNA)		•	5

^{*}The biohazard warning applies to Surveys HC6 and HC6X.

Program Information

- HC6 Three swab specimens and two 1.0-mL simulated urine specimens
- HC6X Three swab specimens; two 1.0-mL simulated urine specimens in duplicate
- Three shipments per year



- HC7 Five 1.5-mL simulated body fluid specimens; designed for Cepheid users
- Three shipments per year



Human Papillomavirus HPV			
Analyte	Program Code Challenges/Shipment		
	HPV		
Human papillomavirus		2	

For laboratories using Digene, SurePath, and/or ThinPrep collection media, see page 274.

Program Information

- Two simulated cervical specimens contained in Digene transport media
- For Digene Hybrid Capture only
- Two shipments per year

Bacterial Strain Typing, Staphylococcus BSTS					
Analyte Program Code Challenges/Shipment					
	BSTS				
Staphylococcus 2					

Program Information

- Two sets of loops with diluents
- Two shipments per year



Vector-Borne Disease—Molecular VBDM				
Analyte Program Code Challenges/Shipme				
VBDM				
Zika virus	I	3		

Program Information

- Three 1.5-mL liquid specimens
- Two shipments per year

(September 1)

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Nucleic Acid Amplification, Organisms IDO, IDN			
Analyte/Procedure	Program Code Challenges/Shipment		
	IDO	IDN	
Bordetella pertussis/parapertussis	ı	•	1
Legionella pneumophila/ Chlamydophila pneumoniae*			1
Methicillin-resistant Staphylococcus aureus		•	1
Molecular typing (bacterial isolates)	ı		1
Mycobacterium tuberculosis	ı		1
Mycoplasma pneumoniae	ı		1
Vancomycin-resistant Enterococcus			1

^{*}Legionella pneumophila/Chlamydophila pneumoniae will be included in the following shipments:

- Shipment A: Chlamydophila pneumoniae
- Shipment B: Legionella pneumophila

- IDO Seven liquid or swab simulated clinical isolate specimens and two diluents
- IDN Six liquid or swab simulated clinical isolate specimens and two diluents; designed for international laboratories that cannot receive MTB
- Two shipments per year



Nucleic Acid Amplification, Viruses ID1, ID1T			
Analyte	Program Code		Challenges/Shipment
	ID1	ID1T	
Cytomegalovirus			1
Enterovirus			1
Epstein-Barr virus			1
Herpes simplex virus			1
Human herpesvirus 6			1
Human herpesvirus 8			1
Parvovirus B19			1
Varicella-zoster virus			1
BK virus			1
JC virus			1

Program Information

- ID1- Eight 1.0-mL liquid specimens
- ID1T Two 1.0-mL liquid specimens
- Two shipments per year



Nucleic Acid Amplification, Respiratory ID2				
Analyte	Program Code Challenges/Shipmen			
	ID2			
Adenovirus	I	1		
Coronavirus/Rhinovirus*		1		
Human metapneumovirus	I	1		
Influenza virus*	I	1		
Parainfluenza virus		1		
Respiratory syncytial virus (RSV)		1		

^{*}Coronavirus/Rhinovirus and Influenza virus will be included in the following shipments:

- Shipment A: Coronavirus and Influenza A
- Shipment B: Rhinovirus and Influenza B

Influenza A, Influenza B, and RSV by Nucleic Acid Amplification ID3						
Analyte	Program Code Challenges/Shipmen					
ID3						
Influenza A	ı	5				
Influenza B	■ 5					
Respiratory syncytial virus (RSV) ■ 5						

- Six 1.0-mL liquid specimens
- Two shipments per year

- Five 1.0-mL liquid specimens
- Designed for molecular multiplex panel users
- Three shipments per year

)		

Meningitis/Encephalitis Panel IDME					
Analyte	Program Code	Challenges/Shipment			
	IDME				
Escherichia coli K1	•	3			
Haemophilus influenzae		3			
Listeria monocytogenes	ı	3			
Neisseria meningitidis	ı	3			
Streptococcus agalactiae	ı	3			
Streptococcus pneumoniae	I	3			
Cytomegalovirus (CMV)	ı	3			
Enterovirus	ı	3			
Herpes simplex virus 1 (HSV-1)	I	3			
Herpes simplex virus 2 (HSV-2)	ı	3			
Human herpesvirus 6 (HHV-6)	I	3			
Human parechovirus	I	3			
Varicella-zoster virus (VZV)	ı	3			
Cryptococcus neoformans/gattii		3			

- Three 1.0-mL liquid specimens
- Designed for molecular multiplex panel users
- Two shipments per year

Infectious Disease, Respiratory Panel IDR						
Analyte	Program Code	Challenges/Shipment				
	IDR					
Adenovirus		5				
Bocavirus	•	5				
Bordetella (pertussis, parapertussis, bronchiseptica, holmesii)	ı	5				
Chlamydophila pneumoniae		5				
Coronavirus	•	5				
Human metapneumovirus	I	5				
Influenza A		5				
Influenza B	1	5				
Legionella pneumophila		5				
Mycoplasma pneumoniae		5				
Parainfluenza		5				
Parainfluenza 4		5				
Respiratory syncytial virus (RSV)	ı	5				
Rhinovirus/Enterovirus		5				

- Five 1.0-mL liquid specimens
- Designed for molecular multiplex panel users
- Three shipments per year

Gastrointestina	l Panel GIP	
Analyte	Program Code	Challenges/Shipment
	GIP	
Adenovirus NEW		3
Astrovirus NEW		3
Campylobacter	ı	3
Clostridium difficile, toxin A/B		3
Cryptosporidium		3
Cyclospora cayetanensis NEW	ı	3
Enteroaggregative <i>E coli</i> (EAEC) NEW		3
Enteropathogenic <i>E coli</i> (EPEC) NEW		3
Enterotoxigenic <i>E. coli</i> (ETEC) LT/ST	•	3
Entamoeba histolytica	•	3
Escherichia coli 0157		3
Giardia	•	3
Norovirus GI/GII		3
Plesiomonas shigelloides NEW		3
Rotavirus A	ı	3
Salmonella		3
Sapovirus NEW		3
Shiga-like toxin producing E. coli (STEC) stx1/stx2	ı	3
Shigella	1	3
Shigella/Enteroinvasive E. coli (EIEC)		3
Vibrio cholerae NEW		3
Yersinia enterocolitica NEW		3

- Three 1.0-mL simulated stool specimens
- Designed for molecular multiplex panel users
- Not available to international customers due to United States export law restrictions
- Two shipments per year

Blood Culture Panel	GNB	C, GPE	BC
Procedure	Progra	m Code	Challenges/Shipment
	GNBC	GPBC	
Identification of gram-negative organisms such as Acinetobacter, Citrobacter, Enterobacter, Proteus, Haemophilus, Klebsiella, Neisseria, Pseudomonas, Serratia, E. coli, and common resistance mechanisms isolated from positive blood culture bottles	ı		3
Identification of gram-positive organisms such as Staphylococcus, Streptococcus, Enterococcus, Listeria, and common resistance mechanisms isolated from positive blood culture bottles			3

These Surveys are not for the inoculation of blood culture bottles.

- Three 1.0-mL liquid simulated blood culture fluid specimens
- For laboratories using molecular multiplex panels
- Two shipments per year

Infectious Disease Serology

Analytes/procedures in **bold** type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Infectious Disease Serology VR3, VR3M				
Analyte	Progra	m Code	Challenges/Shipment	
	VR3	VR3M		
Cytomegalovirus (CMV) – IgG, IgM, and total antibodies	•		1	
Epstein-Barr virus (EBV) – VCA – IgG, IgM EBNA – IgG, IgM, and total antibodies EA – IgG	1		1	
Helicobacter pylori – IgG, IgA, and total antibodies	•		1	
Herpes simplex virus (HSV) – IgG antibody			1	
Mycoplasma pneumoniae — IgG, IgM, and total antibodies	•		1	
Mumps – IgG		•	1	
Rubeola virus (English measles) – IgG antibody	•		1	
Toxoplasma gondii — IgG, IgM, and total antibodies			1	
Varicella-zoster virus – IgG and total antibodies			1	

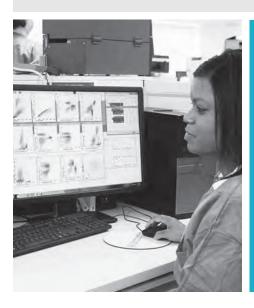
Program Information

- VR3 Eight 0.5-mL lyophilized defibrinated plasma specimens
- VR3M One 0.5-mL lyophilized defibrinated plasma specimen
- Two shipments per year

Tick-Transmitted Diseases TTD						
Analyte	Program Code Challenges/Sh					
	TTD					
Antibodies to tick-transmitted disease organisms	ı	3				

- Three 0.4-mL liquid specimens
- Designed for the detection of antibodies to Borrelia burgdorferi, Babesia microti, and Anaplasma phagocytophilum
- Two shipments per year

16 Immunology and Flow Cytometry



Immunology and flow cytometry testing is changing at a rapid pace—so is our proficiency testing.

Introducing three new programs for 2018:

- Testing of alpha-2-macroglobulin to evaluate patients with pancreatitis and nephrotic syndrome (A2MG)
- Detecting plasma cell neoplasms by flow cytometry (PCNEO)
- Testing for minimal residual disease by flow cytometry utilizing in silico challenges (BALL)

Immunology and Flow Cytometry

Immunology	202
Flow Cytometry	
New Programs NEW	
Alpha-2-Macroglobulin (A2MG)	204

Alpha-2-Macroglobulin (A2MG)	204
Flow Cytometry—B-ALL Minimal Residual Disease (BALL)	
Flow Cytometry—Plasma Cell Neoplasms (PCNEO)	211

Discontinued Programs

Heavy Chain/Light Chain Analysis (HCA)

Immunology

Analytes/procedures in bold type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Immunology ANA, ASO, CRP, HCG, IM, RF/RFX, RUB/RUBX, IL									
Analyte			Pı	ogram	Code)			Challenges/ Shipment
	ANA	AS0	CRP	HCG	IM	RF/ RFX	RUB/ RUBX	IL	
Antinuclear antibody (ANA)*	1							•	5
Antinuclear antibody ANA dry challenge									1
Antistreptolysin O (ASO)*		•							5
C-reactive protein, qualitative/quantitative								ı	2
hCG, serum (qualitative/ quantitative)	■ 5								
Infectious mononucleosis	I 5								
Rheumatoid factor*	■ ■ 5								
Rubella (IgG)*									5

^{*}Antinuclear antibody, Antistreptolysin O, Rheumatoid factor, and Rubella are regulated analytes and are graded for both qualitative and quantitative methods. Semiquantitative and/or titer results for these analytes are ungraded/educational in this Survey and do not meet regulatory requirements.

Program Information

- ANA and RUB Five 0.5-mL serum specimens
- ANA Three educational pattern interpretation dry challenges per year
- ASO, HCG, and RF Five 1.0-mL serum specimens
- CRP Two 0.5-mL serum specimens; not appropriate for high-sensitivity CRP (hsCRP) methods
- IM Five 0.6-mL serum specimens
- RFX All Survey RF specimens in duplicate
- RUBX All Survey RUB specimens in duplicate
- IL All immunology specimens except RFX and RUBX
- Three shipments per year



Immunology, General IG/IGX							
Analyte	Program Code	Challenges/Shipment					
	IG/IGX						
Alpha₁-antitrypsin	1	5					
Complement C3	ı	5					
Complement C4	1	5					
Haptoglobin	1	5					
IgA	I	5					
IgE	1	5					
IgG	1	5					
IgM	I	5					
Total kappa/lambda ratio		5					

- IG Ten 1.0-mL serum specimens
- IGX All Survey IG specimens in duplicate
- · Three shipments per year



Immunology, Special; Immunology Special, Limited; and H. pylori IgG Antibody S2, S4, S5

Analyte	Pro	gram C	ode	Challe	nges/Shi	pment
	S2	S4	S5	Α	В	С
Anticentromere antibody	1			1		1
Anti-DNA antibody double-stranded	ı	•		1	1	1
Antiglomerular basement membrane (GBM), IgG antibody	•				1	1
Antimitochondrial antibody	1			1	1	1
Antineutrophil cytoplasmic antibody (ANCA, anti-MPO, anti-PR3)	•			1	1	
Anti-RNP antibody				1	1	1
Anti-Sm antibody				1	1	1
Anti-Sm/RNP antibody	ı			1	1	1
Antismooth muscle antibody				1	1	1
Anti-SSA antibody	1			1	1	1
Anti-SSB antibody				1	1	1
Anti-SSA/SSB antibody				1	1	1
Antithyroglobulin antibody				1	1	1
Antithyroid microsomal antibody	ı			1	1	1
Antithyroid peroxidase antibody				1	1	1
Ceruloplasmin	1			1	1	1
Haptoglobin	ı			1	1	1
Helicobacter pylori, IgG antibody				1	1	
				2	2	
IgD	I	ı		1	1	1
IgG		•		1	1	1
IgG subclass proteins	ı	•		1	1	1
Prealbumin (transthyretin)	I	ı		1	1	1
Total kappa/lambda ratio	ı	•		1	1	1
Transferrin	•	•		1	1	1

Survey S2 is not appropriate for antimitochondrial antibody assays that are specific for the M2 antibody. Refer to Survey H on page 204.

Infectious Mononucleosis, Waived IMW Analyte **Program Code** Challenges/Shipment IMW ı 3 Infectious mononucleosis, waived

Program Information

- S2 A minimum of seven (0.5- to 1.0-mL/vial) serum specimens
- S4 A minimum of three (0.5- to 1.0-mL/vial) serum specimens
- S2 and S4 Three shipments per year
- S5 Two 1.0-mL serum specimens; two shipments per year



- Three 0.6-mL serum specimens
- Two shipments per year

Antichromatin antibody

Alpha-2-Macroglobulin A2MG				
Analyte	Program Code	Challenges/Shipment		
	A2MG			
Alpha-2-macroglobulin	I	3		

Antichromatin Antibody ACA				
Analyte	Program Code	Challenges/Shipment		
	ACA			

Antifilamentous Actin IgG Antibody FCN Analyte **Program Code** Challenges/Shipment **FCN** Antifilamentous actin (f-actin) 3 IgG antibody

Antihistone Antibody AHT			
Program Code	Challenges/Shipment		
AHT			
I	3		
	Program Code		

Antimitochondrial M2 Antibody H			
Analyte	Program Code	Challenges/Shipment	
	Н		
Antimitochondrial M2 antibody (AMA-M2)	ı	2	

Autoimmune Gastritis Markers APC				
Analyte	Program Code	Challenges/Shipment		
	APC			
Antiparietal cell antibody	I	2		
Anti-intrinsic factor antibody		2		

Program Information

- Three 0.5-mL serum specimens
- Two shipments per year

Program Information

- Three 0.5-mL serum specimens
- Two shipments per year

Program Information

- Three 0.5-mL serum specimens
- Two shipments per year

Program Information

- Three 0.5-mL serum specimens
- Two shipments per year

Program Information

- Two 1.0-mL serum specimens
- Two shipments per year

- Two 1.0-mL serum specimens
- Two shipments per year

Antiphospholipid Antibody ACL			
Analyte	Program Code	Challenges/Shipment	
	ACL		
Anticardiolipin antibody (polyclonal, lgG, lgM, and lgA)	ı	3	
Beta-2-glycoprotein I (polyclonal, lgG, lgM, and lgA)	ı	3	

- Three 0.5-mL lyophilized serum specimens
- Two shipments per year

Antiphosphatidylserine Antibody APS			
Analyte	Program Code	Challenges/Shipment	
	APS		
Anticardiolipin antibody (polyclonal, IgG, IgM, and IgA)	ı	3	
Antiphosphatidylserine antibody (IgG, IgM, and IgA)	ı	3	
Beta-2-glycoprotein I (polyclonal, IgG, IgM, and IgA)	ı	3	

Program Information

- Three 0.5-mL lyophilized serum specimens
- Two shipments per year

Antiribosomal P Antibody ARP					
Analyte Program Code Challenges/Shipmen					
ARP					
Antiribosomal Pantibody	I	3			

Program Information

- Three 0.5-mL serum specimens
- Two shipments per year

Anti-Saccharomyces cerevisiae Antibody ASC			
Analyte Program Code Challenges/Shipme			
	ASC		
Anti-Saccharomyces cerevisiae antibody (IgG and IgA)	ı	2	

- Two 1.0-mL serum specimens
- Two shipments per year

Celiac Serology CES, CESX				
Analyte	Program Code		Challenges/ Shipment	
	CES	CESX		
Antiendomysial antibody (IgA and IgG)	ı		3	
Antiendomysial antibody screen (IgA and IgG)	ı		3	
Antigliadin antibody (IgA and IgG)	ı		3	
Antideamidated gliadin peptide (DGP) antibody (IgA and IgG)			3	
Anti-DGP antibody screen (IgA and IgG)	ı		3	
Antitissue transglutaminase (tTG) antibody (IgA and IgG)	ı		3	
Anti-DGP and anti-tTG antibody screen (IgA and IgG)			3	

- CES Three 0.3-mL serum specimens
- CESX All Survey CES specimens in triplicate
- Two shipments per year

Cyclic Citrullinated Peptide Antibody (Anti-CCP) CCP					
Analyte Program Code Challenges/Shipment					
CCP					
Anti-CCP		2			

Cytokines CTKN

Program Information

- Two 1.0-mL serum specimens
- Two shipments per year



Analyte Program Code Challenges/Shipment Ly CTKN - Ti

3 Interferon (IFN)-gamma Interleukin (IL)-1 beta 3 IL-2 3 IL-6 3 IL-8 3 IL-10 3 П 3 Tumor necrosis factor (TNF)-alpha Vascular endothelial growth factor (VEGF) 3

- Nine 2.0- to 3.0-mL lyophilized serum specimens
- Two shipments per year

Diagnostic Allergy SE				
Analyte/Procedure Program Code Challenges/Ship				
	SE			
IgE, multiallergen screen, qualitative	•	5		
IgE, total		5		
Specific allergens	•	25		
Educational challenges		2 per year		

Diagnostic Allergy SE				
Analyte/Procedure Program Code Challenges/				
	SE			
IgE, multiallergen screen, qualitative		5		
IgE, total	ı	5		
Specific allergens	•	25		
Educational challenges	•	2 per year		

allergens

Program Information • Five 2.0-mL serum specimens

• Three 0.5-mL liquid serum specimens

• Includes common allergens from North America as well as less frequently tested

• Three shipments per year

• Two shipments per year

High-Sensitivity C-Reactive Protein HSCRP					
Analyte	nalyte Program Code Challenges/Shipment				
	HSCRP				
High-sensitivity C-reactive protein	3				

Liver-Kidney Microsomal Antibody (Anti-LKM) LKM				
Analyte	te Program Code Challenges/Shipment			
	LKM			
Anti-LKM		2		

M. tuberculosis-Stimulated Infection Detection QF					
Analyte Program Code Challenges/Shipment					
	QF				
M. tuberculosis	•	2			

• Two shipments per year

Program Information • Two 1.0-mL serum specimens

Program Information

- Two 1.0-mL lyophilized specimens and one lyophilized mitogen control
- · For use with the QuantiFERON®-TB Gold and Gold Plus methods only
- Two shipments per year

Rheumatic Disease Special Serologies RDS				
Analyte Program Code Challenges/Shipment				
	RDS			
Anti-Jo-1 (antihistidyl t-RNA synthetase)		1		
Anti-Scl-70 (anti-DNA topoisomerase)		1		

- Two 1.0-mL serum specimens
- Two shipments per year



Syphilis Serology G			
Analyte	Program Code	Challenges/Shipment	
	G		
Syphilis		5	

Use with VDRL, RPR, MHA-TP/TP-PA/PK-TP/TPHA, EIA, CMIA, multiplex flow immunoassay, TP-LIA IgG, FTA-ABS, and USR methods. Laboratories performing syphilis serology on CSF specimens may also use this Survey.

Program Information

- Five 1.5-mL serum specimens
- Three shipments per year



Total Hemolytic Complement CH50					
Analyte Program Code Challenges/Shipm					
	CH50				
Total hemolytic complement, 50% lysis	1	2			
Total hemolytic complement, 100% lysis ■ 2					

Program Information

- Two 0.5-mL lyophilized serum specimens
- Two shipments per year

Viscosity V			
Analyte	Program Code	Challenges/Shipment	
	V		
Viscosity	•	2	

Program Information

- Two 10.0-mL serum specimens
- Two shipments per year

Serum Free Light Chains SFLC					
Analyte Program Code Challenges/Shipm					
	SFLC				
Kappa serum free light chain		3			
Lambda serum free light chain		3			
Kappa/lambda serum free light chain ratio and ratio interpretation					

- Three 1.0-mL serum specimens
- Two shipments per year

Flow Cytometry

Analytes/procedures in bold type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Flow Cytometry FL, FL1, FL2				
Procedure	Program Code Challenges/Shipment			
	FL	FL1	FL2	
DNA content and cell cycle analysis	•		ı	3
Lymphocyte immunophenotyping	ı			3

These Surveys are not appropriate for hematology analyzers with monoclonal antibody analysis.

Program Information

- FL1 Three 1.5-mL whole blood specimens
- FL2 Three 1.1-mL specimens; two fixed cell line specimens and one calibrator for DNA content and cell cycle analysis
- FL All Survey FL1 and FL2 specimens
- Three shipments per year

Flow Cytometry—Immunophenotypic Characterization of Leukemia/Lymphoma FL3					
Procedure Program Code Challenges/Shipment					
FL3					
Leukemia/lymphoma	I	2			

Program Information

- Two 2.5-mL whole blood specimens and/or cell lines simulating leukemia/ lymphoma; images of tissue sections, bone marrow, and/ or peripheral blood smears with clinical histories
- Online whole slide images powered by DigitalScope® technology
- Two shipments per year

Flow Cytometry, CD34+ FL4		
Analyte	Program Code	Challenges/Shipment
	FL4	
CD34+		2

- Two 1.5-mL stabilized human CD34+ specimens
- · Two shipments per year

Flow Cytometry, Interpretation Only FL5				
Procedure Program Code Challenges/Shipm				
	FL5			
Flow cytometry, interpretation only of leukemia/lymphoma	ı	3		

Survey FL5 is for laboratories that receive flow cytometry analyses from referring laboratories to perform the interpretation of patient results.

Program Information

- Three cases consisting of gated dot plots, clinical histories, and pertinent laboratory data, as well as images of tissue sections, bone marrow, and/or peripheral blood smears
- Online whole slide images powered by DigitalScope technology
- Two shipments per year

Flow Cytometry—B-ALL Minimal Residual Disease BALL			
Analyte	Program Code	Challenges/Shipment	
	BALL		
B-ALL minimal residual disease		3	

Survey BALL is intended for laboratories that currently or will begin to perform minimal residual disease testing (rare event analysis) for B lymphoblastic leukemia/lymphoma. The cases presented will be based on Children's Oncology Group (COG) approved B-ALL MRD method.

Minimum Requirements

For ungated list mode files, each challenge will include 2-3 "virtual tubes"
performed by a 6-color method. The participant will download the files from a CAP
website and analyze the data on a MAC or PC using standard software, including
FlowJo, FACSDiva, Kaluza, Woodlist, etc. The files will be large as each tube will
have collected hundreds of thousands of events. Boolean gating will be necessary
to see if there is an atypical population.

- Two cases consisting of gated dot plots, clinical histories, pertinent laboratory data, and digital images (included for pertinent cases)
- One case with clinical history, pertinent laboratory data, digital images, and ungated list mode files; allows users to examine gating strategies and interpret antibody staining patterns; files are in standard format (see Minimum Requirements)
- Online whole slide images powered by Digital Scope technology, if applicable
- Two shipments per year

Flow Cytometry—Plasma Cell Neoplasms PCNE0			
Analyte	Program Code	Challenges/Shipment	
	PCNEO		
Plasma cell neoplasms	ı	3	

Survey PCNEO is especially helpful for laboratories that have leukemia/lymphoma assays that target plasma cell neoplasms, including cytoplasmic light chain staining.

Program Information

- One 2.5-mL whole blood specimen and/or cell line simulating a plasma cell neoplasm with clinical history and pertinent laboratory data
- Two cases consisting of gated dot plots, clinical histories, and pertinent laboratory data
- Each challenge includes images of tissue sections, bone marrow, and/or peripheral blood smears
- Online whole slide images powered by DigitalScope technology
- Two shipments per year

Flow Cytometry—Immunophenotypic Characterization of Paroxysmal Nocturnal Hemoglobinuria PNH

Analyte	Program Code	Challenges/Shipment
	PNH	
PNH RBC analysis	I	2
PNH WBC analysis	I	2

Additional Information

- The PNH Survey complies with the recommendations from the Guidelines for the Diagnosis and Monitoring of Paroxysmal Nocturnal Hemoglobinuria and Related Disorders by Flow Cytometry for RBC and WBC analysis. Due to the unique nature of these human, donor-based materials, the shipping dates are subject to change. If this should occur, the CAP will provide notification prior to the originally scheduled shipping date.
- This Survey is appropriate for high-sensitivity testing (≤0.01% PNH type clone in red cells and/or granulocytes).

- Two 0.5-mL whole blood specimens for RBC and WBC analysis
- · Two shipments per year

Fetal Red Cell Detection HBF			
Procedure	Program Code	Challenges/Shipment	
	HBF		
Kleihauer-Betke and flow cytometry	I	2	
Rosette fetal screen	I	2	
Acid elution whole slide image	ı	1	

- Two 1.2-mL liquid whole blood specimens
- Not designed for F cell quantitation
- Two online whole slide images per year with optional grids for cell counting
- Powered by DigitalScope® technology
- Two shipments per year

Rare Flow Antigen Validation RFAV1, RFAV2			
Analyte	Program Code Challenges/Shipment		
	RFAV1	RFAV2	
CD1a	•		1
CD103		ı	1

Surveys RFAV1 and RFAV2 do not meet the regulatory requirements for proficiency testing.

Additional Information

These Surveys meet the CAP Accreditation Checklist item FLO.23737, which requires semiannual testing of antigens.

ZAP-70/CD49d Analysis by Flow Cytometry ZAP70			
Analyte	Program Code	Challenges/Shipment	
	ZAP70		
Zeta chain-associated protein kinase 70	ı	3	
CD49d		3	

Additional Information

- This Survey tests for intracellular ZAP-70 staining of a cell line. It allows for assessment of the laboratory's staining techniques and the antibody clone used for ZAP-70 detection.
- CD49d is an important prognostic marker for CLL by flow cytometry. This Survey allows assessment of the laboratory's ability to detect CD49d.

Program Information

- RFAV1 One 4.5-mL cell line specimen
- RFAV2 One 1.0-mL stabilized cell specimen
- Two shipments per year

- Three 4.5-mL cell line specimens
- Two shipments per year

Transfusion Medicine, Viral Markers, and Parentage Testing



Utilize the CAP's Participant Summary Reports to take your laboratory to the next level.

- Compare your results and methods against large peer groups for greater diagnostic confidence.
- Review the extensive discussions to further educate staff on testing trends and best practices.
- Earn continuing education (CE) credit with content that aligns with the proficiency testing challenge.

Transfusion Medicine, Viral Markers, and Parentage Testing

Transfusion Medicine	214
Viral Markers	
Parentage Testing	
New Programs NEW	

17

Transfusion Medicine

Analytes/procedures in **bold** type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Transfusion Medicine J, J1				
Procedure	Progra	Program Code Challenges/Shipment		
	J	J1		
ABO grouping			5	
Rh typing			5	
Antibody detection			5	
Antibody identification			5	
Compatibility testing			5	
Red blood cell antigen typing			1	

Program Information

- J Five 2.0-mL 3%-4% red blood cell suspensions; five 3.0-mL corresponding serum specimens; one 2.0-mL donor red blood cell suspension
- J1 Five 2.0-mL 3%-4% red blood cell suspensions; five 3.0-mL corresponding serum specimens
- · Three shipments per year



Transfusion Medicine—Educational Challenge JE1		
Procedure	Program Code	Challenges/Shipment
	JE1	
Educational challenge	ı	1

- One educational challenge, which may consist of a paper challenge and/or wet specimen for ABO grouping, Rh typing, antibody detection, antibody identification, compatibility testing, antigen typing, and/or direct antiglobulin testing
- Must order in conjunction with Survey J
- Three shipments per year



Electronic Crossmatch EXM				
Procedure	Challenges/Shipment			
	EXM			
Electronic crossmatch	ı	3		

Survey EXM assists laboratories in monitoring the performance of their electronic crossmatching system.

Program Information

- Three simulated, ISBT-128 labeled donor unit challenges and three corresponding red blood cell suspensions
- Must order in conjunction with Survey J
- Two shipments per year



Transfusion Medicine—Automated JAT					
Procedure Program Code Challenges/Shipment					
	JAT				
ABO grouping	I	5			
Antibody detection	I	5			
Antibody identification	■ 5				
Compatibility testing	I	5			
Rh typing	■ 5				

Program Information

- Five bar-coded 4.0-mL 18%-20% whole blood specimens and one 4.0-mL 18%-20% whole blood specimen for compatibility testing
- Three shipments per year



Transfusion Medicine—Automated Education Challenge JATE1						
Procedure Program Code Challenges/Shipment						
JATE1						
Eduational challenge 1						

- One educational challenge, which may consist of a paper challenge and/or wet specimen for ABO grouping, Rh typing, antibody detection, antibody identification, and/or compatibility testing
- Must order in conjunction with Survey JAT
- Three shipments per year



Electronic Crossmatch, Automated EXM2							
Procedure Program Code Challenges/Shipment							
	EXM2						
Electronic crossmatch	ronic crossmatch						

Survey EXM2 assists laboratories in monitoring the performance of their electronic crossmatching system.

Program Information

- Three simulated, ISBT-128 labeled donor unit challenges and three corresponding red blood cell suspensions
- Must order in conjunction with Survey JAT
- Two shipments per year



ABO Subgroup Typing ABOSG					
Procedure	Program Code Challenges/Shipment				
	ABOSG				
ABO subgroup typing	I	3			
Rh typing		3			

ABO Subgroup Typing ABOSG			
Procedure	Program Code	Challenges/Shipment	
	ABOSG		
ABO subgroup typing	I	3	
Rh typing	ı	3	
Titl Cyping	<u>-</u>		

Red Blood Cell Antigen Genotyping RAG				
Procedure Program Code Challenges/Shipme				
	RAG			
Red blood cell antigen genotype with predictive phenotype	ı	3		

Program Information

- Three 2.0-mL 3%-4% red blood cell suspensions; three 2.0-mL corresponding serum specimens
- Two shipments per year

- Three 2.0-mL whole blood specimens
- Two shipments per year

Red Blood Cell Antigen Typing RBCAT					
Procedure Program Code Challenges/Shipment					
	RBCAT				
Red blood cell antigen typing		2			

Survey RBCAT is for donor centers and transfusion laboratories performing non-automated/manual red cell phenotyping for the management of patients with complex serology (ie, alloimmunization, sickle cell disease, warm autoimmune hemolytic anemia). Challenges will include antigens such as Rh, Kell, MNSs, Duffy, and Kidd blood group system.

Antibody Titer ABT, ABT1, ABT2, ABT3					
Procedure	Program Code Challenges/Shipment				
	ABT	ABT1	ABT2	ABT3	
Anti-A titer	•	ı			1
Anti-B titer					1
Anti-D titer					1

Program Information

- Two 2.0-mL 2%-4% red blood cell suspensions
- · Two shipments per year

- ABT One 2.0-mL plasma specimen for anti-A titer with one corresponding titer cell (3%-4% red blood cell suspension); one 2.0-mL plasma specimen for anti-D titer with one corresponding titer cell (3%-4% red blood cell suspension)
- ABT1- One 2.0-mL plasma specimen for anti-A titer with one corresponding titer cell (3%–4% red blood cell suspension)
- ABT2 One 2.0-mL plasma specimen for anti-D titer with one corresponding titer cell (3%–4% red blood cell suspension)
- ABT3 One 2.0-mL plasma specimen for anti-B titer with one corresponding titer cell (3%–4% red blood cell suspension)
- · Two shipments per year

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Transfusion-Related Cell Count TRC					
Procedure Program Code Challenges/Shipment					
	TRC				
Platelet count (platelet-rich plasma)	ı	5			
WBC count	I	4			
Dry challenge	ı	2			

WBC counts must be performed using a Nageotte chamber, fluorescence microscopy, or by flow cytometry.

Program Information

- Five 1.2-mL suspensions of platelet-rich plasma
- Two 1.0-mL vials leukocytereduced platelet material
- Two 1.0-mL vials leukocytereduced red blood cells
- Three shipments per year

Direct Antiglobulin Testing DAT			
Procedure	Program Code Challenges/Shipment		
	DAT		
Direct antiglobulin testing	ı	3	

Program Information

- Three 2.0-mL 3%-4% red blood cell suspensions
- For use with manual method
- Two shipments per year

Eluate Survey ELU			
Procedure	Program Code Challenges/Shipme		
	ELU		
Antibody elution		2	

Program Information

- Two 2.0-mL 50% red blood cell suspensions
- Two shipments per year

Fetal Red Cell Detection HBF			
Procedure Program Code Challenges/Shipm			
	HBF		
Kleihauer-Betke and flow cytometry	I	2	
Rosette fetal screen	I	2	
Acid elution whole slide image	ı	1	

- Two 1.2-mL liquid whole blood specimens
- · Not designed for F cell quantitation
- Two online whole slide images per year with optional grids for cell counting
- Powered by DigitalScope® technology
- Two shipments per year

Platelet Serology PS			
Procedure Program Code Challenges/Shipmen			
	PS		
Antibody detection	I	3	
Platelet crossmatch	ı	3	
Platelet antibody identification	ı	3	

A low concentration of sodium azide may be present in the specimens and may affect lymphocytotoxicity methods.

Program Information

- Three 3.0-mL plasma specimens
- For use with solid-phase red cell adherence, flow cytometry, and EIA/ELISA methods
- Two shipments per year

Transfusion Medicine Comprehensive—Competency Assessment TMCA Procedure Program Code Challenges/Shipment TMCA ABO grouping 2

Procedure Program Code Challenges/Shipment TMCA TMCA ABO grouping ■ 2 Antibody detection ■ 2 Antibody identification ■ 2 Compatibility testing ■ 2 Rh typing ■ 2

Survey TMCA does not meet the regulatory requirements for proficiency testing.

Program Information

- Two 2.0-mL 3%-4% red blood cell suspensions
- Two 3.0-mL corresponding serum specimens
- One 2.0-mL donor 3% red blood cell suspension
- Three shipments per year; order shipments individually or for an entire year

Direct Antiglobulin Test—Competency Assessment TMCAD				
Procedure Program Code Challenges/Shipment				
TMCAD				
Direct antiglobulin testing 2				

Survey TMCAD does not meet the regulatory requirements for proficiency testing.

- Two 2.0-mL 3%-4% red blood cell suspensions
- Two shipments per year; order shipments individually or for an entire year

Eluate Competency Assessment TMCAE				
Procedure	Procedure Program Code Challenges/Shipme			
	TMCAE			
Antibody elution	ı	2		

Survey TMCAE does not meet the regulatory requirements for proficiency testing.

Program Information

- Two 2.0-mL 50% red blood cell suspensions
- Two shipments per year; order shipments individually or for an entire year

Fetal Red Cell Quantitation—Competency Assessment TMCAF					
Procedure	ocedure Program Code Challenges/Shipment				
	TMCAF				
Kleihauer-Betke, flow cytometry	I	2			
Rosette fetal screen	I	2			
Acid elution whole slide image	I	1			

Survey TMCAF does not meet the regulatory requirements for proficiency testing.

Program Information

- Two 1.2-mL whole blood specimens
- Two online whole slide images per year with optional grids for cell counting
- Powered by DigitalScope technology
- Two shipments per year; order shipments individually or for an entire year

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Cord Blood and Stem Cell Processing CBT, SCP			
Procedure	Progra	m Code	Challenges/Shipment
	СВТ	SCP	
Absolute CD3		I	2
Absolute CD34			2
Absolute CD45			2
Bacterial culture		I	2
%CD3+			2
%CD34+		•	2
%CD45+		ı	2
BFU-E		ı	2
CFU-E		ı	2
CFU-GEMM		ı	2
CFU-GM			2
Total CFC		ı	2
Fungal culture		ı	2
Hematocrit		ı	2
Hemoglobin		ı	2
Mononuclear cell count		ı	2
Nucleated red cells		ı	2
Number of CD34 positive events		ı	2
Number of CD45 positive events	1		2
Total nucleated cells		ı	2
Viability		ı	2
WBC count		ı	2

- · Because these materials are human donor-based, the ship date is subject to change. If this should occur, notification will be provided prior to the scheduled date. In some instances, the program may ship in two installments.
- Due to material stability, no replacements will be available.
- See International Shipping information section in the Ordering Information Supplement regarding additional dangerous goods shipping fees.

Program Information

- CBT Two 2.5-mL cord blood specimens; designed for assays required for the production of umbilical cord blood stem cell programs
- SCP Two 3.0-mL peripheral blood specimens; designed for laboratories that process and assess the suitability of stem cells
- · Two shipments per year



Refer to the Ordering Information provided for information regarding additional dangerous goods and related fees.

Bacterial Detection in Platelets BDP, BDP5			
Procedure	Program Code Challenges/Shipment		
	BDP	BDP5	
Bacterial culture and detection systems			2
Bacterial culture and detection systems		ı	5

- The Centers for Medicare & Medicaid Services (CMS) requires proficiency testing for bacterial detection/identification. Please select the appropriate program for your laboratory based on the information below.
- Survey BDP is designed for donor centers/laboratories that are associated with a CMS-certified microbiology laboratory with the same CLIA number and are participating in an approved proficiency testing program for bacterial detection.
- Survey BDP5 is designed for donor centers/laboratories that are performing bacterial detection for the purposes of platelet unit screening and are not associated with a CMS-certified microbiology laboratory with the same CLIA number.
- See International Shipping information section in the Ordering Information Supplement regarding additional dangerous goods shipping fees.

Program Information

- BDP Two lyophilized pellet specimens with diluents; two shipments per year
- BDP5 Five lyophilized pellet specimens with diluents; three shipments per year



Bacterial Detection in Platelets, Rapid BDPV, BDPV5				
Procedure	dure Challenges/Shipment			
Program Code				
	BDPV BDPV5			
CMS certified Rapid immunoassay	2	5		

Additional Information

- The Centers for Medicare & Medicaid Services (CMS) requires proficiency testing for bacterial detection in platelets.
- Survey BDPV is designed for donor centers/laboratories that are associated with a CMS-certified microbiology laboratory with the same CLIA number and are participating in an approved proficiency testing program for bacterial detection.
- Survey BDPV5 is designed for donor centers/laboratories that are performing bacterial detection for the purposes of platelet unit screening and are not associated with a CMS-certified microbiology laboratory with the same CLIA number.
- See International Shipping information section in the Ordering Information Supplement regarding additional dangerous goods shipping fees.

Program Information

- BDPV Two frozen specimens; two shipments per year
- BDPV5 Five frozen specimens; three shipments per year
- For use with methods such as Verax Biomedical





Refer to the Ordering Information provided for information regarding additional dangerous goods and related fees.

Expanded Transfusion Medicine Exercises ETME1				
Procedure Program Code Challenges/Shipment				
ETME1				
Expanded challenges		2		

Survey ETME1 is an educational opportunity that offers:

- More challenging and/or complex antibody identification
- · Comprehensive case studies in transfusion medicine
- Simulated collaboration with other professionals, including those within or outside your institution
- A method for determining the laboratory's ability to recognize and integrate problem solving skills in transfusion medicine

The wet challenge may consist of specimens for ABO grouping, Rh typing, antibody detection, antibody identification, compatibility testing, antigen typing, direct antiglobulin testing, antibody titer, and/or antibody elution.

Program Information

- ETME1 One paper challenge and one wet challenge consisting of a serum specimen(s) and/or red blood cell suspensions
- · Two shipments per year

Make critical transfusion decisions with confidence

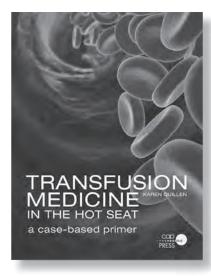
Transfusion Medicine in the Hot Seat is a valuable educational resource for pathology trainees and pathologists practicing transfusion medicine. The text presents a total of 26 realistic transfusion scenarios divided into three sections:

- Antibodies
- Blood Components
- Complications

The short-case format makes the information easily accessible and can serve as the basis for a transfusion medicine curriculum in clinical pathology.

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Item number: PUB224 Softcover; 123 pages

Viral Markers

Analytes/procedures in **bold** type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Viral Markers—Series 1 VM1			
Analyte	Program Code	Challenges/Shipment	
	VM1		
Anti-HAV (total: IgM and IgG)	1	5	
Anti-HAV (IgG)	1	5	
Anti-HBc (total: IgM and IgG)	1	5	
Anti-HBs	1	5	
Anti-HBs, quantitative	1	5	
Anti-HCV		5	
Anti-HIV-1		5	
Anti-HIV-1/2	1	5	
Anti-HIV-2		5	
HBsAg	ı	5	

Do not use Survey VM1 with rapid anti-HCV, anti-HIV-1, or anti-HIV-1/2 kits. See page 225 for Surveys appropriate for rapid methods.

Viral Markers—Series 2 VM2				
Analyte Program Code Challenges/Shipme				
	VM2			
Anti-HBe		5		
HBeAg	■ 5			

Viral Markers—Series 3 VM3				
Analyte	Program Code Ch			
	VM3	_		
Anti-CMV	ı	3		
Anti-HTLV-I/II		3		
HIV-1 p24 antigen		3		

Viral Markers—Series 4 VM4						
Analyte Program Code Challenges/Shipment						
	VM4					
Anti-Trypanosoma cruzi (Chagas disease)	se) 1 2					

Program Information

- Five 3.5-mL plasma specimens
- Three shipments per year

Program Information

- Five 3.5-mL plasma specimens
- Three shipments per year

Program Information

- Three 3.5-mL plasma specimens
- Two shipments per year

- Two 1.0-mL plasma specimens
- Two shipments per year

Viral Markers—Series 5 VM5						
Analyte Program Code Challenges/Shipment						
	VM5					
Anti-HAV (IgM)	■ 5					
Anti-HBc (IgM)	/ I) ■ 5					

Viral Markers—Series 6 VM6					
Analyte Program Code Challenges/Shipmen					
	VM6				
Anti-HIV-1/2	I	5			
HIV-1 p24 antigen	I 5				

Anti-HIV 1/2 AHIV, AHIVW				
Analyte/Procedure	Program Code		Challenges/Shipment	
	AHIV	AHIVW		
Anti-HIV-1, Anti-HIV-2, Anti-HIV-1/2			5	
Anti-HIV-1, Anti-HIV-1/2, waived methods only			2	

Anti-HCV, Rapid Methods, Waived RHCVW						
Analyte/Procedure	Challenges/Shipment					
	RHCVW					
Anti-HCV, waived methods only	ı	3				

Program Information

- Five 1.5-mL plasma specimens
- Three shipments per year

Program Information

- Five 0.5-mL plasma specimens
- For use with methods such as the Abbott ARCHITECT HIV Combo, Bio-Rad GS HIV Combo, and Alere Determine HIV Combo assays
- Three shipments per year

Program Information

- AHIV Five 0.5-mL plasma specimens; three shipments per year
- AHIVW Two 0.5-mL plasma specimens; two shipments per year

- Three 0.5-mL plasma specimens
- Two shipments per year

Nucleic Acid Testing NAT				
Analyte Program Code Challenges/Shipment				
	NAT			
HBV	I	5		
HCV ■		5		
HIV	5			
West Nile virus ■ 5				

Program Information

- Five 6.0-mL plasma specimens
- Designed for blood donor centers performing nucleic acid testing on donor units
- Compatible with HIV, HCV, and HBV multiplex assays
- Three shipments per year

Vector-Borne Disease—Molecular VBDM					
Analyte	Program Code Challenges/Shipment				
	VBDM				
Zika virus	us I 3				

Program Information

- Three 1.5-mL liquid specimens
- Two shipments per year

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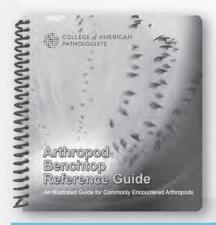
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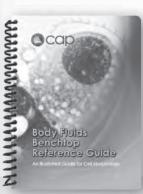
Parentage Testing

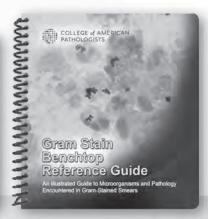
Parentage/Relationship Test—Filter Paper PARF				
Analyte/Procedure Program Code Challenges/Shipr				
	PARF			
Calculation challenge (paper challenge)		1		
DNA testing (PCR)		4		

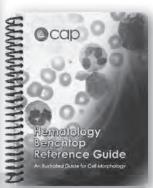
- Three blood-stained filter paper paternity trio specimens; two buccal swabs for a second allegedfather challenge
- Reporting for short tandem repeats (STRs), Y-STRs, as well as the conclusions provided
- Three shipments per year

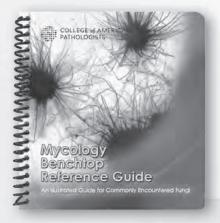
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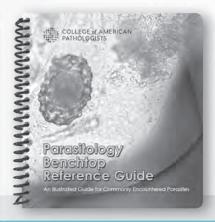


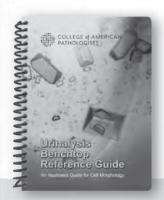












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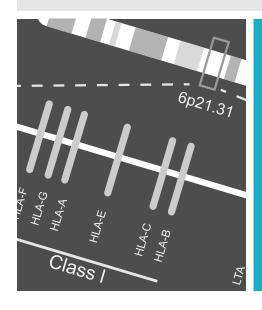
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18 Histocompatibility



CAP laboratory accreditation checklists are updated annually to keep you current with changes in laboratory medicine.

Updates in the 2017 Histocompatibility Checklist requirements help you comply with:

- United Network for Organ Sharing (UNOS) bylaws
- Foundation for the Accreditation of Cellular Therapy (FACT) standards
- CLIA regulations

New Analyte/Drug Additions NEW

DQA1*02

DQA1*03

DQA1*05

DQB1*02:01

DQB1*02:02

Discontinued Programs

HLA Molecular Typing (ML, DL) HLA Serologic Typing (Class I/II) (ABO, HLAS, HLAS1)

Histocompatibility

Analytes/procedures in **bold** type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

HLA Crossmatching, Antibody Screen, and Antibody Identification (Class I) MX1B, MX1C, MX1E

Procedure	Program Code			Challenges/Shipment
	MX1B	MX1C	MX1E	
Crossmatching				8
Antibody screen				4
Antibody identification				4

Additional Information

Blood donor screening is now a reporting option for antibody screening results. This change covers the use of HLA testing in blood centers/hospital laboratories for the purpose of donor qualification.

Program Information

- MX1B Four 0.25-mL plasma specimens; two (approximately 1.0 x 10⁶ cells) purified peripheral blood lymphocyte specimens
- MX1C Four 0.50-mL plasma specimens; two (approximately 4.0 x 10⁶ cells) purified peripheral blood lymphocyte specimens
- MX1E Four 0.30-mL plasma specimens; must be ordered in conjunction with Survey MX1B or MX1C
- Three shipments per year

HLA Crossmatching, Antibody Screen, and Antibody Identification (Class II) MX2B, MX2C, MX2E

Procedure	Pro	gram Co	de	Challenges/Shipment
	MX2B MX2C MX2E			
Crossmatching		ı		4
Antibody screen		ı		2
Antibody identification		ı		2

Additional Information

Blood donor screening is now a reporting option for antibody screening results. This change covers the use of HLA testing in blood centers/hospital laboratories for the purpose of donor qualification.

Program Information

- MX2B Two 0.25-mL plasma specimens; two (approximately 7.2 x 10⁶ cells) purified peripheral blood lymphocyte specimens
- MX2C Two 0.50-mL plasma specimens; two (approximately 9.6 x 10⁶ cells) purified peripheral blood lymphocyte specimens
- MX2E Two 0.30-mL plasma specimens; must be ordered in conjunction with Survey MX2B or MX2C
- Three shipments per year

For laboratories conducting BOTH Class I and Class II HLA testing, see next page.

HLA Crossmatching, Antibody Screen, and Antibody Identification (Class I/II) Combinations MXB, MXC

Procedure	Corresponding Survey	Progra	am Code
		MXB	MXC
Crossmatching, antibody screen, and antibody identification (Class I)	MX1B*	•	
Crossmatching, antibody screen, and antibody identification (Class II)	MX2B*		
Crossmatching, antibody screen, and antibody identification (Class I)	MX1C*		ı
Crossmatching, antibody screen, and antibody identification (Class II)	MX2C*		ı

^{*}See page 230 for specimen and analyte information.

Program Information

- MXB Class I: four 0.25-mL plasma specimens, two purified peripheral blood lymphocyte specimens; Class II: two 0.25-mL plasma specimens, two purified peripheral blood lymphocyte specimens
- MXC Class I: four 0.50-mL plasma specimens, two purified peripheral blood lymphocyte specimens; Class II: two 0.50-mL plasma specimens, two purified peripheral blood lymphocyte specimens
- Three shipments per year

Class I & II HLA Molecular Typing DML					
Procedure	Program Code	Challenges/Shipment			
	DML				
Molecular HLA-A, -B, and -C typing (Class I)	I	5			
Molecular HLA-DR, -DQ, and -DP typing (Class II)	ı	5			

HLA-B27 Typing B27				
Procedure	Program Code	Challenges/Shipment		
	B27			
HLA-B27 typing		5		

Program Information

- Ten approximately 1.0-mL whole blood specimens in CPD-A
- Serologic equivalents and MICA reporting available
- Three shipments per year

- Five 2.0-mL whole blood specimens in CPD-A
- Two shipments per year

Antibody Titer ABT, ABT1, ABT2, ABT3						
Procedure		Program Code Challenges/Shipment				
	ABT	ABT1	ABT2	ABT3		
Anti-A titer					1	
Anti-B titer					1	
Anti-D titer	ı				1	

Program Information

- ABT One 2.0-mL plasma specimen for anti-A titer with one corresponding titer cell (3%-4% red blood cell suspension); one 2.0-mL plasma specimen for anti-D titer with one corresponding titer cell (3%-4% red blood cell suspension)
- ABT1- One 2.0-mL plasma specimen for anti-A titer with one corresponding titer cell (3%–4% red blood cell suspension)
- ABT2 One 2.0-mL plasma specimen for anti-D titer with one corresponding titer cell (3%–4% red blood cell suspension)
- ABT3 One 2.0-mL plasma specimen for anti-B titer with one corresponding titer cell (3%-4% red blood cell suspension)
- Two shipments per year

Monitoring Engraftment ME						
Procedure Program Code Challenges/Shipmen						
ME						
Stem cell monitoring engraftment	I	3				

- Five 1.0-mL whole blood specimens
- Designed for laboratories supporting stem cell transplant and laboratories monitoring chimerism after organ transplantation
- Three shipments per year

HLA Disease Ass	sociation-D	rug Risk	DADR1, DADR2
Analyte	Progra	m Code	Challenges/Shipment
	DADR1	DADR2	
HLA-A*31:01			3
HLA-B*13:01			3
HLA-B*15:02			3
HLA-B*57:01			3
HLA-B*58:01			3
HLA-A*29:01		ı	3
HLA-A*29:02			3
HLA-DQA1*04:01			3
HLA-DQA1*05:01		ı	3
HLA-DQB1*03:02			3
HLA-DQB1*06:02			3
HLA-DRB1*03:01		•	3
HLA-DRB1*03:02			3
HLA-DRB1*04:02			3
HLA-DRB1*04:03		•	3
HLA-DRB1*04:06			3
HLA-DRB1*08:02			3
HLA-DRB1*08:04			3
HLA-DRB1*14:04			3
HLA-DRB1*14:05			3
HLA-DRB1*14:08			3
HLA-DRB1*15:01		•	3
HLA-DRB1*15:02		•	3
DQA1*02 NEW			3
DQA1*03 NEW		•	3
DQA1*05 NEW		•	3
DQB1*02:01 NEW			3
DQB1*02:02 NEW			3

These Surveys will challenge the laboratory to accurately identify the presence or absence of alleles associated with a variety of disease states (listed below) and/or the adverse reactions to specific drugs.

DADR1

- o Carbamezepine induced Stevens-Johnson syndrome (CSJ)
- o Allopurinol Stevens-Johnson syndrome (ASJ)
- O Hypersensitivity to abacavir (HA)
- O Dapsone hypersensitivity (DH)

DADR2

- o Celiac disease (CD)
- o Narcolepsy (N)
- o Pemphigus vulgaris (PV)
- o Psoriasis (P)
- o Antiglomerular basement membrane disease (ABM)
- o Birdshot retinochoroidopathy (BR)
- Oldiopathic myopathy (IM)

- Three 0.1-mL specimens, each containing 200 µg/mL of human DNA in media
- Two shipments per year

Atlas of Transplant Pathology

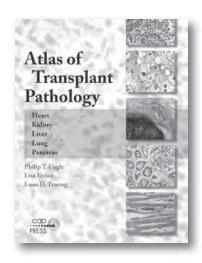
This atlas serves as a handy resource for practical interpretation of solid organ transplant biopsies and other specimens by general pathologists as well as subspecialists.

Includes over 600+ photomicrographs and tables.

Available in print and ebook formats.

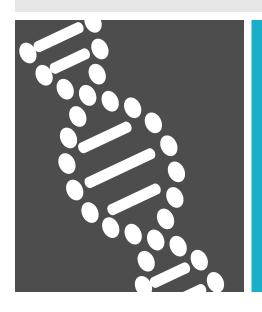
View sample pages and order online:

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- ebooks at ebooks cap org



Item number: PUB124 254 pages; 2015

Genetics and Molecular Pathology



We support quality assurance in NGS.

- Molecular tumor profiling using cell free DNA (CFDNA)
- Noninvasive prenatal screening of aneuploidies (NIPT)
- IGHV somatic hypermutation (IGHV)
- Customized in silico programs: NGS Undiagnosed Disorders-Exome (NGSE) and NGS Bioinformatics Somatic Variant Validated Material (NGSBV)
- Transcriptome analysis using RNASeq (RNA)
- Educational program for germline variant interpretation (VIP/VIP1)

Genetics and Molecular Pathology

Cytogenetics	236
Biochemical and Molecular Genetics	
Next-Generation Sequencing	246
Molecular Oncology—Solid Tumors	
Molecular Oncology—Hematologic	254
New Programs NEW	
Cell Free DNA (CFDNA)	252
IGHV Mutation Analysis (IGHV)	
Next-Generation Sequencing Bioinformatics Somatic Validated Materials (NGSBV)	249
Next-Generation Sequencing Undiagnosed Disorders—Exome (NGSE)	248
Noninvasive Prenatal Testing (NIPT)	245
RNA Sequencing (RNA)	252
Variant Interpretation Only (VIP/VIP1)	244
New Analyte/Drug Additions	
Molecular Hematologic Oncology (MHO2, MHO3)	254

Discontinued Programs

BCR/ABL1 p210

Cytogenetics

Analytes/procedures in bold type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

CAP/ACMG Cytogenetics CY, CYBK					
Analyte/Procedure	Program Code Challenges/Shipment				
	CY	CYBK			
Chromosome abnormality			6		
Karyotype nomenclature			6		
Educational challenge, ungraded			1 per year		

Additional Information

Each challenge includes a case history and images of metaphase cells that are representative of each case. Each mailing will include three constitutional and three neoplastic challenges.

Program Information

- CY Online images of metaphase cells; delivered two times a year; your CAP shipping contact will be notified <u>via email</u> when the activity is available
- CYBK Prints of metaphase cells; two shipments per year



CAP/ACMG Fluorescence In Situ Hybridization CYF, CYI					
Disease/Procedure	Progra	m Code	Challenges	/Shipment	
	CYF	CYI	Α	В	
Constitutional and Hematologic Disorders					
FISH for constitutional disorder - slides	1		1	1	
FISH for constitutional disorder - paper/photograph challenge	ı		2	2	
FISH for hematologic disorder - slides	ı		1	1	
FISH for hematologic disorder - paper/photograph challenge	ı		2	2	
Urothelial Carcinoma					
FISH for urothelial carcinoma			2	2	

Additional Information

CYF 2018-A:

Constitutional disorder - Williams syndrome critical region (two slides)

Constitutional disorder - (two paper/photograph challenges)

Hematologic disorder - BCR/ABL1 (two slides)

Hematologic disorder - (two paper/photograph challenges)

• CYF 2018-B:

Constitutional disorder - SRY (two slides)

Constitutional disorder - (two paper/photograph challenges)

Hematologic disorder - CLL panel (four slides)

Hematologic disorder - (two paper/photograph challenges)

• CYF is prepared from cell suspension samples. For FISH in paraffin-embedded tissues, see page 237.

- CYF Four slides and four paper/photograph challenges
- CYI Two 250-µL cell samples suspended in ethanol from two different specimens; participants use FISH to detect chromosome abnormalities
- Two shipments per year



CAP/ACMG Fluorescence In Situ Hybridization for Paraffin-Embedded Tissue CYH, CYJ, CYK, CYL

Analyte/Procedure	F	Program Code			Challenges/ Shipment	
	СҮН	CYJ	CYK	CYL	Α	В
Breast Cancer						
HER2 gene amplification					10	10
Brain/Glioma Tissue						
1p/19q					1	1
Solid Tumor						
FOXO1 gene rearrangement			•		1	
DDIT3 gene rearrangement						1
Lymphoma Tissue						
CCND1 (CyclinD1) gene rearrangement					1	
BCL2 gene rearrangement						1

- · CYH Two unstained, fivecore tissue microarray slides equivalent to 10 paraffinembedded breast tissue specimens; two H&E stained tissue microarray slides will also be provided
- CYJ Four unstained slides; one H&E stained slide
- CYK, CYL Two unstained slides; one H&E stained slide
- · All CYJ, CYK, CYL specimens will be 4.0-micron tissue sections mounted on positively charged glass slides
- Two shipments per year



CAP/ACMG Constitutional Microarray CYCGH						
Procedure	Program Code	Challenges/Shipment				
	CYCGH					
Cytogenomic microarray analysis for constitutional abnormality	ı	2				
Educational paper/photograph challenge for constitutional abnormality	ı	1				

Participants will identify and characterize gains or losses and the cytogenetic location of abnormalities detected.

This Survey is not appropriate for low resolution arrays that are designed to detect only aneuploidy.

CAP/ACMG Oncology Microarray CYCMA Procedure Program Code Challenges/Shipment CYCMA Cytogenomic microarray analysis for oncologic abnormality Educational paper/photograph challenge for oncologic abnormality 1

Additional Information

Participants will identify and characterize gains or losses and the cytogenetic location of abnormalities detected.

Program Information

- Two 3.0-µg DNA specimens; one paper/photograph challenge
- Two shipments per year



- One 3.0-ug DNA specimen; one paper/photograph challenge
- Two shipments per year



Biochemical and Molecular Genetics

Analytes/procedures in **bold** type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

CAP/ACMG Biochemical Genetics BGL, BGL1					
Analyte/Procedure	Progra	m Code	Challenges/Shipment		
	BGL	BGL1			
Acylcarnitines, qualitative and quantitative	ı		1		
Amino acids, qualitative and quantitative	ı		1		
Carnitine, qualitative and quantitative			3		
Glycosaminoglycans (mucopolysaccharides), qualitative and quantitative	ı		1		
Organic acids, qualitative and quantitative	ı		1		
Educational challenge			1		

Program Information

- BGL -
 - Acylcarnitines: One 0.1-mL plasma specimen
 - Amino acids: One 1.0-mL plasma or 2.0-mL urine specimen
 - Glycosaminoglycans (mucopolysaccharides): One 2.0-mL urine specimen
 - Organic acids: One 7.5-mL urine specimen
 - Educational challenge: Will consist of any one of the BGL analytes
- BGL1 Three 0.3-mL serum specimens
- Two shipments per year



CAP/ACMG Alpha-1 Antitrypsin Genotyping AAT					
Analyte/Procedure	Program Code	Challenges/Shipment			
	AAT				
Alpha-1 antitrypsin (SERPINA1) genotyping	ı	3			

This Survey will test for the M, S, and Z alleles.

CAP/ACMG Apolipoprotein E Genotyping APOE						
Analyte/Procedure	Program Code Challenges/Shipment					
	APOE					
Apolipoprotein E (APOE) genotyping	ı	3				

This Survey is designed for laboratories utilizing APOE testing for hyperlipoproteinemia type III and Alzheimer diseases and will test for APOE e2, APOE e3, and APOE e4.

Program Information

- Three 10.0-µg extracted DNA specimens
- Two shipments per year



- Three 10.0-µg extracted DNA specimens
- Two shipments per year



Σ	
and	
etics	
Gene	

CAP/ACMG BRCA1/2 Sequencing BRCA						
Analyte/Procedure	Program Code	Challenges/Shipment				
	BRCA					
BRCA1/2 DNA sequencing and variant interpretation	ı	3				
BRCA1/2 duplication/deletion analysis	ı	3				

Program Information

- Three 10-µg extracted DNA specimens
- Two shipments per year



Additional Information

- Test your skill at reporting and interpreting DNA sequence variants for BRCA1/2 using standard nomenclature.
- · Receive a summary and discussion of responses, including comments on the variant nomenclature and known or expected outcomes from identified variants.
- Primers are not included; laboratories are expected to utilize the primers used in routine clinical testing.

CAP/ACMG Hemoglobinopathies Genotyping HGM						
Analyte/Procedure Program Code Challenges/Shipment						
	HGM					
Alpha-thalassemia	I	3				
Beta-thalassemia	I	3				
Hemoglobin S/C	ı	3				

Program Information

- Three 50.0-µg extracted **DNA** specimens
- Two shipments per year



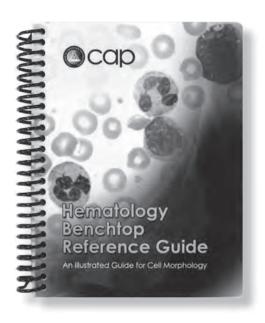
Hematology Benchtop Reference Guide (HBRG)

- More than 50 different cell identifications, including common and rare cells
- Detailed descriptions for each cell morphology
- Six tabbed sections for easy reference
 - Erythrocytes
 - Erythrocyte Inclusions
 - o Granulocytic (Myeloid) and Monocytic Cells
 - Lymphocytic Cells
 - Platelets and Megakaryocytic Cells
 - o Microorganisms and Artifacts
- A durable and water-resistant format to withstand years of benchtop use—5" x 61/2"

Choose code HBRG on your Surveys order form.

Or, view sample pages and order online:

- · printed books at estore.cap.org
- ebooks at ebooks cap org



CAP/ACMG Molecular Genetics MGL1, MGL2, MGL3, MGL4, MGL5

Program Code						
Disease/Gene	Program Code MGL1 MGL2 MGL3 MGL4 MGL5					Challenges/ Shipment
	MGL1	WGL2	MGL3		MGL5	
Bloom syndrome						3
BRCA1/2			•			3
Canavan						3
Connexin 26			•			3
Cystic fibrosis		•				3/2(MGL5)
DMD/Becker						3
Factor V Leiden						3
Familial dysautonomia						3
Fanconi anemia complementation group C						3
Fragile X						3
Friedreich ataxia		ı				3
Gaucher						3
Glycogen storage disease type IA						3
Hemochromatosis						3
Hemoglobin S/C		1				3
Huntington						3
Methylene tetrahydrofolate reductase (MTHFR) c.665C>T (677C>T) and c.1286A>C (1298A>C)	•					3
Mucolipidosis IV						3
Multiple endocrine neoplasia type 2 (MEN2)			•			3
Myotonic dystrophy		ı				3
Niemann-Pick type A/B						3
Plasminogen activator inhibitor (PAI)-1						3
Prader-Willi/Angelman syndrome						3
Prothrombin						3
RhD		ı				3
Spinal muscular atrophy		ı				3
0.1						3
Spinocerebellar ataxia		-				O

Additional Information

- The BRCA1/2 program (module MGL3) is designed for laboratories testing for the three Ashkenazi Jewish founder mutations.
- Module MGL4 is designed for laboratories testing for diseases/disorders related to Ashkenazi Jewish ancestry.
- The Prader-Willi/Angelman syndrome program is designed for laboratories using methylation techniques for analysis.

- MGL1, MGL2, MGL3, MGL4 - Three 50.0-µg extracted DNA specimens
- MGL5 Two 50.0-μg extracted DNA specimens
- Two shipments per year



CAP/ACMG Inherited Metabolic Diseases IMD1, IMD2, IMD3				
Analyte/Procedure	Program Code Challenges/Shipr			
	IMD1	IMD2	IMD3	
Mitochondrial DNA deletion syndromes	•			3
MCAD				3
Mitochondrial cytopathies*			ı	3

^{*}Includes disorders/diseases such as Leber hereditary optic neuropathy and myoclonus epilepsy with ragged red fibers (MERRF).

Program Information

- IMD1 Three 50.0-µL DNA specimens (50.0 ng/ µL DNA PCR product that encompasses the entire mitochondrial genome)
- IMD2, IMD3 Three 50.0-µg extracted DNA specimens
- Two shipments per year



CAP/ACMG Molecular Genetics Sequencing SEC, SEC1						
Procedure	Progra	m Code	Challenges/Shipment			
	SEC	SEC1				
DNA sequencing interpretation challenge	1		3			
DNA sequencing	3					

Additional Information

- Test your skill at interpreting and reporting DNA sequence variants for inherited diseases using standard nomenclature.
- Receive a summary and discussion of responses, including comments on nomenclature, known or expected outcomes from identified variants, and teaching points about genes/disorders represented.
- Results for both programs (SEC, SEC1) must be submitted online through e-LAB Solutions Suite.

- SEC One CD-ROM
 containing DNA sequence
 electropherogram files with a
 range of variants, suitable for
 base-calling and analysis
 using a range of commercial
 or public domain software
 programs; also includes
 nomenclature/variant
 references
- SEC1 Three 10.0-µg extracted DNA specimens; forward and reverse lyophilized primers are provided
- Two shipments per year



Pharmacogenetics PGX, PGX1, PGX2, PGX3					
Analyte/Procedure		Prograi	Challenges/ Shipment		
	PGX	PGX1	PGX2	PGX3	
CYP2C19	•				3
CYP2C9	•				3
CYP2D6					3
CYP3A4	ı				3
CYP3A5					3
SLC01B1 (rs4149056)	1				3
VKORC1	1				3
IL28B (rs12979860)					3
HLA-B*15:02			1		3
HLA-B*57:01			1		3
DPYD				1	3
TPMT				1	3
UGT1A1				ı	3

- UGT1A1 (PGX3 Survey) tests the laboratory's ability to detect variants in the TATA repeat sequence in the UGT1A1 promotor (eg, UGT1A1*28 with seven TA repeats). The ability to detect variants in other regions of the UGT1A1 gene is not part of this program.
- Survey PGX2 is designed for laboratories that provide *HLA-B*57:01* testing to identify risk of hypersensitivity to abacavir and HLA-B*15:02 testing to identify risk of hypersensitivity to carbamazepine. The intended response is qualitative (presence/absence of the allele). This Survey is not appropriate for laboratories that perform molecular HLA typing. For HLA typing proficiency testing, please consult the HLA Molecular Typing (DML) Survey.

CAP/ACMG Rett Syndrome (MECP2) RETT						
Analyte/Procedure	Program Code	Challenges/Shipment				
	RETT					
MECP2 genotyping	I	3				
MECP2 duplication/deletion analysis	I	3				

Program Information

- Three 25.0-µg extracted DNA specimens
- Includes allele detection (genotyping) and/or interpretive challenges
- Two shipments per year

- Three 10.0-µg extracted DNA specimens
- Two shipments per year



CAP/ACMG Thrombophilia Mutations TPM					
Analyte/Procedure Program Code Challenges/Shipmer					
	ТРМ				
Factor II	I	3			
Factor V		3			

This Survey is designed for the Cepheid GeneXpert factor II and factor V assays. DNA extraction for other assays/methods is NOT recommended.

CAP/ACMG Thrombophilia Mutations TPM			
Analyte/Procedure Program Code Challenges/Shipment			
ТРМ			
	3		
ı	3		
	Program Code		

Program Information

- Three 250-µL synthetic whole blood specimens
- · Two shipments per year



Red Blood Cell Antigen Genotyping RAG		
Procedure	Program Code	Challenges/Shipment
	RAG	
Red blood cell antigen genotype with predictive phenotype		3

Program Information

- · Three 2.0-mL whole blood specimens
- · Two shipments per year

Variant Interpretation Only VIP/VIP1			
Analyte/Procedure	Program Code	Challenges/Shipment	
	VIP/VIP1		
Variant interpretation online case review	ı	3	

Additional Information

VIP educates pathologists, PhDs, genetic counselors, technologists, and any other laboratory staff with an interest in germline variant interpretation to assess and improve their diagnostic skills. All cases will comply with the 2015 ACMG standards and guidelines for the interpretation of sequence variants and will include:

- · A clinical history with relevant laboratory data
- · Results of ancillary studies, where appropriate
- · Case discussion and discussion of interpretive criteria
- A variety of germline variants, diseases, and disorders
- · Five questions per case

- VIP Three germline diagnostic challenges; reporting with CME/CE credit is available for one pathologist, MD, PhD, technologist, or genetic counselor
- · VIP1 Reporting option with CME/CE credit for each additional pathologist, MD, PhD, technologist, or genetic counselor (within the same institution); must order in conjunction with Survey VIP
- · Earn a maximum of 3 CME credits (AMA PRA Category 1 Credits™) per pathologist/ MD/PhD and a maximum of 3 CE credits per technologist/ genetic counselor for completion of an entire year
- · One online activity per year; your CAP shipping contact will be notified via email when the activity is available



Noninvasive Pre	enatal Testing	NIPT NEW
Analyte	Program Code	Challenges/Shipment
	NIPT	
Cell-free DNA screening for fetal aneuploidy	•	3

Program Information

- Three maternal plasma samples
- · Two shipments per year

Give the CAP's complimentary Sample Exchange Registry service a try!

Sign up for this unique and complimentary service for those rare analytes for which proficiency testing is not yet available. This service now includes all clinical laboratory disciplines.

- The CAP connects labs performing testing for which no formal proficiency testing is available.
- There is no charge for this service.
- · Participate at any time, no contract required.
- A minimum of three labs performing the same analyte test must participate before the CAP can facilitate the sample exchange.
- Each individual laboratory will receive its own results along with an anonymized summary report for all participants.

Register today! Visit cap.org and search Sample Exchange Registry.

Next-Generation Sequencing

Analytes/procedures in bold type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Next-Generation Sequencing NGS		
Procedure	Program Code	Challenges/Shipment
	NGS	
Next-generation sequencing	I	1

Additional Information

Laboratories will have the ability to test up to 200 preselected chromosomal positions within various genes; for a full list of genes in this program, please go to cap.org. Under the Laboratory Improvement tab, click on Catalog and Ordering Information. The list is located under the PT Order Supplements header.

Program Information

- One 10.0-µg extracted DNA specimen
- Methods-based challenge for germline variants for laboratories using gene panels, exome, and whole genome sequencing
- · Results for this program must be submitted online through e-LAB Solutions Suite
- Two shipments per year

hree 1.0-µg DNA (50 ng/µL)

Program Information

Next-Generation Sequencing—Solid Tumor NGSST					
Procedure Program Code Challenges/Shipment					
NGSST					
Next-generation sequencing		3			

Additional Information

This is a methods-based proficiency challenge for laboratories performing targeted next-generation sequencing of cancer genes or mutation hotspots in solid tumors. Laboratories will be asked to identify somatic single nucleotide variants and small insertions or deletions in the following genes: AKT1, ALK, APC, ATM, BRAF, CDH1, CTNNB1, EGFR, ERBB2, FBXW7, FGFR2, GNAQ, GNAS, HRAS, IDH1, KIT, KRAS, MET, NRAS, PDGFRA, PIK3CA, PTEN, SMAD4, SMARCB1, SMO, SRC, STK11, TP53.

	 Three 1.0-μg DNA (50 ng
nt	specimens
	• Two shipments per year

Program Information

- Three 1.0-μg DNA (50 ng/μL) specimens
- · Two shipments per year

Next-Generation Sequencing—Hematologic Malignancies NGSHM				
Procedure Program Code Challenges/Shipment				
NGSHM				
Next-generation sequencing				

Additional Information

This is a methods-based proficiency challenge for laboratories performing targeted next-generation sequencing of genes or mutation hotspots in hematologic malignancies. Laboratories will be asked to identify somatic single nucleotide variants and small insertions or deletions in the following genes: ASXL1, ATM, BRAF, CALR, CEBPA, CREBBP, CSF3R, DNMT3A, EZH2, FLT3, IDH1, IDH2, JAK2, KIT, KMT2D, MPL, MYD88, NOTCH1, NPM1, SF3B1, SRSF2, TET2, TP53, U2AF1.

19

Next-Generation Sequencing Bioinformatics NGSB1, NGSB2

Procedure	Program Code		Challenges/Shipment
	NGSB1 NGSB2		
Illumina TruSeq Amplicon Cancer Panel	ı		1
Ion Torrent AmpliSeq Cancer Hotspot v2			1

Additional Information:

- This in silico bioinformatics program is designed to complement and augment somatic variant wet bench NGS proficiency testing programs by testing a greater diversity of variants at a greater range of variant allele fractions.
- The BAM and/or FASTQ files are platform-specific and may not be compatible with other instruments/software.
- Laboratories will be asked to identify somatic single nucelotide variants and small insertions/deletions/indels in the following genes: ABL1, AKT1, ALK, APC, ATM, BRAF, CDH1, CDKN2A, CSF1R, CTNNB1, EGFR, ERBB2, ERBB4, FBXW7, FGFR1, FGFR2, FGFR3, GNA11, GNAQ, GNAS, HNF1A, HRAS, IDH1, JAK3, KDR, KIT, KRAS, MET, MLH1, MPL, NOTCH1, NPM1, NRAS, PDGFRA, PIK3CA, PTEN, PTPN11, RB1, RET, SMAD4, SMARCB1, SMO, SRC, STK11, TP53, VHL.

- Sequencing files containing somatic variants to be downloaded into your laboratory bioinformatics pipeline for analysis and reporting; file sizes range from 100MB to 1GB
- NGSB1 FASTQ file format for the Illumina TruSeq Amplicon Cancer Panel
- NGSB2 BAM and FASTQ file formats for the Ion Torrent AmpliSeq Cancer Hotspot v2 Panel
- This is an online only program, delivered two times a year; your CAP shipping contact will be notified via email when the activity is available

Next-Generation Sequencing Undiagnosed Disorders—Exome NGSE Analyte/Procedure Program Code Challenges/Shipment NGSE Exome analysis for germline undiagnosed disorders

Additional Information/Minimum Requirements

- This in silico based Survey will assess the ability of the laboratory to identify germline variants responsible for a provided clinic phenotype as is encountered in an undiagnosed disease scenario.
- Laboratories are expected to provide an exome sequencing data file that has been generated using one of the following sources: a specimen from the NGS Survey program (see page 246) or from one of the NIST Reference Material cell lines: RM 8398 (NA12878), RM 8391, RM 8392, or RM 8393.
- FASTQs or unaligned BAMs should be submitted along with a BED file describing
 the regions targeted and interrogated by your laboratory. Additionally, >90% of
 exons targeted and interrogated by your laboratory must have a minimum read
 coverage of 10X.
- Laboratories can transfer and download files from most modern browsers/ operating systems:
 - o Internet Explorer (IE) 11
 - o Safari The two latest, released versions on Mac OS X and iOS
 - o Firefox The two latest, released versions
 - o Chrome The two latest, released versions
 - o Windows 7 (32-bit and 64-bit), 8 (64-bit), and 10 (32-bit and 64-bit)
- Due to the extremely large file sizes, a minimum allowable transfer speed
 of 20Mbps will be needed to ensure the successful transfer of sequencing
 files between laboratories and CAP; however, 40 Mbps and higher is strongly
 recommended. Note: Laboratories should check with their technology department
 for allowable transfer speeds to determine estimated transfer time and browser/
 operating system access.
- Laboratories must comply with the above requirements to participate in this
 program. Additional information regarding how and where to provide your
 laboratory's exome file will be sent closer to the ship date.

- One in silico mutagenized exome sequencing file to be downloaded and analyzed by your laboratory bioinformatics pipeline
- The exome sequencing file will be accompanied by a clinical history, relevant laboratory data, and results of ancillary studies, where appropriate
- Two online activities per year; your CAP shipping contact will be notified <u>via email</u> when the activity is available

Next-Generation Sequencing Bioinformatics Somatic Validated Materials NGSBV

Analyte/Procedure	Program Code	Challenges/Shipment
	NGSBV	
Somatic in silico mutagenized sequencing file	ı	1

Additional Information/Minimum Requirements

- This in silico program is designed to optimize bioinformatics pipelines, augment validations, and assist with pipeline verification after changes to NGS/ bioinformatics processes. This is not traditional proficiency testing and no results will be returned to the CAP; information regarding the variants introduced will be sent along with the mutagenized file.
- Laboratories are expected to provide a gene panel or exome sequencing data file that has been generated using one of the following sources: a specimen from the NGS Survey program (see page 246) or from one of the following NIST Reference Material cell lines: RM 8398 (NA12878), RM 8391, RM 8392, or RM 8393.
- FASTQs or unaligned BAMs should be submitted along with a BED file describing the regions targeted and interrogated by your laboratory.
- The mutagenized sequencing file will contain up to 50 somatic variants (depending
 on the size of the panel/exome provided) at allele fractions from 5% to 50% and will
 include both single nucleotide variants and insertions/deletions, the latter ranging
 from 1-15bp. All variants will be modeled based on actual somatic mutations from
 the COSMIC database.
- Laboratories can transfer and download files from most modern browsers/ operating systems:
 - o Internet Explorer (IE) 11
 - o Safari The two latest, released versions on Mac OS X and iOS
 - o Firefox The two latest, released versions
 - o Chrome The two latest, released versions
 - o Windows 7 (32-bit and 64-bit), 8 (64-bit), and 10 (32-bit and 64-bit)
- Due to the extremely large file sizes, a minimum allowable transfer speed
 of 20Mbps will be needed to ensure the successful transfer of sequencing
 files between laboratories and CAP; however, 40 Mbps and higher is strongly
 recommended. Note: Laboratories should check with their technology department
 for allowable transfer speeds to determine estimated transfer time and browser/
 operating system access.
- Laboratories must comply with the above requirements to participate in this
 program. Additional information regarding how and where to provide your
 laboratory's sequencing file will be sent closer to the ship date.

- One challenge containing in silico mutagenized somatic variants to be downloaded into your bioinformatics pipeline
- Gene panel or exome files may be submitted
- Two online activities per year; your CAP shipping contact will be notified <u>via email</u> when the activity is available

10

Molecular Oncology—Solid Tumors

Analytes/procedures in bold type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Microsatellite Instability (HNPCC) MSI			
Procedure Program Code Challenges/Shipme			
	MSI		
Microsatellite instability testing (DNA amplification)		3	
MLH1 promoter methylation analysis	I	1	

Laboratories performing DNA mismatch repair assessment by immunohistochemistry methods should see Survey MMR on page 267.

Program Information

- Two 10.0-micron unstained paraffin section slides and one H&E slide; two photograph challenges
- For laboratories performing molecular testing using PCR
- · Two shipments per year

IGHV Mutation Analysis IGHV		
Analyte/Procedure	Program Code	Challenges/Shipment
	IGHV	
IGHV		3

Additional Information

- Sequence analysis of the clonal immunoglobulin heavy chain V gene (*IGHV*) to determine somatic hypermutation (SHM) status.
- · Any sequencing method may be used.
- Report V-gene allele, percent similarity and mutation status (SHM).

In Situ Hybridization ISH, ISH2			
Analyte/Procedure	Program Code		Challenges/Shipment
	ISH	ISH2	
Epstein-Barr virus (EBV)			4
Human papillomavirus (HPV)			4
Kappa/Lambda (IGK/IGL)			4
HER2 (ERBB2) gene amplification (brightfield)			10

Laboratories performing FISH for interphase chromosomal targets in paraffin sections refer to the Cytogenetics Surveys, page 237.

Program Information

- Three 20-µg DNA specimens (200 ng/µL)
- Two shipments per year

- ISH
 - EBV, HPV: Three 4-core tissue microarray slides and one H&E slide (each)
 - Kappa/Lambda: Four
 4-core tissue microarray
 slides and one H&E slide
- ISH2 Two 5-core tissue microarray slides in duplicate
- Two shipments per year

Program Information

sections

• Three 10.0-micron paraffin

• Two shipments per year

DNA Extraction & Amplification FFPE MH05					
Procedure Program Code Challenges/Shipment					
	MH05				
DNA purification	I	1			

Additional Information

Methods-based proficiency challenge to examine DNA purification from formalin-fixed, paraffin-embedded tissues (FFPET). Laboratories will be able to purify DNA from FFPET sections and amplify control targets using laboratory-provided reagents.

Neoplastic Cellularity NEO				
Procedure	Program Code Challenges/Shipmen			
	NEO			
Online assessment of percent neoplastic cellularity	ı	10		

Program Information

- Ten Regions of Interests (ROIs) using whole slide images
- A method-based preanalytic Survey to assess competency for determining percent neoplastic cellularity
- Powered by DigitalScope® technology
- Individual reporting fields for up to five pathologists are available
- Two online activities per year; your CAP shipping contact will be notified <u>via email</u> when the activity is available

Sarcoma Translocation SARC					
Gene	e Program Code Challenges/Shipment				
	SARC				
Sarcoma translocation* (RT-PCR) ■ 3					

^{*}See translocation listing below.

Laboratories performing FISH for sarcoma translocation refer to the Cytogenetics Surveys, page 238.

Program Information

- Snap-frozen cell pellet from which approximately 5.0-µg of RNA can be extracted
- Two shipments per year

Sarcoma Translocation Listing

COL1A1/PDGFB, t(17;22)	EWSR1/FLI1 or EWSR1/ERG	PAX3/F0X01 or PAX7/F0X01
ETV6-NTRK3, t(12;15)	EWSR1/WT1, t(11;22)	SS18/SSX1, t(X;18)
EWSR1/ATF1, t(12;22)	FUS/DDIT3, t(12;16)	SS18/SSX2, t(X;18)
EWSR1/ERG, t(21;22)	PAX3/FOXO1, t(2;13)	SS18/SSX1 or SS18/SSX2
EWSR1/FLI1. t(11:22)	PAX7/F0X01. t(1:13)	



Cell Free DNA	CFDNA	NEW
Analyte/Procedure	Program Code	Challenges/Shipment
	CFDNA	
cfDNA	ı	3

Program Information

- Three 300-ng DNA (10 ng/µL) specimens
- Two shipments per year

Additional Information

- Mix of fragmented cell line gDNA and biosynthetic DNA.
- This is not intended for laboratories that perform circulating tumor cell (CTC) analysis.
- Potential targets included in this Survey are BRAF V600E, EGFR T790M, IDH1 R132C, KRAS G12D, and NRAS Q61R, all within a range of 0.1 to 1.0%.

RNA Sequenci	NEW	
Analyte/Procedure	Program Code	Challenges/Shipment
	RNA	
RNA	I	3

Program Information

- Three 500-ng RNA (10 ng/μL) specimens
- Two shipments per year

Additional Information

- Total RNA from a cell line engineered to contain desired fusion RNA.
- This is for laboratories using RNAseq to detect gene fusion transcripts.
- This is not intended to replace the current Survey (SARC) for reverse transcription (RT)-PCR based detection (see page 251).
- Potential fusion variants include: CD74-ROS1, EML4-ALK, ETV6-NTRK3, FGFR3-TACC3, PAX8-PPARG, SLC45A3-BRAF.

Solid Tumor—Other BRAF, EGFR, KRAS, KIT							
Analyte		Program Code Challenges/Shipment					
	BRAF	BRAF EGFR KRAS KIT					
BRAF	1				3		
EGFR					3		
KRAS		3					
KIT				1	3		
PDGFRA		■ 3					

- · BRAF, EGFR, KRAS -Paraffin-embedded sections or shavings
- KIT/PDGFRA Four 10.0-micron unstained paraffin section slides and one H&E slide, for each specimen
- For laboratories performing molecular testing using PCR
- · Two shipments per year

Multigene Tumor Panel MTP				
Analyte	Program Code	Challenges/Shipment		
	MTP			
BRAF	ı	3		
EGFR	ı	3		
HER2 (ERBB2)	I	3		
KIT	ı	3		
KRAS	ı	3		
NRAS	I	3		
PDGFRA	I	3		
PIK3CA		3		

This program meets CAP Accreditation requirements for BRAF, EGFR, and KRAS for laboratories that perform testing for the detection of somatic single nucleotide variants, insertions, and deletions in these genes. Laboratories performing this testing must enroll in either MTP or the respective single gene PT programs to meet CAP Accreditation requirements; this includes laboratories that perform NGS-based assays, non-NGS-based multiplexed assays or non-multiplexed assays (eg, Sanger sequencing).

Glioma GLI					
Analyte Program Code Challenges/Shipment					
GLI					
MGMT		2			
IDH1, IDH2	I	3			
10q (PTEN) deletion		1			

Program Information

- Three 2.0-µg gDNA (50 ng/μL) specimens for laboratories performing molecular testing on multiple targets
- · Two shipments per year

- Two 2.0-ug gDNA (50 ng/uL) specimens
- · One specimen containing four 10.0-micron unstained paraffin section slides and one H&E slide
- · For laboratories performing molecular testing using PCR
- Two shipments per year

Molecular Oncology—Hematologic

Analytes/procedures in bold type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Molecular Hematologic Oncology MHO, MHO1, MHO2, MHO3, MHO5					
Procedure/Gene		Challenges/ Shipment			
	MH0, MH01	MH02, MH03	MH05		
Lymphoid malignancy genotyp	ing				
IGH	ı			3	
IGH/BCL2 major	ı			3	
IGH/BCL2 minor	ı			3	
IGH/CCND1	•			3	
IGK	ı			3	
TRB	ı			3	
TRG	ı			3	
Myeloid malignancy genotypin	g				
BCR/ABL1 p190 NEW				1	
BCR/ABL1 p210 NEW		•		1	
CBFB/MYH11		•		1	
FLT3 ITD		•		1	
FLT3 TKD		•		1	
JAK2 c.1849G>T(p.V617F)		•		1	
NPM1		•		1	
PML/RARA		•		1	
RUNX1/RUNX1T1		ı		1	
DNA extraction and amplification from formalin-fixed, paraffin-embedded (FFPE) tissue				1	

Program Information

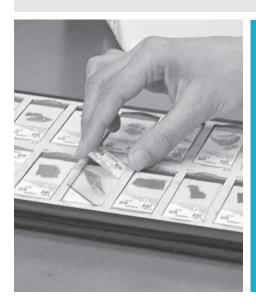
- MHO One sample vial containing purified DNA (200 µg/mL per vial) for each specimen
- MH01 MH0 specimens in duplicate for additional DNA testing
- MHO2 Two sample vials; one with purified DNA containing 200 µg/mL and one with purified RNA containing 400 µg/mL
- MH03 MH02 specimen in duplicate for additional DNA and RNA testing
- MH05 Three 10.0-micron paraffin sections; extraction and amplification from FFPE tissue will be assessed by a method-based challenge
- Two shipments per year; ships on dry ice (dry ice does not apply to MHO5 or international shipments)

Minimal Residual Disease MRD, MRD1, MRD2					
Analyte	1	Program Code		Challenges/ Shipment	
	MRD	MRD1	MRD2		
BCR/ABL1 p190				3	
BCR/ABL1 p210				3	
PMI /RARA				3	

- Three RNA specimens in distilled water
- For laboratories diagnosing and monitoring leukemia tumor burden by measuring the quantity of BCR/ABL1 or PML/RARA fusion transcripts
- Two shipments per year; ships on dry ice

Anatomic Pathology

2 Anatomic Pathology



Evaluate your laboratory's tissue and slide preparation techniques with CAP/NSH HistoQIP Surveys.

- Submit your own laboratory's slides and have the stain quality graded by an expert panel of pathologists, histotechnologists, and histotechnicians.
- Participate in three new programs evaluating:
 - Gynecologic biopsy specimen slide preparation
 - Mismatch repair protein IHC staining
 - IHC staining of non-small cell lung carcinoma

Anatomic Pathology

Surgical Pathology	256
General Immunohistochemistry	
Predictive Markers	
Specialty Anatomic Pathology	270
Cytopathology	272
2 , 2	

New Programs NEW



CAP/NSH HistoQIP Mismatch Repair IHC (HQMMR)	. 264
CAP/NSH HistoQIP Non-small Cell Lung Carcinoma IHC (HQNSC)	
CAP/NSH HistoQIP Specialty Series (HQBX4)	

Surgical Pathology

Analytes/procedures in bold type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Performance Improvement Program in Surgical Pathology PIP/PIP1					
Program Code Challenges/Shipment					
	PIP/PIP1				
Surgical pathology case review ■ 10					

Additional Information

PIP educates pathologists in general surgical pathology. This program:

- · Provides a practical approach to continuing education
- Gives pathologists a method to assess their diagnostic skills and compare their performance with that of their peers
- · Features PIP case selections that include:
 - O A variety of neoplastic and nonneoplastic lesions
 - o Inflammatory and infectious disease
 - o Various sites, encompassing a variety of organ systems

Program Information

- PIP Ten diagnostic challenges/H&E stained glass slides with clinical history; reporting with CME credit is available for one pathologist; for each additional pathologist, order PIP1
- PIP1 Reporting option with CME credit for each additional pathologist (within the same institution); must order in conjunction with Survey PIP
- Earn a maximum of 40 CME credits (AMA PRA Category 1 Credits™) per pathologist for completion of an entire year
- This activity meets the ABP MOC Part IV Practice Performance Assessment requirements
- Four shipments per year

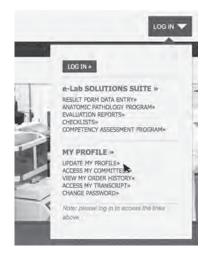


Have you created or updated your CAP Profile?

Each laboratory staff member should have their own profile. Your profile is transferrable when you leave your current position. Use it to maintain information about yourself, including:

- Business affiliations
- Certifications
- · Contact preferences
- Inspector-related information
- · Personal contact information
- Specialties and skills
- Addresses

To create or update your profile, visit cap.org, log in, and click on UPDATE MY PROFILE.



Online Performance Improvement Program in Surgical Pathology PIPW/PIPW1				
Program	Program Code Challenges/Shipment			
PIPW/PIPW1				
Surgical pathology case review	I	10		

PIPW educates pathologists in general surgical pathology.

- Pathologists can assess their diagnostic skills and compare their performance with that of their peers.
- · Included PIPW case selections feature:
 - O A variety of neoplastic and nonneoplastic lesions
 - o Inflammatory and infectious disease
 - O Various sites, encompassing a variety of organ systems
- See system requirements on page 13.

Program Information

- PIPW Ten diagnostic challenges/whole slide H&E images with clinical history; reporting with CME credit is available for one pathologist; for each additional pathologist, order PIPW1
- PIPW1 Reporting option with CME credit for each additional pathologist (within the same institution); must order in conjunction with Survey PIPW
- Earn a maximum of 40 CME credits (AMA PRA Category 1 Credits™) per pathologist for completion of an entire year
- This activity meets the ABP MOC Part IV Practice Performance Assessment requirements
- Powered by DigitalScope® technology
- Four online activities per year; your CAP shipping contact will be notified <u>via</u> <u>email</u> when the activity is available



Test your diagnostic skills as a pathologist with CPIP

Online, hands-on and interactive, the Clinical Pathology Improvement Program (CPIP) enables pathologists to sharpen their diagnostic skills in real time by working through an actual case. Each month, you will receive a new scenario, including slide images and clinical background. As the case unfolds, more information is revealed, just as in the laboratory. Participants who successfully complete the posttest may apply their earned credits to their MOC SAM requirements. Enjoy a full year of CPIP and earn up to 15 CME/SAM credits.

Choose code CPIP/CPIP1 on your Surveys order form.

Virtual Biopsy Program VBP/VBP1				
Program	Program Code Challenges/Shipment			
	VBP/VBP1			
Online biopsy case review		5		

VBP educates pathologists to assess and improve their diagnostic skills in surgical pathology.

- Cases may include gross, radiographic, or endoscopic images.
- Cases are from selected organ systems and may include a variety of specimen types (eg, core biopsies, endoscopic biopsies, curettings, aspirate smears). Activities with their corresponding topics are:
 - o 2018-A Liver Biopsy
 - o 2018-B Soft Tissue Biopsy
 - o 2018-C Prostate Biopsy
 - o 2018-D Surgical Pathology Biopsy (various sites)
- See system requirements on page 13.

- VBP Five diagnostic challenges/whole slide images with clinical history; reporting with CME/SAM credit is available for one pathologist; for each additional pathologist, order VBP1
- VBP1 Reporting option with CME/SAM credit for each additional pathologist (within the same institution); must order in conjunction with Survey VBP
- Earn a maximum of 25 CME/SAM credits (AMA PRA Category 1 Credits[™]) per pathologist for completion of an entire year
- This activity meets the ABP MOC Part IV Practice Performance Assessment requirements
- Powered by DigitalScope technology
- Four online activities per year; your CAP shipping contact will be notified <u>via</u> <u>email</u> when the activity is available





Digital Slide Program—Dermatopathology DPATH/DPATH1					
Program Code Challenges/Shipmen					
	DPATH/DPATH1				
Online dermatopathology case review 6					

DPATH educates pathologists, dermatopathologists, and dermatologists to assess and improve their diagnostic skills in dermatopathology.

- · Cases include static images.
- See system requirements on page 13.

Program Information

- DPATH Six diagnostic challenges/whole slide images with clinical history; reporting with CME/SAM credit is available for one pathologist; for each additional pathologist, order DPATH1
- DPATH1 Reporting option with CME/SAM credit for each additional pathologist (within the same institution); must order in conjunction with Survey DPATH
- Earn a maximum of 15 CME/SAM credits (AMA PRA Category 1 Credits™) per pathologist for completion of an entire year
- This activity meets the ABP MOC Part IV Practice Performance Assessment requirements
- Powered by DigitalScope technology
- Two online activities per year; your CAP shipping contact will be notified <u>via email</u> when the activity is available





Experience a new level of pathology education with SAM courses from the CAP

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- Peer reviewed by at least two subject matter experts
- Highly interactive formats with immediate feedback

Visit cap.org and choose the Learning tab, then search SAM Courses.



Hematopathology Online Education HPATH/HPATH1					
Program	Program Code Challenges/Shipment				
	HPATH/HPATH1				
Hematopathology online case review	ı	5			

HPATH educates pathologists, hematolopathologists, and hematologists with an interest in hematopathology to assess and improve their diagnostic skills in hematopathology.

- All cases have been specially selected to highlight important changes in the 2016 revision of the WHO Classification.
- · Clinical history and relevant laboratory data.
- At least one online whole slide image of peripheral blood, bone marrow, spleen, lymph node, or other tissue.
- Results of ancillary studies such as immunohistochemistry, flow cytometry, FISH, karyotyping, and molecular studies, where appropriate.
- Case discussion and discussion of differential diagnoses.
- · Five SAM questions per case.
- See system requirements on page 13.

- HPATH Five diagnostic challenges/online whole slide images with clinical history; reporting with CME/SAM credit is available for one pathologist/ hematologist. For additional pathologist/hematologist, order HPATH1
- HPATH1 Reporting option with CME/SAM credit for each additional pathologist and hematologist (within the same institution); must order in conjunction with Survey HPATH
- Earn a maximum of 12.5 CME/SAM credits (AMA PRA Category 1 Credits™) per pathologist and a maximum of 12.5 CE credits per hematologist for completion of an entire year
- This activity meets the ABP MOC Part IV Practice Performance Assessment requirements
- Powered by DigitalScope technology
- Two online activities per year; your CAP shipping contact will be notified <u>via email</u> when the activity is available





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Touch Imprint/Crush Preparation TICP/TICP1			
Procedure	Program Code	Challenges/Shipment	
	TICP/TICP1		
Online slide and image program in rapid assessment case review	•	4	

- The TICP Program is designed to familiarize surgical pathologists, cytopathologists, and cytotechnologists with the cytomorphologic features of pathologic processes and tumors in touch imprints and crush or scrape preparations. These specimens are prepared either for intraoperative consultation (frozen section) or rapid on-site evaluation (ROSE) of tissue biopsies for adequacy and/or interpretation. Participants will learn to make an immediate adequacy assessment, assign the process to a general category, and triage the specimen to appropriate ancillary studies. Participants will review digital whole slides of the TICP preparations (hematoxylin & eosin, modified Wright-Giemsa, and/or Papanicolaou stains), static images of the preparation and ancillary studies, and clinical history/radiographic findings to reach a diagnosis. Each case has a complete description of entities in the differential diagnosis along with a discussion of the correct interpretation.
- · Participants will receive immediate feedback on interpretations, ancillary studies, and case-related adequate assessment.
- The 2018 cases will be comprised of specimens from the central nervous system and soft tissue.
- See system requirements on page 13.

- · TICP Four online assessment challenges with clinical history; for each additional pathologist or cytotechnologist, order TICP1
- TICP1 Reporting option with CME/SAM/CE credit for each additional pathologist/ technologist (within the same institution); must order in conjunction with Survey TICP
- Earn a maximum of 10 CME/SAM credits (AMA PRA Category 1 Credits™) per pathologist and a maximum of 10 CE credits per cytotechnologist for completion of an entire year
- This activity meets the ABP MOC Part IV Practice Performance Assessment requirements
- Online whole slide images powered by DigitalScope technology
- Two online activities per year; your CAP shipping contact will be notified via email when the activity is available





Cancer Staging Improvement Program PCSP/PCSP1		
Program	Program Code	Challenges/Shipment
	PCSP/PCSP1	
Online surgical pathology cancer case review	ı	4

PCSP educates pathologists in cancer case review and reporting. This program features four challenging online cases to stage and classify a cancer as defined by the *American Joint Committee on Cancer (AJCC) Cancer Staging Manual*, 8th edition, and synoptically report out using the CAP Cancer Protocols.

- · Cases review challenges in interpretation, staging, and cancer reporting.
- See system requirements on page 13.

- PCSP One mailing containing four diagnostic challenges/whole slide H&E images with clinical history; reporting with CME/ SAM credit is available for one pathologist; for each additional pathologist, order PCSP1
- PCSP1 Reporting option with CME/SAM credit for each additional pathologist (within the same institution); must order in conjunction with Survey PCSP
- Earn a maximum of 5 CME/SAM credits (AMA PRA Category 1 Credits™) per pathologist for completion of an entire year
- Powered by DigitalScope technology
- This activity meets the ABP MOC Part IV Practice Performance Assessment requirements
- One online activity per year; your CAP shipping contact will be notified via email when the activity is available





CAP/NSH HistoQIP HQIP			
Stain/Tissue	Program Code Challenges/Shipment		
	HQIP	Α	В
H&E – Fallopian tube resection	ı	1	
H&E – Small intestine resection	ı	1	
IHC – SMA (small intestine resection)	ı	1	
IHC – Pancytokeratin (fallopian tube)	ı	1	
Special Stain – Congo red (tissue with amyloid)	ı	1	
H&E – Gallbladder resection	ı		1
H&E – Melanoma from skin resection	ı		1
IHC – Ki67 (tonsil resection)	ı		1
IHC – S100 (melanoma from skin resection)	ı		1
Special Stain – Mucicarmine (gallbladder resection)	ı		1

Program Information

- Participant laboratories may submit up to five stained and coverslipped glass slides (one from each category) per mailing
- Includes photographs and online learning assessment questions
- · Two shipments per year





Additional Information

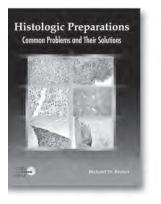
HistoQIP improves the preparation of histologic slides in all anatomic pathology laboratories. In this educational program, participants will receive an evaluation specific to their laboratory, an education critique, and a Participant Summary that includes peer comparison data, evaluators' comments, and performance benchmarking data. An expert panel of pathologists, histotechnologists, and histotechnicians will evaluate submitted slides for histologic technique using uniform grading criteria.

Learn the secret to good slide technique

Histologic Preparations: Common Problems and Their Solutions is a how-to guide to good slide preparation. Building on data and images from the CAP/NSH HistoQIP program, the book presents photographic examples of well-prepared slides followed by numerous examples of associated problems and their solutions. The text contains troubleshooting techniques for the most common artifacts and problems incurred in routine histologic preparations, including fixation and processing; microtomy; frozen sections; hematoxylin-eosin, trichrome, reticulin, elastin, basement membrane, mucin, amyloid, immunohistochemical, and Gram stains; and mycobacteria, Helicobacter pylori, sprirochetes, and fungi.

View sample pages and order online:

- · printed books at estore.cap.org
- · ebooks at ebooks.cap.org



Item number: PUB123 Softcover; 168 pages; 300+ photomicrographs, figures, and tables; 2009

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CAP/NSH HistoQIP—IHC HQIHC			
Stain/Tissue	Program Code	Challenge	es/Shipment
	HQIHC	Α	В
IHC – CK20 (bladder biopsy)	ı	1	
IHC – Progesterone receptor (cervical biopsy)	ı	1	
IHC – CD34 (skin, punch biopsy)	ı	1	
IHC – CD138 (stomach biopsy)	ı	1	
IHC – CD3 (colon biopsy)	ı		1
IHC – EMA (endometrium)	ı		1
IHC – S100 (skin, excisional biopsy)			1
IHC – p504s (prostate biopsy with carcinoma)			1

Program Information

- Participants may submit up to four IHC stained and coverslipped glass slides (one from each category) per mailing
- Two shipments per year



Additional Information

HistoQIP – IHC improves the preparation of immunohistochemistry slides in all anatomic laboratories involved in the handling of gastrointestinal, dermatologic, and urological tract biopsies. Participants will receive an evaluation specific to their laboratory and a Participant Summary. An expert panel of pathologists, histotechnologists, and histotechnicians will evaluate submitted slides for histologic technique using uniform grading criteria.

CAP/NSH HistoQIP Mismatch Repair IHC



HQMMR			
Stain/Tissue	Program Code	Challenges	s/Shipment
	HQMMR	Α	В
H&E – Colon adenocarcinoma		1	
IHC – MLH1 (colon adenocarcinoma)		1	
IHC – MSH2 (colon adenocarcinoma)	•	1	
IHC – MSH6 (colon adenocarcinoma)	•	1	
IHC – PMS2 (colon adenocarcinoma)	•	1	
H&E – Endometrial adenocarcinoma	•		1
IHC – MLH1 (endometrial adenocarcinoma)	•		1
IHC – MSH2 (endometrial adenocarcinoma)	•		1
IHC – MSH6 (endometrial adenocarcinoma)	•		1
IHC – PMS2 (endometrial adenocarcinoma)	ı		1

Additional Information

This program augments efforts to improve the preparation of H&E and immunohistochemical slides in all anatomic pathology laboratories involved in the handling of colonic and endometrial tumors performing mismatch repair IHC.

- Participants may submit up to four IHC and one H&E stained coverslipped glass slides (one from each category) per mailing
- · Two shipments per year



CAP/NSH HistoQIP Non-small Cell Lung Carcinoma IHC HQNSC

Stain/Tissue	Program Code	Challenge	s/Shipment
	HQNSC	Α	В
H&E – Lung adenocarcinoma	ı	1	
IHC – TTF-1 (lung adenocarcinoma)	ı	1	
IHC – Napsin-A (lung adenocarcinoma)	•	1	
H&E – ALK (positive lung adenocarcinoma)	ı	1	
IHC – ALK (positive lung adenocarcinoma)	ı	1	
H&E – Lung squamous cell carcinoma	•		1
IHC – p40/p63 (lung squamous cell carcinoma)	ı		1
IHC – CK5/6 (lung squamous cell carcinoma)	ı		1
H&E – PD-L1 (positive lung squamous cell carcinoma)	ı		1
IHC – PD-L1 (positive lung squamous cell carcinoma)	ı		1

Program Information

- · Participants may submit up to four IHC and one H&E stained coverslipped glass slides (one from each category) per mailing
- · Two shipments per year



Additional Information

This program augments efforts to improve the preparation of H&E and immunohistochemical slides in all anatomic pathology laboratories involved in the handling of non-small cell lung carcinoma.

CAP/NSH HistoQIP Biopsy Series HQIPBX			
Stain/Tissue	Program Code	Challenge	s/Shipment
	HQIPBX	Α	В
H&E – Bladder biopsy	•	1	
H&E – Cervical biopsy	•	1	
H&E – Skin punch biopsy	•	1	
H&E – Stomach biopsy	•	1	
H&E – Colon biopsy	•		1
H&E – Endometrial biopsy	•		1
H&E – Prostate needle biopsy			1
H&E – Breast core biopsy	•		1

Additional Information

The HistoQIP Biopsy Series is an additional program to improve the preparation of histologic slides in all anatomic pathology laboratories. Participants will receive an evaluation specific to their laboratory and a Participant Summary. An expert panel of pathologists, histotechnologists, and histotechnicians will evaluate submitted slides for histologic technique using uniform grading criteria.

- · Participant laboratories may submit up to four H&E stained and coverslipped glass slides (one from each category) per mailing
- Two shipments per year



CAP/NSH HistoQIP Specialty Series HQBX1, HQBX2, HQBX3, HQBX4 Challenges/ Stain/Tissue **Program Code** Shipment HQBX1 HQBX2 HQBX3 HQBX4 NEW Α В Gastrointestinal Biopsy Module H&E - Colon biopsy 1 1 1 1 H&E - Esophageal biopsy 1 1 H&E – Small intestinal biopsy 1 H&E - Stomach biopsy 1 Dermatologic Biopsy Module H&E - Alopecia 1 H&E - Skin excisional biopsy 1 1 (large excision) 1 H&E - Skin punch biopsy 1 H&E - Skin shave biopsy 1 1 Urogenital Tract Biopsy Module H&E - Bladder biopsy ī 1 1 (nonneoplastic) H&E - Bladder biopsy (with П 1 1 carcinoma) H&E - Prostate needle biopsy 1 1 (nonneoplastic) H&E - Prostate needle biopsy ī 1 1 (with carcinoma) **Gynecological Biopsy** 1 1 H&E - Cervical biopsy H&E - Endometrial biopsy 1 1 H&E - Cone/Leep biopsy 1 1

Additional Information

H&E - Vagina biopsy

The HistoQIP Specialty Series includes modules to improve the preparation of histologic slides in all anatomic pathology laboratories involved in the handling of gastrointestinal, dermatologic, gynecologic, and urogenital tract biopsies. Participants will receive an evaluation specific to their laboratory and a Participant Summary. An expert panel of pathologists, histotechnologists, and histotechnicians will evaluate submitted slides for histologic technique using uniform grading criteria.

- HQBX1, HQBX2, HQBX3, HQBX4 - Participants may submit up to four H&E stained and coverslipped glass slides (one from each category) per mailing
- · Two shipments per year



General Immunohistochemistry

Analytes/procedures in **bold** type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

BRAF V600E BRAFV			
Procedure Program Code Challenges/Shipment			
BRAFV			
BRAF V600E	I	10	

Program Information

- One 10-core tissue microarray slide
- · One shipment per year

Immunohistochemistry MK			
Procedure	Program Code	Challenges/Shipment	
	MK		
Immunohistochemistry	ı	16	

The MK program allows laboratories to compare their assay methodology and results with all participating laboratories.

Program Information

- · Seven glass slides with unstained tissue sections from four separate cases; additional slides provided for an H&E stain and negative control
- Two shipments per year

DNA Mismatch Repair MMR			
Procedure Program Code Challenges/Shipment			
	MMR		
DNA mismatch repair by immunohistochemistry	I	1	

If your laboratory performs DNA mismatch repair by molecular methods, see the MSI program on page 250.

_		
Program	Inform	atior
1 10514111		MCIO!

- Four 4.0-micron unstained paraffin section slides and one H&E slide for the immunohistochemical analysis of DNA mismatch repair proteins MLH1, MSH2, MSH6, and PMS2
- Two shipments per year

PD-L1 PDL1			
Procedure Program Code Challenges/Shipment			
	PDL1		
PD-L1		10	

- One 10-core tissue microarray slide
- · One shipment per year

CD117, CD20 Immunohistochemistry Tissue Microarray PM1, PM3			
Analyte	Program Code Challenges/Shipment		
	PM1	PM3	
CD117			10
CD20		I	10

For ER/PgR testing, see the PM2 program on page 269.

Program Information

- One 10-core tissue microarray slide per predictive marker
- · One shipment per year

Immunohistochemistry Tissue Microarray Series PM5				
Analyte	yte Program Code Challenges/Shipment			
PM5				
Glypican-3	I	10		
Мус	ı	10		

Program Information

- Two 10-core tissue microarray slides, one for Glypican-3 and one for Myc
- · One shipment per year

Additional Information

Each year, the PM5 program will feature two different markers for immunohistochemistry laboratories to evaluate assay performance on a variety of tissue and/or tumor types.

Anaplastic Lymphoma Kinase IHC PM6				
Procedure Program Code Challenges/Shipm				
PM6				
Anaplastic lymphoma kinase IHC (ALK)	ı	10		

Program Information

- One 10-core tissue microarray slide
- · One shipment per year

Let the CAP connect you to the IHC samples you need

CAP Immunohistochemistry (IHC) Validation Program

- The CAP will facilitate the exchange of tissue samples once a sufficient number of laboratories performing the same marker are identified.
- Samples will be exchanged twice a year based on availability.
- Each laboratory will receive its own individual results along with an anonymized summary report for all participants.

Sign up for this complimentary service to access those hard-to-obtain specimens.

To get started, visit cap.org and search Sample Exchange Registry to learn more and download a Contact Information Form.

Predictive Markers

Analytes/procedures in **bold** type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

HER2 Immunohistochemistry HER2			
Analyte Program Code Challenges/Shipment			
HER2			
HER2	I	20	

Program Information

- Two 10-core tissue microarray slides
- · Two shipments per year

Additional Information

The HER2 program fulfills the proficiency testing requirement stated in the ASCO/CAP HER2 Testing Guideline. Due to the unique nature of these human, donor-based materials, the shipping date is subject to change. If this should occur, the CAP will provide notification prior to the originally scheduled shipping date.

Gastric HER2 GHER2					
Analyte Program Code Challenges/Shipment					
	GHER2				
HER2	I	10			

Additional Information

The interpretive criteria for HER2 immunohistochemistry performed on gastroesophageal adenocarcinomas differs significantly from breast carcinoma. The GHER2 program will help participating laboratories understand these differences.

ER/PgR Immunohistochemistry Tissue Microarray PM2					
Analyte	Program Code Challenges/Shipment				
	PM2				
Estrogen receptor (ER)	•	20			
Progesterone receptor (PgR)		20			

Additional Information

The PM2 program fulfills the proficiency testing requirement stated in the ASCO/CAP ER/PgR Testing Guideline. Due to the unique nature of these human, donor-based materials, the shipping date is subject to change. If this should occur, the CAP will provide notification prior to the originally scheduled shipping date.

Program Information

- One 10-core tissue microarray slide
- Two shipments per year

- Four 10-core microarray slides, two for ER and two for PgR
- Two shipments per year

Specialty Anatomic Pathology

Analytes/procedures in bold type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Autopsy Pathology AUP/AUP1				
Procedure	Program Code Challenges/Shipment			
	AUP/AUP1			
Autopsy online case analysis	ı	5		

Each case includes case description, gross and/or microscopic images, and case discussion with sample death certificate, key teaching points, and current references.

Program Information

- AUP Online activity providing five cases; reporting with CME/SAM credit is available for one pathologist; for each additional pathologist, order AUP1
- Includes the option to download program content
- AUP1 Reporting option with CME/SAM credit for each additional pathologist (within the same institution); must order in conjunction with Survey AUP
- Earn a maximum of 12.5 CME/SAM credits (AMA PRA Category 1 Credits™) per pathologist
- This activity meets the ABP MOC Part IV Practice Performance Assessment requirements
- · Two online activities per year



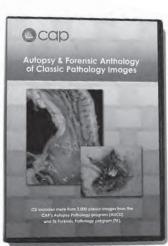


Autopsy & Forensic Anthology of Classic Pathology Images (AFA)

This CD-ROM offering contains more than 2,000 classic images from the CAP's 1992–2010 Autopsy Pathology (AUCD) and the 1990–2010 Forensic Pathology (FR) programs.

- View images in three different modes:
 - List—Cases listed by program, year, patient age, and diagnosis
 - Browse—Images filtered by category and/or image type
 - o Search—Access images via key words or word fragments
- Customize the anthology by adding your own images and categories
- Use only a standard Web browser—AFA runs on any operating system—with no software to install

Choose code AFA on your Surveys order form.



Neuropathology Program NP/NP1				
Program Code Challenges/Shipment				
	NP/NP1			
Neuropathology online case review	ı	8		

The Neuropathology program helps anatomic pathologists, neuropathologists, and trainees assess and improve their diagnostic skills and learn about new developments in neuropathology. Each shipment of this educational program includes eight cases that cover the spectrum of neoplastic and nonneoplastic disorders affecting the central and peripheral nervous systems, including infectious, degenerative, developmental, demyelinating, traumatic, toxic-metabolic, vascular, and neuromuscular diseases. In addition, each mailing will include a mini-symposium that focuses on a specific problem area in neuropathology, which relates to four of the eight cases.

- · NP Online activity providing eight cases and a minisymposium; reporting with CME/SAM credit is available for one pathologist; for each additional pathologist, order NP1
- Includes option to download program content
- NP1 Reporting option with CME/SAM credit for each additional pathologist (within the same institution); must order in conjunction with Survey NP
- · Earn a maximum of 10 CME/SAM credits (AMA) PRA Category 1 Credits[™]) per pathologist
- · This activity meets the ABP MOC Part IV Practice Performance Assessment requirements
- Powered by DigitalScope technology
- · Two online activities per year





Cytopathology

Analytes/procedures in bold type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Glass Slide Gynecologic Cytopathology PT Program with Glass Slide PAP Education PAP PT

Slide Type		Program Code					es Per Year
	PAPCPT	PAPKPT	PAPMPT	PAPLPT	PAPJPT	Proficiency Testing	Education
Conventional							
SurePath							
ThinPrep						10	10
Individual Participant Response Form	APAPCPT	APAPKPT	APAPMPT	APAPLPT	APAPJPT		

Ordering Information

You will receive one shipment for proficiency testing (10 slides) and two additional shipments for your education (five slides each).

Follow these steps to order your PAP Proficiency Testing and PAP Education:

- 1. Choose the following:
 - a. Slide Type program code (refer to table above)
 - b. PAP Education series shipment dates (choose one)
 - Series 1
 - O A mailing ships February
 - OB mailing ships August
 - Series 2
 - A mailing ships May
 - o B mailing ships November
 - c. Add the PAP Education series number after the slide type program code (eg, PAPCPT1, PAPCPT2).
- 2. Order one Individual Participant Response Form code for each participating pathologist/cytotechnologist. Also include the PAP Education Series number after the program code (eg, APAPCPT1).
- 3. Select primary testing session option with two alternative date options using the Gynecologic Cytology Proficiency Testing Order Details Form.
- 4. Order PPTENR only if you are a laboratory possessing a CLIA license to perform gynecologic cytology where all personnel are performing proficiency testing at another CLIA location.

Additional Information

- Participants can receive laboratory reference interpretations and performance for the PAP Education slides within 20 minutes by fax.
- The PAP Education component meets the CAP Laboratory Accreditation Program requirement for participation in a peer educational program.

- Ten glass slides for proficiency testing and ten glass slides for education
- APAPCPT/APAPKPT/
 APAPMPT/APAPLPT/
 APAPJPT Reporting option
 with CME/CE credit for each
 additional pathologist/
 cytotechnologist (within the
 same institution); must order
 in conjunction with Survey
 PAPCPT/PAPKPT/PAPMPT/
 PAPLPT/PAPJPT
- Earn a maximum of 8 CME credits (AMA PRA Category 1 Credits™) per pathologist and a maximum of 8 CE credits per cytotechnologist for completing all challenges
- This activity meets the ABP MOC Part IV Practice Performance Assessment requirements
- Three shipments per year; one shipment for proficiency testing (10 slides) and two shipments for education (five slides each)



Cytopathology Glass Slide Education Program PAPCE, PAPJE, PAPKE, PAPLE, PAPME Series 1 or 2

Slide Type	Program Code				Education Challenges Per Year	
	PAPCE	PAPKE	PAPME	PAPLE	PAPJE	
Conventional	ı				ı	
SurePath						
ThinPrep						10
Individual Participant Response Form	APAPCE	APAPKE	APAPME	APAPLE	APAPJE	10

Ordering Information

Follow these steps to order your PAP Education:

- 1. Choose the following:
 - a. Slide Type program code (refer to table above)
 - b. PAP Education series shipment dates (choose one)
 - Series 1
 - O A mailing ships February
 - o B mailing ships August
 - Series 2
 - O A mailing ships May
 - OB mailing ships November
 - c. Add the PAP Education series number after the slide type program code (eg, PAPCE1, PAPCE2)
- 2. Order one Individual Participant Response Form code for each participating pathologist/cytotechnologist. Also include the PAP Education series number after the program code (eg, APAPCE1).

Additional Information

- · Participants can receive laboratory reference interpretations and performance for the PAP Education slides within 20 minutes by fax.
- The PAP Education component meets the CAP Laboratory Accreditation Program requirement for participation in a peer educational program.

- · Ten glass slides for education
- APAPCE/APAPJE/APAPKE/ APAPLE/APAPME - Reporting option with CME/CE credit for each additional pathologist/cytotechnologist (within the same institution); must order in conjunction with Survey PAPCE/PAPJE/ PAPKE/PAPLE/PAPME
- · Earn a maximum of 8 CME credits (AMA PRA Category 1 Credits™) per pathologist and a maximum of 8 CE credits per cytotechnologist for completing all challenges
- · This activity meets the ABP MOC Part IV Practice Performance Assessment requirements
- Two shipments (five slides each)



Human Papillomavirus (High Risk) for Cytopathology CHPVD, CHPVM, CHPVK, CHPVJ

Analyte/Procedure		Program Code			Challenges/Shipment
	CHPVD	CHPVM	CHPVK	CHPVJ	
HPV	•	•			5
High-risk HPV genotyping (optional)					5

Additional Information

- Each laboratory should choose the Survey that best reflects the transport media
 received in its facility. For Survey CHPVJ, participants must provide results for all
 three media types. If your laboratory receives two types of media, order the Survey
 that is most appropriate for your specific laboratory (CHPVD, CHPVM, or CHPVK).
- For laboratories that perform HPV genotyping using ThinPrep PreservCyt Transport medium on site, Survey CHPVM and select samples of Survey CHPVJ provide an opportunity to report specific HPV genotypes.
- The CAP does not report genotyping responses to the CMS.

Program Information

- Five simulated cervical specimens
- CHPVD Digene® Specimen Transport Medium[™] (STM)
- CHPVM ThinPrep PreservCyt® Transport Medium
- CHPVK SurePath
 Preservative Fluid Transport
 Medium and corresponding
 vial of diluent
- CHPWJ Combination of Digene, ThinPrep PreservCyt, and SurePath transport mediums
- Three shipments per year

Practice Management Resources

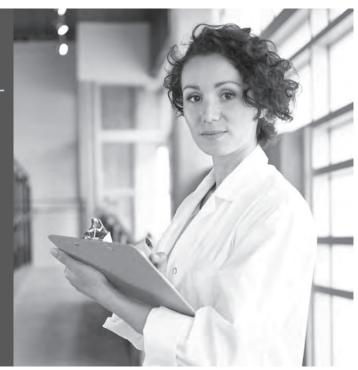
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- 2015 Practice Management Participant



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Touch Imprint/Crush Preparation TICP/TICP1				
Procedure	Program Code	Challenges/Shipment		
	TICP/TICP1			
Online slide and image program in rapid assessment case review	ı	4		

- The TICP Program is designed to familiarize surgical pathologists, cytopathologists, and cytotechnologists with the cytomorphologic features of pathologic processes and tumors in touch imprints and crush or scrape preparations. These specimens are prepared either for intraoperative consultation (frozen section) or rapid on-site evaluation (ROSE) of tissue biopsies for adequacy and/or interpretation. Participants will learn to make an immediate adequacy assessment, assign the process to a general category, and triage the specimen to appropriate ancillary studies. Participants will review digital whole slides of the TICP preparations (hematoxylin & eosin, modified Wright-Giemsa, and/or Papanicolaou stains), static images of the preparation and ancillary studies, and clinical history/radiographic findings to reach a diagnosis. Each case has a complete description of entities in the differential diagnosis along with a discussion of the correct interpretation.
- Participants will receive immediate feedback on interpretations, ancillary studies, and case-related adequate assessment.
- The 2018 cases will be comprised of specimens from the central nervous system and soft tissue.
- See system requirements on page 13.

- TICP Four online assessment challenges with clinical history; for each additional pathologist or cytotechnologist, order TICP1
- TICP1 Reporting option with CME/SAM/CE credit for each additional pathologist/ technologist (within the same institution); must order in conjunction with Survey TICP
- Earn a maximum of 10 CME/SAM credits (AMA PRA Category 1 Credits™) per pathologist and a maximum of 10 CE credits per cytotechnologist for completion of an entire year
- This activity meets the ABP MOC Part IV Practice Performance Assessment requirements
- Online whole slide images powered by DigitalScope technology
- Two online activities per year; your CAP shipping contact will be notified via email when the activity is available





Nongynecologic Cytopathology Education Program NGC/NGC1					
Procedure Program Code Challenges/Shipment					
	NGC/NGC1				
Nongynecologic cytopathology case review – glass slides		5			
Nongynecologic cytopathology case review — online	ı	5 per year			

- The Nongynecologic Cytopathology Education (NGC) program is an interlaboratory
 educational opportunity to assess participants' screening and interpretive skills.
 The NGC program is unsuitable for proficiency testing as these cases are chosen for
 their educational value. Cases may incorporate static online images that
 incorporate radiology and multiple aspects of pathology to enhance the
 interpretation.
- Participants can access laboratory reference interpretations and performance for the glass slides within 20 minutes by fax, providing rapid educational feedback, peer comparison, and additional review time.
- Additional online advanced education cases provide immediate feedback on interpretation selection, follow-up recommendations, and case-related educational questions.
- See system requirements on page 13.

- NGC Five glass slides; five online advanced education cases; one laboratory response form and two individual response forms
- NGC1 Reporting option with CME/CE credit for each additional pathologist/ cytotechnologist (within the same institution); must order in conjunction with Survey NGC
- Earn a maximum of 25 CME credits (AMA PRA Category 1 Credits™) per pathologist and a maximum of 25 CE credits per cytotechnologist for completing the glass slides and online cases
- This activity meets the ABP MOC Part IV Practice Performance Assessment requirements
- Online whole slide images powered by DigitalScope technology
- Four shipments per year



Digital Slide Program in Fine-Needle Aspiration FNA/FNA1					
Procedure Program Code Challenges/Shipment					
	FNA/FNA1				
Online program in fine-needle aspiration case review	•	5			

- This program focuses on FNA diagnostic dilemmas in practice. Online cases, which consist of whole slide images and static images, provide immediate feedback on interpretation selection, ancillary studies selection, and case-related educational questions.
- · Cases will focus on mediastinum and salivary topics.
- See system requirements on page 13.

- FNA Five online diagnostic challenges; for each additional pathologist or cytotechnologist, order FNA1
- FNA1 Reporting option with CME/CE credit for each additional pathologist/ cytotechnologist (within the same institution); must order in conjuction with Survey FNA
- Earn a maximum of 10 CME credits (AMA PRA Category 1 Credits[™]) per pathologist and a maximum of 10 CE credits per cytotechnologist
- · This activity meets the ABP MOC Part IV Practice Performance Assessment requirements
- Online whole slide images powered by DigitalScope technology
- Two online activities per year; your CAP shipping contact will be notified via email when the activity is available



Fine-Needle Aspiration Glass Slide FNAG/FNAG1				
Procedure	Program Code	Challenges/Shipment		
	FNAG/FNAG1			
Fine-needle aspiration glass slide case review	ı	5		

- The Fine-Needle Aspiration Glass Slide Education program is an interlaboratory educational opportunity to assess participants' screening and interpretive skills.
 FNAG cases may include more than one slide of varying stains and/or preparations used on fine-needle aspirations.
- Cases may include static online images that incorporate radiology and multiple aspects of pathology to support the interpretation.
- Participants can access laboratory reference interpretations and performance for the glass slides within 20 minutes by fax, providing rapid educational feedback, peer comparison, and additional review time.

- FNAG Five cases consisting of glass slides and selected online images, representing a variety of conditions; one laboratory response form and two individual response forms
- FNAG1 Reporting option with CME/CE credit for each additional pathologist/ cytotechnologist (within the same institution); must order in conjunction with Survey FNAG
- Earn a maximum of 10 CME credits (AMA PRA Category 1 Credits™) per pathologist/ resident and a maximum of 10 CE credits per cytotechnologist
- This activity meets the ABP MOC Part IV Practice Performance Assessment requirements
- Two shipments per year



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- 2,471 CAP inspections
- 23,196 laboratory sites using CAP proficiency testing

Discontinued Programs

Forensic Mitochondrial DNA Analysis (FIDM)

Forensic Sciences

Analytes/procedures in bold type are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Forensic Identity, Nuclear DNA Analysis FID					
Procedure	Program Code Challenges/Ship				
	FID				
Forensic nuclear DNA analysis	ı	3			

The American Society of Crime Laboratory Directors/Laboratory Accreditation Board Proficiency Review Committee (ASCLD/LAB PRC) has approved Survey FID.

Program Information

- Simulated forensic case work includes reference standards for all suspects and victims along with evidentiary material such as vaginal swabs, semen stains, and crime scene blood stains
- · Two shipments per year

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Let your peers, patients, and the public know you've earned the CAP accreditation certification mark.

Proudly display the mark. It distinguishes you as one of almost 8,000 laboratories worldwide that have attained CAP accreditation, the most respected and recognized laboratory accreditation in the world.

oontaining oix oadd otaaidd
illustrating gross and/or
microscopic slides and
questions related to
medicolegal decision
making; CME or CE credit is
available for one pathologist
or investigator. For each
additional pathologist/
investigator, order FR1
FR1 - Additional pathologist
or investigator (within the
same institution) reporting
came incritition) reporting

containing six case studies

Program Information

• FR - Online activity

- FRT Additional pathologist or investigator (within the same institution) reporting option with CME or CE credit; must order in conjunction with Survey FR
- Includes option to download program content
- Pathologists can earn a maximum of 12 CME credits (AMA PRA Category 1 Credits™) for completion of an entire year
- This activity meets the ABP MOC Part IV Practice Performance Assessment requirements
- Members of the American Board of Medicolegal Death Investigators, analysts, and technologists can earn a maximum of 12 CE credits for completion of an entire year
- Two online activities per year



Forensic Pathology FR/FR1 Procedure Program Code Challenges/Shipment FR/FR1 Forensic pathology cases ■ 6

Additional Information

- Cases may include or reflect anthropologic materials, ballistics, dental identification, DNA identification, environmental pathology, forensic evidence, injury pattern, medicolegal issues, toxicology, and trace evidence.
- FR/FR1 is for hospital-based pathologists, forensic pathologists, residents, fellows, and medical examiners/coroners. This educational program is also designed for investigators, analysts, and technicians/technologists.

Vitreous Fluid, Postmortem VF				
Analyte	Program Code	Challenges/Shipment		
	VF			
Acetone		3		
Chloride	I	3		
Creatinine	I	3		
Ethanol	I	3		
Glucose	ı	3		
Potassium	I	3		
Sodium	I	3		
Vitreous urea nitrogen		3		

Program Information

- Three 5.0-mL synthetic vitreous fluid specimens
- For forensic and other toxicology laboratories that perform quantitative analysis of vitreous fluid
- Conventional and International System of Units (SI) reporting offered
- Two shipments per year

Find a practical guide to toxicology laboratory operations with this resource

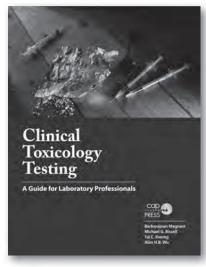
Clinical Toxicology Testing A Guide for Laboratory Professionals

Complex issues face the laboratory director or pathologist who offers toxicology services. This thorough reference book will guide both experienced physicians and those in training through the pharmacological principles, testing menus, and methodologies for toxicology testing.

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Item number: PUB220 Softcover; 304 pages; 2012

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Forensic Toxicology, Criminalistics FTC				
Analyte	Program Code	Challenges/Shipment		
	FTC			
See drug listing below	ı	4		

The American Society of Crime Laboratory Directors/Laboratory Accreditation Board Proficiency Review Committee (ASCLD/LAB PRC) has approved Survey FTC.

Program Information

- Three 20.0-mL whole blood specimens and one 20.0-mL synthetic urine specimen
- · For crime and hospital laboratories that have forensic toxicology divisions performing qualitative and quantitative analysis of drugs in whole blood specimens along with a urine qualitative challenge
- Two shipments per year

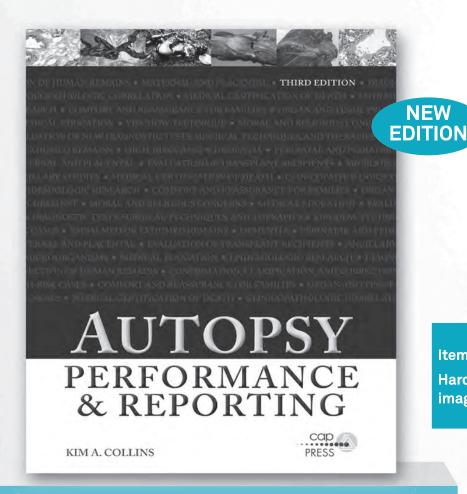
FTC Program Drug Listing

Challenges will include a mix of drugs from the list below.

6-acetylmorphine (6-AM)	Ecgonine ethyl ester	Norfluoxetine
7-aminoclonazepam	Ecgonine methyl ester	Norketamine
7-aminoflunitrazepam	Ephedrine	Norpropoxyphene
Acetaminophen	Fentanyl*	Norsertraline
Alpha-hydroxyalprazolam	Fluoxetine	Nortriptyline
Alprazolam	Flurazepam*	Oxazepam
Amitriptyline	Gamma-hydroxybutyrate (GHB)	Oxycodone
Amphetamine	Hydrocodone	Oxymorphone
Benzoylecgonine	Hydromorphone	Paroxetine
Brompheniramine	Imipramine	Phencyclidine
Butalbital	Ketamine	Phenethylamine
Carisoprodol	Lorazepam	Phenobarbital
Chlorpheniramine	Lysergic acid diethylamide (LSD)	Phentermine
Clonazepam	Meperidine*	Phenytoin
Cocaethylene	Meprobamate	Propoxyphene
Cocaine	Methadone	Pseudoephedrine
Codeine	Methadone metabolite (EDDP)	Salicylate
Cyclobenzaprine*	Methamphetamine	Secobarbital
Delta-9-THC	Methylenedioxyamphetamine (MDA)	Sertraline
Delta-9-THC-COOH	Methylenedioxymethamphetamine	Temazepam
Desipramine	(MDMA)	Tramadol*
Desmethylcyclobenzaprine	Morphine*	Trazodone
Dextromethorphan	N-desmethyltramadol	Zolpidem
Diazepam	Nordiazepam	
Diphenhydramine	Nordoxepin	
Doxepin	Norfentanyl	*and/or metabolite(s)

Refer to Section 9, Toxicology, for a more comprehensive selection of toxicology offerings.

Take a modern approach to autopsy pathology



Item number: PUB126

Hardcover; 472 pages; 1,000+ color images and tables; 2017

With more than 1,000 high-quality color images, the new edition of *Autopsy Performance & Reporting* is completely updated to include:

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22 Analyte/Procedure Index



Our programs are supported by 500 experts in laboratory medicine.

These experts spend countless hours monitoring testing trends to:

- Determine specimen specifications to challenge participants.
- Keep our offerings contemporary with new analytes and programs.
- Provide peer-reviewed CME, SAM, and CE education.

Analyte/Procedure Index

The following Analyte/Procedure Index is a comprehensive listing of analytes and corresponding CAP program options.

Analytes/procedures **in bold type** whose corresponding program codes are **bold** are regulated for proficiency testing by the Centers for Medicare & Medicaid Services (CMS).

Laboratories must perform five challenges three times per year (as noted by boldface) for analytes that are regulated by the CMS.

The **X** in the LAP ENR column denotes the CAP programs that can be used to fulfill the proficiency testing enrollment requirements for CAP-accredited laboratories. Refer to the program description in this catalog to determine compatibility with your specific methodologies.

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
1,5-anhydroglucitol		AG	1,5-Anhydroglucitol	71
1,25 dihydroxy		BMV1	Bone Markers and	86
vitamin D			Vitamins	
3-methoxytyramines		N/NX	Urine Chemistry, Special	69
4-hydroxytriazolam		DFC	Drug-Facilitated Crime	107
5-hydroxyindoleacetic acid, qualitative		N/NX	Urine Chemistry, Special	69
5-hydroxyindoleacetic acid, quantitative	Х	N/NX	Urine Chemistry, Special	69
6-acetylmorphine (6-AM)		DMPM	Drug Monitoring for Pain Management	106
		FTC	Forensic Toxicology, Criminalistics	104
		OFD	Oral Fluid for Drugs of Abuse	100
		Т	Toxicology	96
		UDC	Forensic Urine Drug Testing, Confirmatory	99
		UT	Urine Toxicology	96
7-aminoclonazepam		DFC	Drug-Facilitated Crime	107
		DMPM	Drug Monitoring for Pain Management	106
		FTC	Forensic Toxicology, Criminalistics	104
		Т	Toxicology	96
		UT	Urine Toxicology	96
7-aminoflunitrazepam		DFC	Drug-Facilitated Crime	107
		FTC	Forensic Toxicology, Criminalistics	104
		Т	Toxicology	96
		UT	Urine Toxicology	96
10q (PTEN) deletion		GLI	Glioma	253
11-dehydrothromboxane B2		TBX	11-Dehydrothromboxane B2	166
11-deoxycortisol		Y/YY	Ligand Assay, Special	84
17-hydroxycorticosteroids		N/NX	Urine Chemistry, Special	69
17-hydroxyprogesterone	Χ	Y/YY	Ligand Assay, Special	84
17-ketosteroids		N/NX	Urine Chemistry, Special	69

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
25-OH vitamin D	Х	ABVD	Accuracy-Based Vitamin D	85
		LN40	Vitamin D Cal Ver/Lin	127
	Х	VITD	25-OH Vitamin D	84
50:50 mixing study, APTT		CGE/CGEX	Coagulation, Extended	159
		CGS1	Coag Special, Series 1	160- 161
50:50 mixing study, PT		CGE/CGEX	Coagulation, Extended	159
		CGS1	Coag Special, Series 1	160- 161
ABO grouping	Х	J,J1	Transfusion Medicine	214
	Х	JAT	Transfusion Medicine, Automated	215
		JATE1	Transfusion Medicine, Automated, Educational	215
		TMCA	Transfusion Medicine, Competency Assessment	219
ABO subgroup typing		ABOSG	ABO Subgroup Typing	216
Acetaminophen	X	CZ, CZ2X,	Chemistry and TDM	56-
		CZX, Z		58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		FTC	Forensic Toxicology, Criminalistics	104
		LN3	TDM Cal Ver/Lin	119
	Х	SDS	Serum Drug Screen	101
		Т	Toxicology	96
		UDS, UDS6	Urine Drug Screen	98
		UT	Urine Toxicology	96
Acetone	X	AL1	Whole Blood Alcohol/ Ethylene Glycol/ Volatiles	101
	Х	AL2	Serum Alcohol/Ethylene Glycol/Volatiles	101
		SDS	Serum Drug Screen	101
		VF	Vitreous Fluid, Post- mortem	101
Acid-fast smear	Х	E	Mycobacteriology	183

Analyte/Procedure	LAP	Program	Description	Pg
	ENR			
Acid-fast smear (cont.)	Х	E1	Mycobacteriology, Ltd	183
Acid phosphatase	Χ	C3, C3X, CZ,	Chemistry and TDM	56-
		CZ2X, CZX		58
		CZQ	Quality Cross Check,	43
Activated clotting time	X	CT, CT1, CT2,	Chemistry and TDM ACT	162
Activated clotting time	^	CT3, CT5	ACI	102
		CTQ, CT1Q,	Quality Cross Check,	50
		CT2Q, CT3Q,	ACT	
		CT5Q		
		P0C14,	Competency Activated	54
		POC16	Clotting Time	
Activated partial	X	POC16	Basic Coagulation	158
thromboplastin time	^	ССБ	Dasic Coagulation	130
		CGE/CGEX	Coagulation, Extended	159
	Χ	CGL	Coagulation, Limited	158
		CGLQ	Quality Cross Check,	49
			Coagulation, Limited	
		CGS1	Coag Special, Series 1	160-
		CGS3	Coag Special, Series 3	161 160-
		CGSS	Coag Special, Series 3	161
		CGS4	Coag Special, Series 4	160-
				161
		DBGN	Anticoagulant	161
			Monitoring, Dabigatran	
		FNPX	Anticoagulant	161
			Monitoring, Fondaparinux	
		RVBN	Anticoagulant	161
		=	Monitoring, Rivaroxaban	
Activated protein C		CGE/CGEX	Coagulation, Extended	159
resistance				
		CGS2	Coag Special, Series 2	160-
Acylcarnitine		BGL	Biochemical Genetics	161 239
ADAMTS-13		CGS7	ADAMTS-13	160-
ABAWITO TO		0007	7.57.11110	161
Adenovirus		GIP	Gastrointestinal Panel	199
		ID2	Nucleic Acid Amp,	197
			Respiratory	
	X	IDR	Infectious Disease	198
		VI 92	Respiratory Panel	102
	X	VLS2 VR1	Viral Load Virology Culture	193
	X	VR1	Viral Antigen by DFA	191
	X	VR4	Viral Antigen by EIA and	192
	'		Latex	
Adenovirus 40/41		SP, SPN	Stool Pathogen	180
Adjustable micropipette		I	Instrumentation	129
Cal V/L				

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Adrenocorticotropic hormone (ACTH)	Х	TM/TMX	Tumor Markers	89
Alanine aminotrans- ferase (ALT/SGPT)	X	C1, C3, C3X, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		IFS	Interfering Substances	130
		LN2	Chemistry, Lipid,	118
			Enzyme Cal Ver/Lin	
		LN2BV	Chemistry, Lipid, Enzyme all Beckman except AU, Vitros Cal Ver/Lin	118
Albumin	X	C1, C3, C3X, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		FLD	Body Fluid	72
		FLDQ	Quality Cross Check, Body Fluid Chemistry	44
		IFS	Interfering Substances	130
		LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
		LN2BV	Chemistry, Lipid, Enzyme all Beckman except AU, Vitros Cal Ver/Lin	118
-		SPE	Protein Electrophoresis	76
Albumin, CSF	Х	M, OLI	CSF Chemistry and Oligoclonal Bands	74
Albumin, urine		ABU	Accuracy-Based Urine	111
		LN20	Urine Albumin	124
	Х	U	Urine Chemistry, General	68
Albumin: creatinine ratio		ABU	Accuracy-Based Urine	111
		LN20	Urine Albumin Cal Ver/ Lin	124
		U	Urine Chemistry, General	68
		UMC	Urine Albumin Creatinine	151
Alcohol, serum	Х	AL2	Serum Alcohol/Ethylene Glycol/Volatiles	101
		LN11	Serum Ethanol Cal Ver/ Lin	122
Alcohol, whole blood	Х	AL1	Whole Blood Alcohol/ Ethylene Glycol/ Volatiles	101
Aldolase		ADL	Aldolase	71
Aldosterone, serum	Χ	RAP	Renin and Aldosterone	89

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Aldastanaa unina	V	N/NX	Unio a Ob amiator Caracial	00
Aldosterone, urine Alkaline phosphatase	X	C1, C3, C3X,	Urine Chemistry, Special Chemistry and TDM	69 56-
(ALP)	^	CZ, CZ2X,	Chemistry and Tolvi	58
		CZQ	Quality Cross Check,	43
			Chemistry and TDM	
		FLD2	Body Fluid Chemistry 2	73
		IFS	Interfering Substances	130
		LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
		LN2BV	Chemistry, Lipid, Enzyme all Beckman except AU, Vitros Cal Ver/Lin	118
Allergens (specific)		SE	Diagnostic Allergy	207
Alpha-1 antitrypsin	Х	IG/IGX	Immunology, General	202
		LN7	Immunology Cal Ver/Lin	121
Alpha-1 antitrypsin genotyping	Х	AAT	Alpha-1 Antitrypsin Genotyping	239
Alpha-1 globulin		SPE	Protein Electrophoresis	76
Alpha-2 globulin		SPE	Protein Electrophoresis	76
Alpha-2-antiplasmin		CGE/CGEX	Coagulation, Extended	159
Alpha-2-macroglobulin		A2MG	Alpha-2-Macroglobulin	204
Alpha-fetoprotein (AFP), amniotic fluid	Х	FP/FPX	Maternal Screen	87
Alpha-fetoprotein (AFP), serum	Х	FP/FPX	Maternal Screen	87
	Χ	K/KK	Ligand Assay, General	82
		LN5	Ligand Assay Cal Ver/Lin	119-
		LNEC	1:	120
		LN5S	Ligand Assay, Siemens Cal Ver/Lin	119- 120
Alpha-hydroxyalprazolam		DFC	Drug-Facilitated Crime	107
		DMPM	Drug Monitoring for Pain Management	106
		FTC	Forensic Toxicology, Criminalistics	104
		T	Toxicology	96
		UDC	Forensic Urine Drug Testing, Confirmatory	99
		UT	Urine Toxicology	96
Alpha-thalassemia		HGM	Hemoglobinopathies, Molecular Methods	240
Alprazolam		DMPM	Drug Monitoring for Pain Management	106
		FTC	Forensic Toxicology, Criminalistics	104
		T	Toxicology	96
		OFD	Oral Fluid for Drugs of Abuse	100
		UT	Urine Toxicology	96

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Aluminum	Х	R	Trace Metals	78
		TMU	Trace Metals, Urine	103
		TMWB	Trace Metals, Whole	103
			Blood	
Amikacin	Х	CZ, CZ2X, CZX, Z	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		LN3	TDM Cal Ver/Lin	119
Amino acids, qualitative	Х	BGL	Biochemical Genetics	239
Amino acids, quantitative		BGL	Biochemical Genetics	239
Amitriptyline		DFC	Drug-Facilitated Crime	107
		FTC	Forensic Toxicology,	104
			Criminalistics	
		T	Toxicology	96
		UT	Urine Toxicology	96
	Х	ZT	TDM, Special	60
Ammonia		C3, C3X, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		LN32	Ammonia Cal Ver/Lin	126
Amniotic fluid leakage (nitrazine)		AFL	Amniotic Fluid Leakage	146
Amobarbital		DFC	Drug-Facilitated Crime	107
Amphetamine		DFC	Drug-Facilitated Crime	107
		DMPM	Drug Monitoring for Pain Management	106
		FTC	Forensic Toxicology, Criminalistics	104
		OFD	Oral Fluid for Drugs of Abuse	100
		T	Toxicology	96
		UDC	Forensic Urine Drug Testing, Confirmatory	99
		UDS, UDS6	Urine Drug Screen	98
		UT	Urine Toxicology	96
		UTCO	Urine Toxicology Carryover	131
Amphetamine group		DMPM	Drug Monitoring for Pain Management	106
		OFD	Oral Fluid for Drugs of Abuse	100
Delayed until 20	19	Т	Toxicology	96
2 Siay Sa aritir 20		TQP	Toxicology Quality Program	108
		UDS, UDS6	Urine Drug Screen	98
		UT	Urine Toxicology	96
Amylase	X	C1, C3, C3X, CZ, CZ2X, CZX	Chemistry and TDM	56- 58

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Amylase (cont.)		CZQ	Quality Cross Check,	43
			Chemistry and TDM	
		FLD	Body Fluid	72
		FLDQ	Quality Cross Check,	44
			Body Fluid Chemistry	
		IFS	Interfering Substances	130
		LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
		LN2BV	Chemistry, Lipid, Enzyme all Beckman	118
			except AU, Vitros Cal Ver/Lin	
Amylase, pancreatic	X	C1, C3, C3X, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
Amylase, urine		LN6	Urine Chemistry Cal Ver/Lin	120
	Х	U	Urine Chemistry, General	68
Anabasine		NTA	Nicotine and Tobacco Alkaloids	102
Analytical balance		I	Instrumentation	129
Anaplasma phagocytophilum		TTD	Antibody Detection- Tick-Transmitted Diseases	200
Anaplastic lymphoma kinase		PM6	Anaplastic Lymphoma Kinase IHC	268
Androstenedione	Х	Y/YY	Ligand Assay, Special	84
Angiotensin converting enzyme		ACE	Angiotensin Converting Enzyme	71
Anti-A titer		ABT, ABT1	Antibody Titer	217
Anti-B titer		ABT3	Antibody Titer	217
Anti-beta-2-glycoprotein		CGE/CGEX	Coagulation, Extended	159
Antibody detection	Х	J, JAT	Transfusion Medicine	214-
•				215
		JATE1	Transfusion Medicine, Automated, Educational	215
-	X	PS	Platelet Serology	219
	-	TMCA	Transfusion Medicine,	219
			Competency Assessment	
Antibody detection/	Х	MX1B, MX1C,	HLA Analysis, Class I	230-
identification (HLA)		MX1E, MXB, MXC		231
	Х	MX2B, MX2C,	HLA Analysis, Class II	230-
		MX2E, MXB, MXC		231
Antibody identification		ETME1	Expanded Transfusion Medicine Exercises	223

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Antibody identification (cont.)	Х	J, JAT	Transfusion Medicine	214- 215
		JATE1	Transfusion Medicine, Automated, Educational	215
		TMCA	Transfusion Medicine, Competency Assessment	219
Antibody screen (HLA)		MX1B, MX1C, MX1E, MXB, MXC	HLA Analysis, Class I	230– 231
		MX2B, MX2C, MX2E, MXB, MXC	HLA Analysis, Class II	230- 231
Anticardiolipin IgA, qualitative		ACL, APS	Antiphospholipid Antibody	205
Anticardiolipin IgA, quantitative		ACL, APS	Antiphospholipid Antibody	205
Anticardiolipin IgG, IgM, polyclonal; qualitative	Х	ACL, APS	Antiphospholipid Antibody	205
Anticardiolipin IgG, IgM, polyclonal; quantitative		ACL, APS	Antiphospholipid Antibody	205
Anti-CCP		CCP	Cyclic Citrullinated Peptide Antibody	206
Anticentromere antibody		S2	Immunology, Special	203
Antichromatin antibody		ACA	Antichromatin Antibody	204
Anti-CMV, total	Х	VM3	Viral Markers-Series 3	224
	Х	VR3	Infectious Disease Serology	200
Anti-CMV, IgG, IgM	Х	VR3	Infectious Disease Serology	200
Anti-D titer		ABT, ABT2	Antibody Titer	217
Anti-DNA (ds) antibody, qualitative	Х	S2, S4	Immunology, Special	203
Anti-DNA (ds) antibody, quantitative		S2, S4	Immunology, Special	203
Anti-DNA topoisomerase (Scl-70)		RDS	Rheumatic Disease Special Serologies	207
Antideamidated gliadin peptide antibody, IgA, IgG, qualitative	Х	CES, CESX	Celiac Serology	206
Antideamidated gliadin peptide antibody, IgA, IgG, quantitative		CES, CESX	Celiac Serology	206
Antideamidated gliadin peptide antibody screen, IgA, IgG		CES, CESX	Celiac Serology	206
Antideamidated gliadin peptide/tissue transglutaminase antibody screen, IgA, IgG		CES, CESX	Celiac Serology	206
Antiendomysial antibody IgA, qualitative		CES, CESX	Celiac Serology	206

Analyte/Procedure	LAP	Program	Description	Pg
Anatyte/Trocedure	ENR		Description	۱ ۵
Antiendomysial antibody		CES, CESX	Celiac Serology	206
IgA, quantitative				
Antiendomysial antibody		CES, CESX	Celiac Serology	206
IgG, qualitative				
Antiendomysial antibody		CES, CESX	Celiac Serology	206
IgG, quantitative				
Antifilamentous actin IgG		FCN	Antifilamentous Actin	204
antibody			Antibody	
Antifungal susceptibility	Х	F	Mycology and Aerobic	184
testing			Actinomycetes	
	Х	F1	Yeast	184
Antigen detection,		CDF2	Clostridium difficile	178
bacterial			Detection	
	Х	CDF5	Clostridium difficile	178
	.,	_	Detection	
	X	D	Bacteriology	170
	X	D4	Bacteriology, Limited	173
	X	D6	Rapid Group A Strep	175
	Х	D8	Group B Strep	176
	Х	D9	Rapid Group A Strep,	175
			Waived	
	Х	HC1	C. trachomatis by DFA	178
	Х	HC3	C. trachomatis by EIA	178
		LBAS	Legionella pneumophila	176
	X	MC1	Microbiology	173
			Combination	
	Х	MC2	Microbiology	173
	.,		Combination	
	Х	MC4	Urine Colony Count	174
	.,		Combination	
	Х	MC5	Throat Culture/Rapid	174
		500/	Strep Combination	
		POC4	POC Strep Screen	52
		CDAC	Competency	170
		SBAS	Streptococcus	176
	Х	VS	Vaginitis Screen	181
Antigen detection, viral	X	HC2		
Antigen detection, virat	X	VR2	HSV by DFA	192
	^	VKZ	Viral Antigen Detection by DFA	191
	Х	VR4	Viral Antigen Detection	192
		1114	by EIA and Latex	102
Antigliadin antibody IgA,	Х	CES, CESX	Celiac Serology	206
IgG, qualitative		,	21.00	
Antigliadin antibody IgA,		CES, CESX	Celiac Serology	206
IgG, quantitative		,		
Antiglomerular basement	Χ	S2	Immunology, Special	203
membrane, qualitative			0, -100	
Antiglomerular basement		S2	Immunology, Special	203
membrane, quantitative			55.	

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Anti-HAV, IgM	Χ	VM5	Viral Markers-Series 5	225
Anti-HAV, IgG	Х	VM1	Viral Markers-Series 1	224
Anti-HAV, total		VM1	Viral Markers-Series 1	224
Anti-HBc, IgM	Х	VM5	Viral Markers-Series 5	225
Anti-HBc, total	Х	VM1	Viral Markers-Series 1	224
Anti-HBe	Х	VM2	Viral Markers-Series 2	224
Anti-HBs, qualitative	Х	VM1	Viral Markers-Series 1	224
Anti-HBs, quantitative		VM1	Viral Markers-Series 1	224
Anti-HCV	Х	RHCVW	Anti-HCV, Rapid	225
			Methods, Waived	
	Х	VM1	Viral Markers-Series 1	224
Antihistidyl t-RNA		RDS	Rheumatic Disease	207
synthetase (Jo-1)			Special Serologies	
Antihistone antibody		AHT	Antihistone Antibody	204
Anti-HIV-1	Х	AHIV	Anti-HIV Rapid Methods	225
	Х	AHIVW	Anti-HIV Rapid Methods	225
	Х	VM1	Viral Markers-Series 1	224
Anti-HIV-2	Х	VM1	Viral Markers-Series 1	224
	Х	AHIV	Anti-HIV Rapid Methods	225
Anti-HIV-1/2	Х	AHIV	Anti-HIV Rapid Methods	225
	Х	AHIVW	Anti-HIV Rapid Methods	225
	Х	VM1	Viral Markers-Series 1	224
Anti-HIV-1/2, HIV-1 p24 antigen	X	VM6	Viral Markers-Series 6	225
Anti-HTLV-I/II		VM3	Viral Markers-Series 3	224
Anti-Jo-1(antihistidyl		RDS	Rheumatic Disease	207
t-RNA synthetase)			Special Serologies	
Anti-LKM		LKM	Liver-Kidney	207
			Microsomal Antibody	
Antimicrobial susceptibility testing	Х	D	Bacteriology	170
	Х	D2	Urine Cultures	172
	Х	D4	Bacteriology, Limited	173
	Х	D7	Throat, Urine Cultures	172
		MBT	Microbiology Bench	171
			Tools Competency	
	X	MC1	Microbiology	173
-			Combination with GC	
	Х	MC2	Microbiology Combination	173
	V	MOF		17/
	X	MC5	Throat Culture/Rapid Strep	174
Antimitochondrial antibody, qualitative	Х	S2	Immunology, Special	203
Antimitochondrial M2 antibody		Н	Antimitochondrial M2 Antibody	204
Anti-MPO		S2	Immunology, Special	203
Antimüllerian hormone		AMH	Antimüllerian Hormone	84
Antimycobacterial	Х	E	Mycobacteriology	183
susceptibility testing				

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Antimycobacterial susceptibility testing (cont.)		MTBR	Molecular MTB Detection and Resistance	183
Antineutrophil cytoplasmic antibody (ANCA)		S2	Immunology, Special	203
Antinuclear antibody (ANA)	Х	ANA, IL	Immunology	202
Antiparietal cell antibody		APC	Autoimmune Gastritis Markers	204
Antiphospholipid antibody		ACL	Antiphospholipid Antibody	205
		CGE/CGEX	Coagulation, Extended	159
Antiphosphatidylserine antibodies (IgG, IgM, and IgA)		APS	Antiphosphatidylserine Antibodies	205
Anti-PR3		S2	Immunology, Special	203
Antiribosomal P antibody		ARP	Antiribosomal P Antibody	205
Anti-RNP antibody, qualitative	Х	S2	Immunology, Special	203
Anti-RNP antibody, quantitative		S2	Immunology, Special	203
Anti-Saccharomyces cerevisiae antibody		ASC	Anti-Saccharomyces cerevisiae Antibody	205
Anti-Scl-70 (anti-DNA topoisomerase)		RDS	Rheumatic Disease Special Serologies	207
Anti-Sm antibody, qualitative	Х	S2	Immunology, Special	203
Anti-Sm antibody, quantitative		S2	Immunology, Special	203
Anti-Sm/RNP antibody, qualitative	Х	S2	Immunology, Special	203
Anti-Sm/RNP antibody, quantitative		S2	Immunology, Special	203
Antismooth muscle antibody	Х	S2	Immunology, Special	203
Antisperm antibody IgG	Х	ASA	Semen Analysis	154
Anti-SSA antibody, qualitative	Х	S2	Immunology, Special	203
Anti-SSA antibody, quantitative		S2	Immunology, Special	203
Anti-SSB antibody, qualitative	Х	S2	Immunology, Special	203
Anti-SSB antibody, quantitative		S2	Immunology, Special	203
Anti-SSA/SSB antibody, qualitative	Х	S2	Immunology, Special	203
Anti-SSA/SSB antibody, quantitative		S2	Immunology, Special	203
Antistreptolysin O (ASO)	Х	ASO, IL	Immunology	202

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Antithrombin (activity,		CGE/CGEX	Coagulation, Extended	159
Ag)				
		CGS2	Coag Special, Series 2	160- 161
		LN35	Thrombophilia Cal Ver/ Lin	127
Antithyroglobulin antibody, qualitative	Х	S2, S4	Immunology, Special	203
Antithyroglobulin antibody, quantitative		S2, S4	Immunology, Special	203
Antithyroid microsomal,	Х	S2, S4	Immunology, Special	203
qualitative				
Antithyroid microsomal, quantitative		\$2,\$4	Immunology, Special	203
Antithyroid peroxidase, qualitative	Х	S2, S4	Immunology, Special	203
Antithyroid peroxidase, quantitative		S2, S4	Immunology, Special	203
Antitissue transglutaminase antibody IgA, qualitative	Х	CES, CESX	Celiac Serology	206
Antitissue transglutaminase antibody IgA, quantitative		CES, CESX	Celiac Serology	206
Antitissue transglutaminase antibody IgG, qualitative		CES, CESX	Celiac Serology	206
Antitissue transglutaminase antibody IgG, quantitative		CES, CESX	Celiac Serology	206
Anti-Trypanosoma cruzi		VM4	Viral Markers-Series 4	224
Apixaban		APXBN	Anticoagulant Monitoring, Apixaban	161
Apolipoprotein A1	Х	ABL	Accuracy-Based Lipids	110
	Х	C3, C3X, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
Apolipoprotein B	Х	ABL	Accuracy-Based Lipids	110
	Х	C3, C3X, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
Apolipoprotein E (APOE) genotyping	Х	APOE	Apolipoprotein E (APOE) genotyping	239
Aripiprazole		T	Toxicology	96
		UT	Urine Toxicology	96
Arsenic, urine		TMU	Trace Metals, Urine	103
Arsenic, whole blood		TMWB	Trace Metals, Whole Blood	103

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Arthropod identification		тмо	Ticks, Mites, and Other	189
Aspartate aminotransferase (AST/SGOT)	X	C1, C3, C3X, CZ, CZ2X, CZX	Arthropods Chemistry and TDM	56- 58
		CZQ	Quality Cross Check,	43
			Chemistry and TDM	
		IFS	Interfering Substances	130
		LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
		LN2BV	Chemistry, Lipid, Enzyme all Beckman except AU, Vitros Cal Ver/Lin	118
Aspirin assay		PIA, PIAX	Drug-Specific Platelet Aggregation	165
Astrovirus		GIP	Gastrointestinal Panel	199
Atenolol		T	Toxicology	96
		UT	Urine Toxicology	96
Atropine		T	Toxicology	96
		UT	Urine Toxicology	96
Automated WBC differential	X	FH1-FH4, FH6, FH9, FH10, FH13, FH1P-FH4P, FH6P, FH9P, FH10P, FH13P	Hematology, Auto Diff	135
		FH3Q, FH4Q, FH6Q, FH9Q	Quality Cross Check, Automated Hematology Series	47
Autopsy pathology		AUP/AUP1	Autopsy Pathology	270
B-ALL		BALL	B-ALL Minimal Residual Disease	210
B-type natriuretic peptides	Х	BNP	B-Type Natriuretic Peptides, 2 Chall	61
	Х	BNP5	B-Type Natriuretic Peptides, 5 Chall	61
		BNPQ	Quality Cross Check, B-Type Natriuretic Peptides	43
		LN30	B-Type Natriuretic Peptides Cal Ver/Lin	125
	Х	PCARM, PCARMX	Plasma Cardiac Markers	65
		POC12	Competency Plasma Cardiac Markers	53
Babesia microti		TTD	Antibody Detection of Tick-Transmitted Diseases	200
Bacterial antigen detection		CDF2	Clostridium difficile Detection	178

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Bacterial antigen	Х	CDF5	Clostridium difficile	178
detection (cont.)			Detection	
	Х	D	Bacteriology	170
	Х	D4	Bacteriology, Limited	173
	Х	D6	Rapid Group A Strep	175
	Х	HC1	C. trachomatis by DFA	178
	Х	HC3	C. trachomatis by EIA	178
		LBAS	Legionella pneumophila	176
			Antigen Detection	
	Х	MC1	Microbiology	173
			Combination	
	Х	MC2	Microbiology	173
			Combination	
	Х	MC4	Urine Colony Count	174
			Combination	
	Х	MC5	Throat Culture/Rapid	174
			Strep Combination	
		POC4	POC Strep Screen	52
			Competency	
		SBAS	S. pneumoniae Antigen	176
			Detection	
	Х	VS	Vaginitis Screen	181
Bacterial detection in		BDP, BDPV	Bacterial Detection,	222
platelets			Platelets	
	Х	BDP5, BDPV5	Bacterial Detection,	222
			Platelets	
Bacterial identification	Х	D	Bacteriology	170
	Х	D1, D2, D3,	Throat, Urine, GC	172
		D7	Cultures	
	X	D4	Bacteriology, Limited	173
	Х	D8	Group B Strep	176
		DEX	Expanded Bacteriology	171
	Х	HC6/HC6X	C. trachomatis/GC by	194
			Nucleic Acid Amp	
	Х	HC7	C. trachomatis/GC DNA	194
			by NAA	
	X	IDR	Infectious Disease,	198
			Respiratory Panel	
	X	MC1	Microbiology	173
			Combination with GC	
	X	MC2	Microbiology	173
			Combination	
	X	MC4	Urine Colony Count	174
			Combination	
	X	MC5	Throat Culture/Rapid	174
			Strep	
		MBT	Microbiology Bench	171
			Tools Competency	
		MRS	Methicillin-Resistant	179
			Staphylococcus aureus	
			Screen	

,	LAP ENR	Program Code	Description	Pg
Bacterial identification (cont.)		MRS2M	MRSA Screen Molecular, 2 Challenge	179
	X	MRS5	Methicillin-Resistant Staphylococcus aureus Screen	179
	Χ	MRS5M	MRSA Screen, Molecular, 5 Challenge	179
Bacterial strain typing		BSTS	Bacterial Strain Typing- Staphylococcus	195
Bacterial vaginosis screen		BV	Bacterial Vaginosis	181
		VS2	Vaginitis Screen, Virtual Gram Stain	182
Barbiturate group		DMPM	Drug Monitoring for Pain Management	106
		SDS	Serum Drug Screen	101
		T	Toxicology	96
		UDS, UDS6	Urine Drug Screen	98
		UT	Urine Toxicology	96
BCR/ABL1 p190		MH02, MH03	Molecular Hematologic	254
BCR/ABLT P190		IVITIOZ, IVITIOS	Oncology	234
		MRD1	Minimal Residual Disease	254
BCR/ABL1 p210		MH02, MH03	Molecular Hematologic Oncology	254
		MRD	Minimal Residual Disease	254
Bence Jones protein		UBJP	Urinary Bence Jones Protein	76
Benzodiazepine group		DMPM	Drug Monitoring for Pain Management	106
		OFD	Oral Fluid for Drugs of Abuse	100
		SDS	Serum Drug Screen	101
		T	Toxicology	96
Delayed until 20	19	TQP	Toxicology Quality Program	108
		UDS, UDS6	Urine Drug Screen	98
		UT	Urine Toxicology	96
Benzoylecgonine		DFC	Drug-Facilitated Crime	107
, ,		DMPM	Drug Monitoring for Pain Management	106
		FTC	Forensic Toxicology, Criminalistics	104
		OFD	Oral Fluid for Drugs of Abuse	100
		T	Toxicology	96
		UDC	Forensic Urine Drug Testing, Confirmatory	99
		UDS, UDS6	Urine Drug Screen	98
		UT	Urine Toxicology	96

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Benzoylecgonine (cont.)		UTCO	Urine Toxicology	131
			Carryover	
Beta-2-glycoprotein I		ACL, APS	Antiphospholipid Antibody	205
Beta-2-microglobulin, serum	Х	TM/TMX	Tumor Markers	89
Beta-2-microglobulin, urine		CD	Cadmium	102
Beta-hydroxybutyrate	Х	KET	Ketones	64
Beta globulin		SPE	Serum Electrophoresis	76
Beta-thalassemia		HGM	Hemoglobinopathies, Molecular Methods	240
Bile crystal identification		BCR	Bile Crystals	147
Bilirubin, confirmatory urine		DSC	Dipstick Confirmatory	147
Bilirubin, direct	Х	C1, C3, C3X, C4, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
		LN2BV	Chemistry, Lipid, Enzyme all Beckman except AU, Vitros Cal Ver/Lin	118
	Х	NB, NB2	Neonatal Bilirubin	65
Bilirubin, total	Х	C1, C3, C3X, C4, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
-		FLD2	Body Fluid Chemistry 2	73
		IFS	Interfering Substances	130
		LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
		LN2BV	Chemistry, Lipid, Enzyme all Beckman except AU, Vitros Cal Ver/Lin	118
	Χ	NB, NB2	Neonatal Bilirubin	65
Bilirubin, urine	Χ	CMP, CMP1	Clinical Microscopy	144
		CMQ	Quality Cross Check, Urinalysis	48
		DSC	Dipstick Confirmatory	147
	Χ	HCC2	Waived Combination	66
		POC3	POC Urine Dipstick Competency	52
Bioavailable testosterone		DY	Ligand Assay, Special	84
Biochemical genetics		BGL, BGL1	Biochemical Genetics	239
Bioterrorism agents		LPX	Laboratory Preparedness Exercise	180

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
BK virus		ID1T	Nucleic Acid Amp, JC and BK	196
		VLS, VLS2	Viral Load	193
Blood cell identification	Х	BCP, BCP2	Blood Cell Identification	134
		EHE1	Expanded Virtual Peripheral Blood Smear	142
	X	FH1-FH4, FH6, FH9, FH10, FH13, FH1P-FH4P, FH6P, FH9P, FH10P, FH13P	Hematology, Auto Diff	135
	Χ	HEP	Basic Hematology	134
		VPBS	Virtual Peripheral Blood	141
			Smear	
		VBF	Virtual Body Fluid	146
Blood culture	Х	BCS	Blood Culture	176
		GNBC	Gram-Negative Blood Culture Panel	199
		GPBC	Gram-Positive Blood Culture Panel	199
Blood culture		BCS1	Blood Culture	177
Staphylococcus aureus			Staphylococcus aureus	
Blood parasite	Х	ВР	Blood Parasite	188
	Х	Р	Parasitology	187
Blood parasite, rapid		RMAL	Rapid Malaria	188
Bloom syndrome	Х	MGL4	Molecular Genetics	241
Bocavirus		IDR	Infectious Disease Respiratory Panel	198
Body fluid case studies		VBF	Virtual Body Fluid	146
Body fluid (cell count)		ABF1, ABF2, ABF3	Automated Body Fluid	146
Body fluid (cell count)	Х	HFC, HFCI	Hemocytometer Fluid Count	148- 149
Body fluid cell identification		CMP/CMP1	Clinical Microscopy	144
Body fluid (chemistry)		FLD, FLD2	Body Fluid	72- 73
Body fluid crystal identification		BFC	Crystals	147
Body fluid photographs		CMP, CMP1	Clinical Microscopy	144
Bone marrow cell		BMD	Bone Marrow Cell	138
differential			Differential	
Bone marrow cell identification		BMD	Bone Marrow Cell Differential	138
Bone specific alkaline		BMV2	Bone Markers and	86
phosphatase			Vitamins	
Bordetella holmesii	Х	IDR	Nucleic Acid Amp, Organisms	198

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Bordetella parapertussis		BOR	Bordetella pertussis/	177
			parapertussis,	
			Molecular	
		IDN, IDO	Nucleic Acid Amp,	196
			Organisms	
	Х	IDR	Infectious Disease	198
			Respiratory Panel	
Bordetella pertussis		BOR	Bordetella pertussis/	177
			parapertussis,	
			Molecular	
		IDN, IDO	Nucleic Acid Amp,	196
			Organisms	
	X	IDR	Infectious Disease	198
			Respiratory Panel	
Borrelia burgdorferi		TTD	Antibody Detection	200
			of Tick-Transmitted	
			Diseases	
BRAF	Х	BRAF	Mutation Testing	252
	Х	MTP	Multigene Tumor Panel	253
BRAF V600E		BRAFV	BRAF V600E	267
BRCA1/2	Х	MGL3	Molecular Genetics	241
BRCA1/2 sequencing	Х	BRCA	BRCA1/2 Sequencing	240
BRCA1/2 duplication/		BRCA	BRCA1/2 Sequencing	240
deletion analysis				
Brain tissue by FISH		CAN	Fluorescence In Situ Hybrid, Brain/Glioma Tissue	237
Brightfield in situ	Χ	ISH2	In Situ Hybridization	250
hybridization				
Brompheniramine		DFC	Drug-Facilitated Crime	107
		FTC	Forensic Toxicology,	104
			Criminalistics	
		Т	Toxicology	96
		UT	Urine Toxicology	96
Buprenorphine		DMPM	Drug Monitoring for Pain	106
			Management	
		OFD	Oral Fluid for Drugs of	100
			Abuse	
		Т	Toxicology	96
		UDC	Forensic Urine Drug	99
			Testing, Confirmatory	
		UDS, UDS6	Urine Drug Screen	98
		UT	Urine Toxicology	96
Bupropion		T	Toxicology	96
		UT	Urine Toxicology	96
Butalbital		DFC	Drug-Facilitated Crime	107
		DMPM	Drug Monitoring for Pain	106
			Management	
		FTC	Forensic Toxicology,	104
		-	Criminalistics	
		T	Toxicology	96

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Butalbital (cont.)		UDC	Forensic Urine Drug Testing, Confirmatory	99
		UT	Urine Toxicology	96
C. difficile		CDF2	Clostridium difficile Detection	178
	X	CDF5	Clostridium difficile Detection	178
	Х	D	Bacteriology, Antigen Detection	170
		GIP	Gastrointestinal Panel	199
		SP, SPN	Stool Pathogens-Rapid and Molecular	180
CA 15-3		LN34	Tumor Markers Cal Ver/ Lin	126
	Х	TM/TMX	Tumor Markers	89
CA 19-9		FLD	Body Fluid	72
		FLDQ	Quality Cross Check, Body Fluid Chemistry	44
		LN34	Tumor Markers Cal Ver/ Lin	126
	Х	TM/TMX	Tumor Markers	89
CA 27.29	X	TM/TMX	Tumor Markers	89
CA 72-4	X	TM/TMX	Tumor Markers	89
CA 125		LN34	Tumor Markers Cal Ver/ Lin	126
	X	TM/TMX	Tumor Markers	89
Cadmium, urine	X	CD	Cadmium	102
Cadmium, whole blood	Х	CD	Cadmium	102
Caffeine	X	CZ2X, CZX, CZ, Z	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
Calcitonin	X	TM/TMX	Tumor Markers	89
Calcium		ABS	Accuracy-Based Testosterone and Estradiol	111
	Х	C1, C3, C3X, C4, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		FLD2	Body Fluid Chemistry 2	73
		IFS	Interfering Substances	130
		LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
		LN2BV	Chemistry, Lipid, Enzyme all Beckman except AU, Vitros Cal Ver/Lin	118
Calcium, urine		ABU	Accuracy-Based Urine	111
		LN6	Urine Chemistry Cal Ver/Lin	120

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Calcium, urine (cont.)	Χ	U	Urine Chemistry,	68
			General	
Calcium, ionized	X	AQ, AQ2, AQ3, AQ4	Aqueous Blood Gas	92
		AQQ, AQ2Q, AQ3Q, AQ4Q	Quality Cross Check, Critical Care Aqueous Blood Gas Series	46
	Х	C3, C3X, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		LN13C	Blood Gas Cal Ver/Lin	122- 123
		P0C10, P0C11	POC Competency Blood Gases	53
Calcofluor white		FSM	Fungal Smear	186
Campylobacter		CAMP	Campylobacter	177
Campylobacter		GIP	Gastrointestinal Panel	199
Canavan disease	V	MGL4	Molecular Genetics	241
Canavan disease Candida culture	X	F3	Candida Culture	185
	_	-		
Candida sp., DNA probe Cannabinoids	X	VS	Vaginitis Screen See Delta-9-THC-COOH	181
			and Delta-9-THC	136
Carbamazepine	X	CZ, CZ2X, CZX, Z	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		LN3	TDM Cal Ver/Lin	119
		Т	Toxicology	96
		UT	Urine Toxicology	96
Carbamazepine-10,11-epoxide		Т	Toxicology	96
		UT	Urine Toxicology	96
Carbamazepine, free	Х	CZ, CZ2X, CZX, Z	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
Carboxyhemoglobin	Х	SO	Blood Oximetry	94
		SOQ	Quality Cross Check, Blood Oximetry	46
Carisoprodol		DFC	Drug-Facilitated Crime	107
		DMPM	Drug Monitoring for Pain Management	106
		FTC	Forensic Toxicology, Criminalistics	104
		Т	Toxicology	96
		UT	Urine Toxicology	96
Carnitine	Х	BGL1	Biochemical Genetics	239
Casts, urine, semiquantitative		UAA, UAA1	Automated Urinalysis	147

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
CD1a		RFAV1	Rare Flow Antigen	212
			Validation CD1a	
CD3	X	FL, FL1	Lymphocyte Subset	209
			Immunophenotyping	
		LN22	Flow Cytometry Cal	124
		COD	Ver/Lin	221
00/	\ \	SCP	Stem Cell Processing	
CD4	X	FL, FL1	Lymphocyte Subset	209
		LN22	Immunophenotyping Flow Cytometry Cal	124
		LINZZ	Ver/Lin	124
CD8	X	FL, FL1	Lymphocyte Subset	209
020	^	,	Immunophenotyping	200
		LN22	Flow Cytometry Cal	124
			Ver/Lin	
CD20		PM3	Immunohistochemistry	268
CD34		CBT	Cord Blood Testing	221
	Х	FL4	Flow Cytometry CD34+	209
		SCP	Stem Cell Processing	221
CD45		CBT	Cord Blood Testing	221
	Х	FL, FL1	Lymphocyte Subset	209
			Immunophenotyping	
		FL4	Flow Cytometry CD34+	209
		SCP	Stem Cell Processing	221
CD49d		ZAP70	ZAP-70 Analysis by Flow	212
			Cytometry	
CD103		RFAV2	Rare Flow Antigen	212
			Validation, CD103	
CD117 (c-kit)		PM1	Immunohistochemistry	268
CEA		FLD	Body Fluid	72
		FLDQ	Quality Cross Check,	44
	\ \ \	14 1414 140	Body Fluid Chemistry	
	X	K, KK, K2	Ligand Assay, General	82
		LN5	Ligand Assay Cal Ver/Lin	119- 120
		LN5S	Ligand Assay, Siemens	119-
		LINGO	Cal Ver/Lin	120
Cell free DNA		CFDNA	Cell Free DNA	252
OCIL 1100 DIVA		NIPT	Non-invasive Prenatal	87
			Testing	0,
Ceruloplasmin	Х	S2, S4	Immunology, Special	203
CFU-GM		SCP	Stem Cell Processing	221
CH50		CH50	Total Hemolytic	208
			Complement	
CH100		CH50	CH100	208
Chlamydia trachomatis	Χ	HC1	C. trachomatis by DFA	178
	Χ	HC3	C. trachomatis by EIA	178
	Х	HC6, HC6X	C. trachomatis/GC by	194
			Nucleic Acid Amp	
	Х	HC7	C. trachomatis/GC DNA	194
			by NAA	L

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Chlamydia trachomatis (cont.)		VR1	Virology Culture	191
Chlamydophila pneumoniae		IDN, IDO	Nucleic Acid Amp, Organisms	196
	Х	IDR	Infectious Disease, Respiratory Panel	198
Chlordiazepoxide		Т	Toxicology	96
		UT	Urine Toxicology	96
Chloride	Х	AQ, AQ2, AQ3, AQ4	Aqueous Blood Gas	92
		AQQ, AQ2Q, AQ3Q, AQ4Q	Quality Cross Check, Critical Care Aqueous Blood Gas Series	46
	X	C1, C3, C3X, C4, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		FLD2	Body Fluid Chemistry 2	73
		IFS	Interfering Substances	130
		LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
		LN2BV	Chemistry, Lipid, Enzyme all Beckman except AU, Vitros Cal Ver/Lin	118
		LN13C	Blood Gas Cal Ver/Lin	122- 123
		POC10, POC11	POC Competency Blood Gases	53
Chloride, sweat	Х	SW1, SW2, SW3, SW4	Sweat Analysis Series	79
Chloride, urine		LN6	Urine Chemistry Cal Ver/Lin	120
	Х	U	Urine Chemistry, General	68
Chloride, vitreous fluid		VF	Vitreous Fluid, Post- mortem	101
Chlorpheniramine		DFC	Drug-Facilitated Crime	107
		FTC	Forensic Toxicology, Criminalistics	104
		Т	Toxicology	96
		UT	Urine Toxicology	96
Cholesterol		ABL	Accuracy-Based Lipids	110
	Х	C1, C3, C3X, C4, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		FLD	Body Fluid	72
		FLDQ	Quality Cross Check, Body Fluid Chemistry	44

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Cholesterol (cont.)	X	LCW	Ltd Chem, Waived	64
Onotostorot (dont.)	+ ^	LN2	Chemistry, Lipid,	118
			Enzyme Cal Ver/Lin	
		LN2BV	Chemistry, Lipid,	118
			Enzyme all Beckman	
			except AU, Vitros Cal	
		_	Ver/Lin	
Chromium	X	R	Trace Metals	78
		TMWB	Trace Metals, Whole Blood	103
Chromium, urine		TMU	Trace Metals, Urine	103
Chromosomal abnormalities	X	CY, CYBK	Cytogenetics	236
Citalopram		DFC	Drug-Facilitated Crime	107
-		T	Toxicology	96
		UT	Urine Toxicology	96
Citrate		KSA	Kidney Stone Risk	69
			Assessment	
CK isoenzymes	X	CRTI	Cardiac Markers	62
CK-MB	X	CRT, CRTI	Cardiac Markers	62
(immunochemical)				
		IFS	Interfering Substances	130
		LN2	Chemistry, Lipid,	118
		LNODY	Enzyme Cal Ver/Lin	
		LN2BV	Chemistry, Lipid, Enzyme all Beckman	118
			except AU, Vitros Cal	
			Ver/Lin	
	X	PCARM, PCARMX	Plasma Cardiac Markers	65
		POC12	Competency Plasma	53
		10012	Cardiac Markers	55
CK2 (MB)		IFS	Interfering Substances	130
		LN2	Chemistry, Lipid,	118
			Enzyme Cal Ver/Lin	
		LN2BV	Chemistry, Lipid,	118
			Enzyme all Beckman	
			except AU, Vitros Cal Ver/Lin	
Clinical pathology	+	CPIP/CPIP1	Quality Management,	14
improvement program		OI II / OF IF I	Education	1-7
Clomipramine		T	Toxicology	96
		UT	Urine Toxicology	96
Clonazepam		DMPM	Drug Monitoring for Pain	106
			Management	
		FTC	Forensic Toxicology,	104
			Criminalistics	
		Т	Toxicology	96
		UT	Urine Toxicology	96
Clonidine		DFC	Drug-Facilitated Crime	107
Clostridium difficile		CDF2	Clostridium difficile	178
			Detection	

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Clostridium difficile (cont.)	Х	CDF5	Clostridium difficile Detection	178
	Х	D	Bacteriology-Antigen Detection	170
		GIP	Gastrointestinal Panel	199
		SP, SPN	Stool Pathogens-Rapid and Molecular	180
Clozapine		Т	Toxicology	96
		UT	Urine Toxicology	96
		ZE	Therapeutic Drug Monitoring, Extended	60
CMV		ID1	Nucleic Acid Amp, Viruses	196
		LN38	CMV Viral Load Cal Ver/ Lin	127
		VLS, VLS2	Viral Load	193
	X	VM3	Viral Markers-Series 3	224
	X	VR1	Virology Culture	191
	X	VR2	Viral Antigen Detection by DFA	191
	X	VR3	Infectious Disease Serology	200
CO ₂	X	C3, C3X, C4, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		IFS	Interfering Substances	130
		LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
		LN2BV	Chemistry, Lipid, Enzyme all Beckman except AU, Vitros Cal Ver/Lin	118
Cobalt		TMU	Trace Metals, Urine	103
		TMWB	Trace Metals, Whole Blood	103
Cocaethylene		FTC	Forensic Toxicology, Criminalistics	104
		Т	Toxicology	96
		UT	Urine Toxicology	96
Cocaine		DMPM	Drug Monitoring for Pain Management	106
		FTC	Forensic Toxicology, Criminalistics	104
		OFD	Oral Fluid for Drugs of Abuse	100
		Т	Toxicology	96
		UDS, UDS6	Urine Drug Screen	98
		UT	Urine Toxicology	96
Codeine		DFC	Drug-Facilitated Crime	107

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Codeine (cont.)		DMPM	Drug Monitoring for Pain	106
			Management	
		FTC	Forensic Toxicology,	104
		OFD	Criminalistics	400
		OFD	Oral Fluid for Drugs of Abuse	100
		Т	Toxicology	96
		UDC	Forensic Urine Drug	99
		ODC	Testing, Confirmatory	33
		UT	Urine Toxicology	96
Compatibility testing	Х	J, JAT	Transfusion Medicine	214-
,		,		215
		JATE1	Transfusion Medicine	215
			Automated, Educational	
		TMCA	Transfusion Medicine,	219
			Competency	
			Assessment	
Complement C3	Χ	IG/IGX	Immunology, General	202
		LN7	Immunology Cal Ver/Lin	121
Complement C4	Х	IG/IGX	Immunology, General	202
		LN7	Immunology Cal Ver/Lin	121
Complexed PSA	Χ	K/KK	Ligand Assay, General	82
Conductivity, sweat	Х	SW1, SW2, SW3, SW4	Sweat Analysis Series	79
Connexin 26	Χ	MGL3	Molecular Genetics	241
Copper	Χ	R	Trace Metals	78
Copper, urine		TMU	Trace Metals, Urine	103
Copper, whole blood		TMWB	Trace Metals, Whole Blood	103
Coproporphyrins	Χ	N/NX	Urine Chemistry, Special	69
Coronavirus		ID2	Nucleic Acid Amp,	197
			Respiratory	
		IDR	Infectious Disease, Respiratory Panel	198
Cortisol		ABS	Accuracy-Based	111
			Testosterone and	
			Estradiol	
	Χ	C1, C3, C3X, CZ, CZ2X,	Chemistry and TDM	56- 58
		CZX		
		CZQ	Quality Cross Check, Chemistry and TDM	43
	Χ	K/KK	Ligand Assay, General	82
		LN5	Ligand Assay Cal Ver/Lin	119-
				120
		LN5S	Ligand Assay, Siemens	119-
			Cal Ver/Lin	120
Cortisol, salivary		SALC	Salivary Cortisol	77_
Cortisol, urinary free	Χ	N/NX	Urine Chemistry, Special	69
Cotinine		NTA	Nicotine and Tobacco	102
			Alkaloids	L

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Cotinine (cont.)		Т	Toxicology	96
		UT	Urine Toxicology	96
C-peptide	Х	ING	Insulin, Gastrin,	86
' '			C-Peptide, PTH	
C-reactive protein (CRP)	Х	CRP, IL	Immunology	202
		LN12, LN12E	C-Reactive Protein Cal	122
			Ver/Lin	
C-reactive protein, high- sensitivity (hsCRP)	X	HSCRP	High-Sensitivity C-Reactive Protein	64
		LN21	High-Sensitivity C-Reactive Protein Cal	124
Creating kings (CK)	V	C1 C2 C2V	Ver/Lin	FC
Creatine kinase (CK)	X	C1, C3, C3X, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		IFS	Interfering Substances	130
		LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
		LN2BV	Chemistry, Lipid,	118
		LIVEDV	Enzyme all Beckman	110
			except AU, Vitros Cal	
			Ver/Lin	
Creatinine	Х	AQ2, AQ4	Aqueous Blood Gas	92
		AQ2Q, AQ4Q	Quality Cross Check, Critical Care Aqueous	46
	V	04 00 000	Blood Gas Series	FC
	X	C1, C3, C3X, C4, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		FLD	Body Fluid	72
		FLDQ	Quality Cross Check,	44
			Body Fluid Chemistry	
		IFS	Interfering Substances	130
		LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
		LN2BV	Chemistry, Lipid,	118
			Enzyme all Beckman except AU, Vitros Cal Ver/Lin	
		LN24	Creatinine Accuracy Cal Ver/Lin	125
		SC0	Serum Carryover	131
Creatinine, urine		ABU	Accuracy-Based Urine	111
	X	BU	Bone and Mineral, Urine	85
	Х	CD	Cadmium	102
		DAI	Urine Drug Adulterant/ Integrity Testing	98

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Creatinine, urine (cont.)		LN6	Urine Chemistry Cal Ver/Lin	120
		LN20	Urine Albumin Cal Ver/ Lin	124
	X	U	Urine Chemistry, General	68
		UDC	Forensic Urine Drug Testing, Confirmatory	99
	Х	UMC	Urine Albumin/ Creatinine	151
Creatinine, vitreous fluid		VF	Vitreous Fluid, Post- mortem	101
Creatinine, whole blood	Χ	WBCR	Whole Blood Creatinine	66
Crossmatching		EXM, EXM2	Electronic Crossmatch	215- 216
	Х	J, JAT	Transfusion Medicine	214- 215
	Х	MX1B, MX1C, MXB, MXC	HLA Analysis, Class I	230- 231
	Х	MX2B, MX2C,	HLA Analysis Class II	230-
		MXB, MXC		231
		TMCA	Transfusion Medicine, Competency Assessment	219
Cryptococcal antigen detection	Х	CRYP	Cryptococcal Antigen Detection	185
	Х	F	Mycology and Aerobic Actinomycetes	184
	Χ	F1	Yeast	184
Cryptococcus neoformans/gatti		IDME	Meningitis/Encephalitis Panel	198
Cryptosporidium		GIP	Gastrointestinal Panel	199
Cryptosporidium immunoassay, preserved specimen	X	P, P3, P4, P5	Parasitology	187
Crystals, urine (semiquantitative)		UAA	Automated Urinalysis	147
Crystal identification (bile)		BCR	Bile crystals	147
Crystal identification (body fluid)		BFC	Body Fluid Crystals	147
Crystal identification (body fluid, urine and bile)		BFC	Body Fluid Crystals	147
Crystal identification (urine)		URC	Urine Crystals	147
CSF antigen detection	Χ	D	Bacteriology	170
C-telopeptide (CTX)		BMV5	Bone Markers and Vitamin	86
		BU	Bone and Mineral, Urine	85
Cyclic citrullinated peptide antibody		CCP	Anti-cyclic Citrullinated Peptide Antibody	206

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Cyclobenzaprine		DFC	Drug-Facilitated Crime	107
		FTC	Forensic Toxicology,	104
			Criminalistics	
		Т	Toxicology	96
		UT	Urine Toxicology	96
Cyclospora cayatanensis		GIP	Gastrointestinal Panel	199
Cyclosporine	Х	CS	Immunosuppressive Drugs	59
		LN31	Immunosuppressive Drugs Cal Ver/Lin	126
CYP2C9		PGX	Pharmacogenetics	243
CYP2C19		PGX	Pharmacogenetics	243
CYP2D6		PGX	Pharmacogenetics	243
CYP3A4		PGX	Pharmacogenetics	243
CYP3A5		PGX	Pharmacogenetics	243
Cystatin C		CYS	Cystatin C	74
Cystic fibrosis	Х	MGL2, MGL5	Molecular Genetics	241
Cystine		KSA	Kidney Stone Risk Assessment	69
Cytogenomic microarray		CYCGH	Constitutional Microarray Analysis	238
		CYCMA	Cytogenomic Microarray Analysis for Oncologic Abnormality	238
Cytology proficiency testing			See Cytopathology GYN proficiency testing	136
Cytomegalovirus (CMV)		ID1	Nucleic Acid Amp, Viruses	196
		IDME	Meningitis/Encephalitis Panel	198
		LN38	CMV Viral Load Cal Ver/ Lin	127
		VLS, VLS2	Viral Load	193
	X	VM3	Viral Markers-Series 3	224
	X	VR1	Virology Culture	191
	X	VR2	Virology by DFA	191
	X	VR3	Infectious Disease Serology	200
Cytopathology GYN education		PAPCE1	PAP Edu, Conventional	273
		PAPJE1	PAP Edu, All Technologies	273
		PAPKE1	PAP Edu, SurePath	273
		PAPME1	PAP Edu, ThinPrep	273
Cytopathology GYN proficiency testing		PAPCPT	PAP PT, Conventional	272
		PAPJPT	PAP PT, Combination	272
		PAPKPT	PAP PT, SurePath	272
		PAPLPT	PAP PT, Combination	272
		PAPMPT	PAP PT, ThinPrep	272

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Cytopathology, nongynecologic		FNA/FNA1	Fine-Needle Aspiration- Online	277
		FNAG/FNAG1	Fine-Needle Aspiration- Glass	278
		NGC/NGC1	Nongynecologic Cytopath Edu Prgm	276
Cytopreparation differential		HFC	Hemocytometer Fluid Count	148
Dabigatran		DBGN	Anticoagulant Monitoring, Dabigatran	161
D-dimer, qualitative		CGDF	Coagulation, D-dimer/ FDP	158
		CGL	Coagulation, Limited	158
D-dimer, quantitative	Х	CGDF	Coagulation, D-dimer/ FDP	158
	Х	CGL	Coagulation, Limited	158
		CGLQ	Quality Cross Check, Coagulation, Limited	49
		LN42	D-dimer Cal Ver/Lin	128
	Х	PCARM, PCARMX	Plasma Cardiac Markers	65
		POC12	Competency Plasma Cardiac Markers	53
Delta-9-THC		FTC	Forensic Toxicology, Criminalistics	104
		OFD	Oral Fluid for Drugs of Abuse	100
		T	Toxicology	96
		UT	Urine Toxicology	96
Delta-9-THC-COOH		DFC	Drug-Facilitated Crime	107
		DMPM	Drug Monitoring for Pain Management	106
		FTC	Forensic Toxicology,	104
			Criminalistics	
		OFD	Oral Fluid for Drugs of Abuse	100
		Т	Toxicology	96
		UDC	Forensic Urine Drug Testing, Confirmatory	99
		UDS, UDS6	Urine Drug Screen	98
		UT	Urine Toxicology	96
		UTC0	Urine Toxicology Carryover	131
Deoxypyridinoline (DPD)		BU	Bone and Mineral, Urine	85
Dermatopathology		DPATH/ DPATH1	Online Digital Slide Program	259
Dermatophyte identification	Х	F	Mycology and Aerobic Actinomycetes	184
Desalkylflurazepam		T	Toxicology	96
		UT	Urine Toxicology	96
Desipramine		DFC	Drug-Facilitated Crime	107

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Desipramine (cont.)		FTC	Forensic Toxicology,	104
			Criminalistics	
		Т	Toxicology	96
		UT	Urine Toxicology	96
	Х	ZT	TDM, Special	60
Desmethylclomipramine		Т	Toxicology	96
		UT	Urine Toxicology	96
Desmethylcycloben-		FTC	Forensic Toxicology,	104
zaprine			Criminalistics	
		Т	Toxicology	96
		UT	Urine Toxicology	96
Desmethylsertraline		Т	Toxicology	96
		UT	Urine Toxicology	96
Dextromethorphan		DFC	Drug-Facilitated Crime	107
· · · · · · · · · · · · · · · · · · ·		FTC	Forensic Toxicology,	104
			Criminalistics	
		Т	Toxicology	96
		UT	Urine Toxicology	96
DHEA sulfate	X	Y/YY	Ligand Assay, Special	84
DIA (Dimeric inhibin A)	X	FP/FPX	Maternal Screen	87
Diazepam	''	DMPM	Drug Monitoring for Pain	106
2.020 pa			Management	
		FTC	Forensic Toxicology,	104
			Criminalistics	
		OFD	Oral Fluid for Drugs of	100
			Abuse	
		Т	Toxicology	96
		UT	Urine Toxicology	96
Differential, automated	X	FH1-FH4, FH6, FH9, FH10, FH13, FH1P-FH4P, FH6P, FH9P, FH10P, FH13P	Hematology, Auto Diff	135
		FH3Q, FH4Q, FH6Q, FH9Q	Quality Cross Check, Automated Hematology Series	47
Differential (fluid),		HFC, HFCI	Hemocytometer Fluid	148-
manual			Count	149
Differential (blood),		EHE1	Expanded Virtual	142
manual			Peripheral Blood Smear	
		VPBS	Virtual Peripheral Blood Smear	141
Differential (bone marrow), manual		BMD	Bone Marrow Cell Differential	138
Digital slide program in fine-needle aspiration, online		FNA/FNA1	Online Digital Slide Program	277
Digoxin	X	CZ, CZ2X, CZX, Z	Chemistry and TDM	56- 58

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Digoxin (cont.)		CZQ	Quality Cross Check, Chemistry and TDM	43
		LN3	TDM Cal Ver/Lin	119
Digoxin, free	X	CZ, CZ2X, CZX, Z	Chemistry and TDM	56-
		CZQ	Quality Cross Check,	43
Dibudus as dains		Т	Chemistry and TDM	06
Dihydrocodeine		•	Toxicology	96
Dilki		UT	Urine Toxicology	96
Diltiazem		T	Toxicology	96
D'I 1		UT	Urine Toxicology	96
Dilute prothrombin time		CGE/CGEX	Coagulation, Extended	159
Dilute Russell's viper venom time		CGE/CGEX	Coagulation, Extended	159
Dimeric inhibin A (DIA)	Х	FP, FPX	Maternal Screen	87
Diphenhydramine		DFC	Drug-Facilitated Crime	107
		FTC	Forensic Toxicology, Criminalistics	104
		T	Toxicology	96
		UT	Urine Toxicology	96
Diphenylhydantoin			See Phenytoin	
Direct antiglobulin testing	Х	DAT	Direct Antiglobulin Testing	218
		TMCAD	Transfusion Medicine, Competency Assessment	219
Direct bilirubin	X	C1, C3, C3X, C4, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
		LN2BV	Chemistry, Lipid, Enzyme all Beckman except AU, Vitros Cal Ver/Lin	118
	X	NB, NB2	Neonatal Bilirubin	65
Disease association/ drug risk		DADR1, DADR2	Disease Association/ Drug Risk	233
Disopyramide	X	CZ, CZ2X, CZX, Z	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
DMD/Becker	Х	MGL2	Molecular Genetics	241
DNA analysis	Х	DML	HLA Molecular Typing	231
		FID	Forensic Identity	280
		МНО	Molecular Oncology	254
	Х	PARF	Parentage/Relationship	227
DNA content/cell cycle analysis		FL, FL2	Flow Cytometry	209

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
DNA extraction and		MH05	Molecular Oncology	251,
amplification			Hematologic	254
DNA fingerprinting		IDN, IDO	Nucelic Acid Amp,	196
0.1.		,	Organisms	
DNA mismatch repair		HQMMR	HistoQIP Mismatch	264
			Repair IHC	
		MMR	DNA Mismatch Repair	267
DNA sequencing		SEC, SEC1	DNA Sequencing	242
Dopamine	Х	N/NX	Urine Chemistry, Special	69
Doxepin		DFC	Drug-Facilitated Crime	107
		FTC	Forensic Toxicology,	104
			Criminalistics	
		T	Toxicology	96
		UT	Urine Toxicology	96
Doxylamine		DFC	Drug-Facilitated Crime	107
		T	Toxicology	96
		UT	Urine Toxicology	96
DPYD		PGX3	Pharmacogenetics	243
Duloxetine		T	Toxicology	96
		UT	Urine Toxicology	96
Ecgonine ethyl ester		FTC	Forensic Toxicology,	104
			Criminalistics	
		T	Toxicology	96
		UT	Urine Toxicology	96
Ecgonine methyl ester		FTC	Forensic Toxicology,	104
			Criminalistics	
		T	Toxicology	96
		UT	Urine Toxicology	96
E. coli 0157		GIP	Gastrointestinal Panel	199
EGFR-Epidermal growth factor receptor	X	EGFR	Mutation Testing	252
	Х	MTP	Multigene Tumor Panel	253
eGFR		LN24	Creatinine Accuracy	125
			CalVer/Lin	
Electronic crossmatch		EXM, EXM2	Electronic Crossmatch	215-
				216
Electrophoresis	X	HG	Hemoglobinopathy	139
		LPE	Lipoprotein Electrophoresis	76
	Х	M, OLI	CSF Chemistry and Oligoclonal Bands	74
	Х	SPE	Protein Electrophoresis	76
		UBJP	Urinary Bence Jones	76
			Proteins	
Elution, antibody		ELU	Eluate	218
-		TMCAE	Eluate Competency	220
			Assessment	
Embryology		EMB	Embryology	155
Enteroaggregative E. coli (EAEC)		GIP	Gastrointestinal Panel	199

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Enteropathogenic <i>E. coli</i> (EPEC)		GIP	Gastrointestinal Panel	199
Enterotoxigenic <i>E. coli</i> (ETEC)		GIP	Gastrointestinal Panel	199
Enterovirus		ID1	Nucleic Acid Amp, Viruses	196
		IDME	Meningitis/Encephalitis Panel	198
	Х	IDR	Infectious Disease, Respiratory Panel	198
	Χ	VR1	Virology Culture	191
Eosinophils, urine		SCM2	Special Clinical Microscopy	150
Ephedrine		FTC	Forensic Toxicology, Criminalistics	104
		T	Toxicology	96
		UT	Urine Toxicology	96
Epidermal growth factor receptor (EGFR)	Х	EGFR	Mutation Testing	252
	Х	MTP	Multigene Tumor Panel	253
Epinephrine		N/NX	Urine Chemistry, Special	69
Epithelial cells, urine, semiquantitative		UAA1	Automated Urinalysis	147
Epstein-Barr virus (EBV)		ID1	Nucleic Acid Amp, Viruses	196
		ISH	In Situ Hybridization	250
		VLS, VLS2	Viral Load	193
		VR3	Antibody Detection- Infectious Disease Serology	200
ER, PgR by immunohistochemistry	X	PM2	ER, PgR by Immunohistochemistry	269
Erythrocyte sedimentation rate		ESR, ESR1, ESR2, ESR3	Erythrocyte Sedimentation Rate	134
Erythropoietin		EP0	Erythropoietin	86
Escherichia coli K1		IDME	Meningitis/Encephalitis Panel	198
Escherichia coli 0157		GIP	Gastrointestinal Panel	199
Estradiol		ABS	Accuracy-Based Testosterone and Estradiol	111
		LN8	Reproductive Endocrinology Cal Ver/ Lin	121
	Χ	Y/YY	Ligand Assay, Special	84
Estriol, unconjugated (uE3)	Χ	FP/FPX	Maternal Screen	87
	Χ	Y/YY	Ligand Assay, Special	84
Estrogen receptors by immunohistochemistry	X	PM2	ER, PgR by Immunohistochemistry	269

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Ethanol	Х	AL1	Whole Blood Alcohol/ Ethylene Glycol/ Volatiles	101
	X	AL2	Serum Alcohol/Ethylene Glycol/Volatiles	101
		LN11	Serum Ethanol Cal Ver/ Lin	122
Ethanol, urine		UDS, UDS6	Urine Drug Screen	98
Ethanol, vitreous fluid		VF	Vitreous Fluid, Post- mortem	101
ETEC (Enterotoxigenic <i>E. coli</i>)		GIP	Gastrointestinal Panel	199
Ethosuximide	Х	CZ, CZ2X, CZX, Z	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
Ethylene glycol		AL1	Whole Blood Alcohol/ Ethylene Glycol/ Volatiles	101
		AL2	Serum Alcohol/Ethylene Glycol/Volatiles	101
Ethyl glucuronide (EtG)		ETB	Ethanol Biomarkers	102
Ethyl sulfate (EtS)		ETB	Ethanol Biomarkers	102
Euglobulin Test		CGE/CGEX	Coagulation, Extended	159
Everolimus		EV	Everolimus	59
Factor II		CGE/CGEX	Coagulation, Extended	159
Factor II mutation	Х	TPM	Thrombophilia Mutations	244
	Х	MGL1	Molecular Genetics	241
Factor V		CGE/CGEX	Coagulation, Extended	159
Factor V Leiden mutation	Х	MGL1	Molecular Genetics	241
	Х	TPM	Thrombophilia Mutations	244
Factor VII		CGE/CGEX	Coagulation, Extended	159
Factor VIII	Х	CGE/CGEX	Coagulation, Extended	159
	Х	CGS3	Coag Special, Series 3	160- 161
Factor VIII inhibitor		CGE/CGEX	Coagulation, Extended	159
Factor IX		CGE/CGEX	Coagulation, Extended	159
Factor X		CGE/CGEX	Coagulation, Extended	159
Factor XI		CGE/CGEX	Coagulation, Extended	159
Factor XII		CGE/CGEX	Coagulation, Extended	159
Factor XIII		CGE/CGEX	Coagulation, Extended	159
Familial dysautonomia	Х	MGL4	Molecular Genetics	241
Fanconi anemia,	Х	MGL4	Molecular Genetics	241
complementation grp. C				
Fecal fat, qualitative		FCFS	Fecal Fat	75
Fecal lactoferrin		FLAC	Fecal Lactoferrin	178
Fentanyl		DFC	Drug-Facilitated Crime	107
		DMPM	Drug Monitoring for Pain Management	106

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Fentanyl (cont.)		FTC	Forensic Toxicology,	104
			Criminalistics	
		Т	Toxicology	96
		UDS, UDS6	Urine Drug Screen	98
		UT	Urine Toxicology	96
Fern test (vaginal)	Х	CMMP	Clinical Microscopy, Misc	145
Ferritin	Х	C3, C3X, CZ,	Chemistry and TDM	56-
		CZ2X, CZX		58
		CZQ	Quality Cross Check, Chemistry and TDM	43
	Х	K, KK, K2	Ligand Assay, General	82
		LN5	Ligand Assay Cal Ver/Lin	119-
				120
		LN5S	Ligand Assay, Siemens	119-
			Cal Ver/Lin	120
Fetal fibronectin	Х	FF	Fetal Fibronectin	86
Fetal hemoglobin (gastric fluid)		APT	Fetal Hemoglobin	148
Fetal hemoglobin identification	Х	HG	Hemoglobinopathy	139
Fetal membrane rupture		ROM1	Rupture of Fetal Membrane	150
Fetal red cell quantitation	Х	HBF	Fetal Red Cell Detection	218
1		TMCAF	Transfusion Medicine, Competency Assessment	220
Fetal screen (Rosette testing)	Х	HBF	Fetal Red Cell Detection	218
		TMCAF	Transfusion Medicine, Competency Assessment	220
Fibrin monomer		CGE/CGEX	Coagulation, Extended	159
Fibrinogen	Х	CGL	Coagulation, Limited	158
		CGLQ	Quality Cross Check, Coagulation, Limited	49
		LN44	Fibrinogen, Cal Ver/Lin	128
Fibrinogen antigen		CGE/CGEX	Coagulation, Extended	159
Fibrinogen degradation		CGDF	Coagulation, D-dimer/	158
products, plasma		CCI	FDP	
		CGL	Coagulation, Limited	158
		CGLQ	Quality Cross Check, Coagulation, Limited	49
Fibrinogen degradation products, serum		CGDF	Coagulation, D-dimer/ FDP	158
		CGL	Coagulation, Limited	158
		CGLQ	Quality Cross Check, Coagulation, Limited	49
Fine-needle aspiration, digital slide program		FNA/FNA1	Online Digital Slide Program	277

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Fine-needle aspiration,		FNAG/FNAG1	Fine-Needle Aspiration	278
glass slides				
FISH for breast carcinoma hybridization and interpretation on site (HER2 gene amplification)	X	СҮН	Fluorescence In Situ Hybridization, Breast Cancer	237
FISH for brain/glioma		CAN	Fluorescence In Situ Hybridization, Brain/ Glioma Tissue	237
FISH for constitutional and hematologic disorders		CYF	Fluorescence In Situ Hybridization	236
FISH for lymphoma		CYL	Fluorescence In Situ Hybridization, Lymphoma	237
FISH for paraffin- embedded tissue		СҮН	Fluorescence In Situ Hybridization, Breast Cancer	237
		CYJ	Fluorescence In Situ Hybridization, Brain/ Glioma Tissue	237
		СҮК	Fluorescence In Situ Hybridization, Sarcoma Tissue or Pediatric Neoplasm	237
		CYL	Fluorescence In Situ Hybridization, Lymphoma	237
FISH for sarcoma		СҮК	Fluorescence In Situ Hybridization, Sarcoma Tissue or Pediatric Neoplasm	237
FISH for urothelial	Х	CYI	Fluorescence In Situ	236
carcinoma hybridization and interpretation on- site			Hybridization, Urothelial Carcinoma	
Flunitrazepam		T	Toxicology	96
		UT	Urine Toxicology	96
Fluorescent microscope check		I	Instrumentation	129
Fluoxetine		DFC	Drug-Facilitated Crime	107
		FTC	Forensic Toxicology, Criminalistics	104
		T	Toxicology	96
		UT	Urine Toxicology	96
Flurazepam		FTC	Forensic Toxicology, Criminalistics	104
		T	Toxicology	96
		UT	Urine Toxicology	96
Folate, RBC	Х	FOL	RBC Folate	88
Folate, serum	X	K, KK, K2	Ligand Assay, General	82

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Folate, serum (cont.)		LN5	Ligand Assay Cal Ver/Lin	119- 120
		LN5S	Ligand Assay, Siemens Cal Ver/Lin	119- 120
Follicle-stimulating hormone (FSH)		ABS	Accuracy-Based Testosterone, Estradiol	111
		LN8	Reproductive Endocrinology Cal Ver/ Lin	121
	Х	Y/YY	Ligand Assay, Special	84
Fondaparinux		FNPX	Anticoagulant Monitoring, Fondaparinux	161
Forensic identity (DNA)		FID	Forensic Identity	280
Forensic pathology		FR/FR1	Forensic Pathology	281
Forensic toxicology		FTC	Forensic Toxicology, Criminalistics	104
Fragile X	Х	MGL1	Molecular Genetics	241
Free beta hCG		FP1B	First Trimester Maternal Screening, Free Beta	87
Free testosterone	Х	DY	Ligand Assay, Special	84
Friedreich ataxia	Х	MGL2	Molecular Genetics	241
Fructosamine		FT	Fructosamine	75
Fungal culture		CBT	Cord Blood Testing	221
		SCP	Stem Cell Processing	221
Fungal serology		FSER	Fungal Serology	185
Fungus identification	X	F	Mycology and Aerobic Actinomycetes	184
	Х	F1	Yeast	184
	Х	F3	Candida culture	185
Gabapentin		DMPM	Drug Monitoring for Pain Management	106
		Т	Toxicology	96
		UT	Urine Toxicology	96
		ZE	Therapeutic Drug Monitoring, Extended	60
Galactomannan		FGAL	Galactomannan	185
Gamma globulin		M, OL1	CSF Chemistry	74
		SPE	Serum Electrophoresis	76
Gamma glutamyl transferase (GGT)	Х	C3, C3X, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		IFS	Interfering Substances	130
		LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
		LN2BV	Chemistry, Lipid, Enzyme all Beckman except AU, Vitros Cal Ver/Lin	118

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Gamma hydroxybutyrate		DFC	Drug-Facilitated Crime	107
		FTC	Forensic Toxicology,	104
			Criminalistics	
Gardnerella vaginalis, DNA Probe	Х	VS	Vaginitis Screen	181
Gastric occult blood		GOCB	Gastric Occult Blood	148
Gastric pH		GOCB	Gastric Occult Blood	148
Gastrin	Х	ING	Insulin, Gastrin, C-Peptide, PTH	86
Gaucher disease	Х	MGL4	Molecular Genetics	241
Genomic copy number		CYCGH	Constitutional	238
array			Microarray Analysis	
Gentamicin	Х	CZ, CZ2X, CZX, Z	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		LN3	TDM Cal Ver/Lin	119
Giardia		GIP	Gastrointestinal Panel	199
Giardia immunoassay, preserved specimen	Х	P, P3, P4, P5	Parasitology	187
Giemsa stain	Х	ВР	Blood Parasite	188
	Х	Р	Parasitology	187
Glioma by FISH		CYJ	Fluorescence In Situ	237
•			Hybridization, Brain/ Glioma Tissue	
Glucose	Х	AQ2, AQ4	Aqueous Blood Gas	92
		AQ2Q, AQ4Q	Quality Cross Check, Critical Care Aqueous Blood Gas Series	46
	X	C1, C3, C3X, C4, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		FLD	Body Fluid	72
		FLDQ	Quality Cross Check, Body Fluid Chemistry	44
		IFS	Interfering Substances	130
		LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
		LN2BV	Chemistry, Lipid, Enzyme all Beckman except AU, Vitros Cal Ver/Lin	118
		LN13C	Blood Gas Cal Ver/Lin	122- 123
Glucose, CSF	Х	M, OLI	CSF Chemistry and Oligoclonal Bands	74
Glucose, urine	Х	CMP, CMP1	Clinical Microscopy	144
		CMQ	Quality Cross Check, Urinalysis	48

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Glucose, urine (cont.)	Χ	HCC2	Waived Combination	66
		LN6	Urine Chemistry Cal	120
			Ver/Lin	
		POC3	POC Urine Dipstick	52
			Competency	
	Х	U	Urine Chemistry,	68
Clusoss vitrosus fluid		VF	General Vitreous Fluid, Post-	101
Glucose, vitreous fluid		VF	mortem	101
Glucose, whole blood	Х	HCC	Waived Combination	66
		HCC2	Waived Combination	66
	Х	LCW	Ltd Chem, Waived	64
		LN17	Whole Blood Glucose	123
			Cal Ver/Lin	
		POC2	POC Glucose	52
			Competency	
		POC7	POC/Waived Glucose	52
			and Hemoglobin	
		WBGQ	Competency	/0
		WBGQ	Quality Cross Check, Whole Blood Glucose	43
Glucose-6-phosphate		G6PDS	Glucose-6 Phosphate	75
dehydrogenase		GOI DO	Dehydrogenase	7.5
(qualitative and				
quantitative)				
Glutaraldehyde, urine		DAI	Urine Drug Adulterant/ Integrity Testing	98
Glycated serum albumin		GSA	Glycated Serum Albumin	64
Glycogen storage disease type 1A	Х	MGL4	Molecular Genetics	241
Glycohemoglobin	Х	GH2, GH5, GH5I	Hemoglobin A _{1c}	63
		GHQ	Quality Cross Check,	44
			Hemoglobin A _{1c}	
		LN15	Hemoglobin A _{1c} Cal Ver/Lin	123
Glycosaminoglycans (mucopolysaccharides)		BGL	Biochemical Genetics	239
Gram stain	Χ	D	Bacteriology	170
	Х	D2, D3, D7	Throat, Urine, GC Cultures	172
	Х	D4	Bacteriology, Ltd	173
	Χ	D5	Gram Stain	174
	Х	MC1	Microbiology	173
			Combination with GC	
	Х	MC2	Microbiology	173
		1405	Combination	4=:
	Х	MC5	Throat Culture/Rapid Strep	174
		VGS1	Virtual Gram Stain Basic	175

	Code		
	VGS2	Virtual Gram Stain	175
		Advanced	
	VS2		182
.,			
Х	D	Bacteriology	170
Χ	D4	Bacteriology, Limited	173
Χ	D6	<u> </u>	175
Х	D9	Rapid Group A Strep, Waived	175
Х	MC1	Microbiology Combination with GC	173
Х	MC2	Microbiology Combination	173
Χ	MC4	Urine Colony Count Combination	174
Χ	MC5	Throat Culture/Rapid Strep	174
	POC4	POC Strep Screen Competency	52
Χ	D8	Group B Strep	176
Χ	Y/YY	Ligand Assay, Special	84
		See Cytopathology GYN Proficiency Testing	
		See Cytopathology GYN Education	
	IDME	Meningitis/Encephalitis Panel	198
Χ	IG/IGX	Immunology, General	202
Χ	S2/S4	Immunology, Special	203
Χ	VM2	Viral Markers-Series 2	224
Χ	VM1	Viral Markers-Series 1	224
Χ	HBVL, HBVL5	Hepatitis Viral Load	193
Χ	NAT	Nucleic Acid Testing	226
Х	HCV2	Hepatitis Viral Load, Genotyping and Qualitative	193
	LN45	HCV Viral Load Cal Ver/ Lin	127
Χ	NAT	Nucleic Acid Testing	226
	ABL	Accuracy-Based Lipid	110
Х	C1, C3, C3X, C4, CZ, CZ3X, CZX	Chemistry and TDM	56- 58
]	CZQ	Quality Cross Check, Chemistry and TDM	43
Χ	LCW	Ltd Chem, Waived	64
	LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
	X X X X X X X X X X X X X X X X X X X	X D4 X D6 X D9 X MC1 X MC2 X MC4 X MC5 POC4 X D8 X Y/YY IDME X IG/IGX X S2/S4 X VM2 X VM1 X HBVL, HBVL5 X NAT X HCV2 LN45 X NAT ABL X C1, C3, C3X, C4, CZ, CZ3X, CZX CZX CZQ X LCW	Advanced VS2 Vaginitis Screen, Virtual Gram stain X D Bacteriology X D4 Bacteriology, Limited X D6 Rapid Group A Strep X D9 Rapid Group A Strep, Waived X MC1 Microbiology Combination with GC X MC2 Microbiology Combination X MC4 Urine Colony Count Combination X MC5 Throat Culture/Rapid Strep POC4 POC Strep Screen Competency X D8 Group B Strep X Y/YY Ligand Assay, Special See Cytopathology GYN Proficiency Testing See Cytopathology GYN Education IDME Meningitis/Encephalitis Panel X IG/IGX Immunology, General X S2/S4 Immunology, Special X VM2 Viral Markers-Series 2 X VM1 Viral Markers-Series 1 X HBVL, HBVL5 Hepatitis Viral Load X NAT Nucleic Acid Testing X HCV2 Hepatitis Viral Load, Genotyping and Qualitative LN45 HCV Viral Load Cal Ver/ Lin X NAT Nucleic Acid Testing ABL Accuracy-Based Lipid X C1, C3, C3X, C4, CZ, CZ3X, CZX CZQ Quality Cross Check, Chemistry and TDM X LCW Ltd Chem, Waived LN2 Chemistry, Lipid,

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
HDL cholesterol (cont.)		LN2BV	Chemistry, Lipid, Enzyme all Beckman except AU, Vitros Cal Ver/Lin	118
Helicobacter pylori	Χ	HPS	H. pylori Antigen, Stool	178
	Χ	S2, S4	H. pylori IgG Antibody	203
	Х	S5	H. pylori IgG Antibody	203
	Χ	VR3	H. pylori IgG Antibody	200
Hematocrit	Х	AQ, AQ2, AQ3, AQ4	Aqueous Blood Gas	92
		AQQ, AQ2Q, AQ3Q, AQ4Q	Quality Cross Check, Critical Care Aqueous Blood Gas Series	46
		CBT	Cord Blood Testing	221
	X	FH1-FH4, FH6, FH9, FH10, FH13, FH1P-FH4P, FH6P, FH9P, FH10P, FH13P	Hematology, Auto Diff	135
		FH3Q, FH4Q, FH6Q, FH9Q	Quality Cross Check, Automated Hematology Series	47
	Х	FH15	Centrifugal Hematology	135
	Х	HCC2	Waived Combination	66
	Х	HE, HEP	Basic Hematology	134
		POC10, POC11	POC Competency Blood Gases	53
		SCP	Stem Cell Processing	221
	Х	S0	Blood Oximetry	94
		SOQ	Quality Cross Check, Blood Oximetry	46
Hematology case studies		EHE1	Expanded Virtual Peripheral Blood Smear	142
		BMD	Bone Marrow Cell Differential	138
		VPBS	Virtual Periperal Blood Smear	141
Hematopathology online education		HPATH, HPATH1	Hematopathology Online Education	143
Hemochromatosis	Χ	MGL1	Molecular Genetics	241
Hemocytometer fluid count	Х	HFC, HFCI	Hemocytometer Fluid Count	148- 149
Hemoglobin		CBT	Cord Blood Testing	221
	X	FH1-FH4, FH6, FH9, FH10, FH13, FH1P-FH4P, FH6P, FH9P, FH10P, FH13P	Hematology, Auto Diff	135

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Hemoglobin (cont.)		FH3Q, FH4Q,	Quality Cross Check,	47
5 , ,		FH6Q, FH9Q	Automated Hematology Series	
	Х	FH15	Centrifugal Hematology	135
-	Х	HCC	Waived Combination	66
-	Х	HCC2	Waived Combination	66
-	X	HE, HEP	Basic Hematology	134
		LN9	Hematology Cal Ver/Lin	121
=		POC7	POC/Waived Glucose	52
		1 007	and Hemoglobin	02
			Competency	
		SCP	Stem Cell Processing	221
	Х	S0	Blood Oximetry	94
		SOQ	Quality Cross Check, Blood Oximetry	46
Hemoglobin electrophoresis	Х	HG	Hemoglobinopathy	139
Hemoglobin, estimated	Х	AQ, AQ2, AQ3, AQ4	Aqueous Blood Gas	92
		AQQ, AQ2Q,	Quality Cross Check,	46
		AQ3Q, AQ4Q	Critical Care Aqueous	
			Blood Gas Series	
		POC10,	POC Competency Blood	53
		POC11	Gases	
Hemoglobin, plasma		PHG	Plasma Hemoglobin	76
Hemoglobin, urine	Х	CMP, CMP1	Clinical Microscopy	144
		CMQ	Quality Cross Check, Urinalysis	48
	Х	HCC2	Waived Combination	66
		POC3	POC Urine Dipstick	52
			Competency	
Hemoglobin A _{1c}	Х	GH2, GH5, GH5I	Hemoglobin A _{1c}	63
		GHQ	Quality Cross Check,	44
			Hemoglobin A _{1c}	
		LN15	Hemoglobin A _{1c} Cal Ver/Lin	123
Hemoglobin A2 quantitation	Х	HG	Hemoglobinopathy	139
Hemoglobin F quantitation	Х	HG	Hemoglobinopathy	139
Hemoglobin S/C	Х	HGM	Hemoglobinopathies Genotyping	240
	Х	MGL2	Molecular Genetics	241
Hemolytic complement,		CH50	Total Hemolytic	208
total			Complement	
Hemosiderin, urine		SCM1	Special Clinical	150
		222	Microscopy	405
Heparin assay		CGS4	Coag Special, Series 4	160- 161
Heparin-induced		CGE/CGEX	Coagulation, Extended	159
thrombocytopenia				

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Heparin-induced		CGS5	Coag Special, HIT	160-
thrombocytopenia (cont.)				161
		CGS6	Coagulation Special	160- 161
		CGS8	Coag Special, HIT	160- 161
Heparin, low molecular weight		LN36	Heparin Cal Ver/Lin	127
Heparin, unfractionated		LN36	Heparin Cal/Ver Lin	127
Heparin/platelet Factor		CGS5	Coag Special, HIT	160-
IV				161
		CGS6	Coagulation Special	160-
				161
Hepatitis B virus	Х	HBVL, HBVL5	Hepatitis Viral Load	193
Hepatitis C virus	X	HCV2	Hepatitis Viral Load, Genotyping and Qualitative	193
		LN45	HCV Viral Load Cal Ver/ Lin	127
HER2, gastric		GHER2	Gastric HER2	269
HER2 gene amplification by ISH	Х	ISH2	In Situ Hybridization	250
HER2 gene amplification by FISH, hybridization and interpretation on site	X	СҮН	Fluorescence In Situ Hybridization, Breast Cancer	237
HER2 by immunohistochemistry	Х	HER2	HER2 by Immunohistochemistry	269
HER2 by molecular testing		MTP	Multigene Tumor Panel	253
Herpes simplex virus (HSV)	Х	HC2	HSV by DFA	192
	Χ	HC4	HSV Culture	192
		ID1	Nucleic Acid Amp, Viruses	196
		IDME	Meningitis/Encephalitis Panel	198
	Х	VR1	Virology Culture	191
	Х	VR2	Viral Antigen by DFA	191
	Х	VR3	Antibody Detection- Infectious Disease Serology	200
HHV6		ID1	Nucleic Acid Amp, Viruses	196
		IDME	Meningitis/Encephalitis Panel	198
		VLS2	Viral Load	193
HHV8		ID1	Nucleic Acid Amp, Viruses	196
High molecular weight kininogen		CGE/CGEX	Coagulation, Extended	159
High-sensitivity C-reactive protein	X	HSCRP	hsCRP	64

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
High-sensitivity C-reactive protein (cont.)		LN21	High-Sensitivity C-Reactive Protein Cal Ver/Lin	124
Histotechnology quality improvement		HQIP, HQIPBX, HQBX1, HQBX2, HQBX3, HQBX4, HQIHC, HQMMR, HQNSC	HistoQIP	263-266
HIV	Х	HIVG, HV2	HIV Viral Load	193
		LN39	HIV Viral Load Cal Ver/ Lin	127
	Х	NAT	Nucleic Acid Testing	226
HIV genotyping		HIVG	HIV Viral Genotyping	193
HIV-1 p24 antigen	Х	VM3	Viral Markers-Series 3	224
HIV-1 p24 antigen, Anti HIV 1/2	Х	VM6	Viral Markers-Series 6	225
HLA-A, -B, -C antibody identification	X	MX1B, MX1C, MX1E, MXB, MXC	HLA Analysis, Class I	230- 231
	X	MX2B, MX2C, MX2E, MXB, MXC	HLA Analysis, Class II	230- 231
HLA-(Class I/II) crossmatching	X	MX1B, MX1C, MX1E, MXB, MXC	HLA Analysis, Class I	230- 231
	X	MX2B, MX2C, MX2E, MXB, MXC	HLA Analysis, Class II	230- 231
HLA-(Class I/II) antibody screen		MX1B, MX1C, MX1E, MX2B, MX2C, MX2E, MXB, MXC	HLA Analysis, Class I/II	230- 231
HLA-B*1502		PGX2	Pharmacogenetics	243
HLA-B27 typing	Х	B27	HLA-B27 Typing	231
HLA-B*5701		PGX2	Pharmacogenetics	243
		DADR1	Disease Association, Drug Risk	233
HLA-B*57:01		DADR1	Disease Association, Drug Risk	233
HLA-B*58:01		DADR1	Disease Association, Drug Risk	233
HLA-DQA1*03/ DQB1*03:02		DADR2	Disease Association, Drug Risk	233
HLA-DQA1*05/DQB1*02		DADR2	Disease Association, Drug Risk	233
HLA molecular typing	Х	DML	HLA Molecular Typing	231
Homocysteine	Х	HMS	Homocysteine	64

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Homocysteine (cont.)		LN16	Homocysteine Cal Ver/ Lin	123
Homovanillic acid	Х	N/NX	Urine Chemistry, Special	69
HPV (cytopathology), high-risk	Х	CHPVD	Digene Specimen Transport Medium	274
	Χ	CHPVJ	Mixed Medium	274
	Х	CHPVK	SurePath Preservative Fluid Transport Medium	274
	Х	СНРУМ	ThinPrep PreservCyt Transport Medium	274
		HPV	Digene Hybrid Capture Technology Only	195
		ISH	In Situ Hybridization	250
HSV	Χ	HC2	HSV by DFA	192
	Х	HC4	HSV Culture	192
		ID1	Nucleic Acid Amp,	196
			Viruses	
	Х	VR1	Virology Culture	191
	Х	VR2	Viral Antigen by DFA	191
	X	VR3	Antibody Detection- Infectious Disease Serology	200
Human chorionic gonadotropin (hCG), serum	Х	C1, C3, C3X, C4, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
	Х	FP/FPX, FP1T	Maternal Screen	87
	Χ	HCG, IL	Immunology	202
	Х	K/KK	Ligand Assay, General	82
		LN5	Ligand Assay Cal Ver/Lin	119- 120
		LN5S	Ligand Assay, Siemens Cal Ver/Lin	119- 120
		LN8	Reproductive Endocrinology Cal Ver/ Lin	121
		SCO	Serum Carryover	131
Human chorionic gonadotropin (hCG), urine (qualitatitve)	X	CMP, CMP1	Clinical Microscopy	144
		CMQ	Quality Cross Check, Urinalysis	48
	Χ	HCC2	Waived Combination	66
		POC1	POC hCG Competency	52
		POC3	POC Urine Dipstick Competency	52
	Χ	UHCG	Urine HCG	150
Human epididymis protein 4		HUEP	Human Epididymis Protein 4	89

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Human herpesvirus 6		ID1	Nucleic Acid Amp, Viruses	196
		IDME	Meningitis/Encephalitis Panel	198
		VLS2	Viral Load	193
Human herpesvirus 8		ID1	Nucleic Acid Amp, Viruses	196
Human immuno- deficiency virus (HIV)	Х	HIVG, HV2	HIV Viral Load	193
		HIVG	HIV Genotyping	193
		LN39	HIV Viral Load Cal Ver/ Lin	127
Human metapneumovirus		ID2	Nucleic Acid Amp, Respiratory	197
	Х	IDR	Infectious Disease, Respiratory Panel	198
Human papillomavirus (cytology) high-risk	Х	CHPVD	Digene Specimen Transport Medium	274
· · · · · · · · · · · · · · · · · · ·	Х	CHPVJ	Mixed Medium	274
	Х	CHPVK	SurePath Preservative Fluid Transport Medium	274
	Х	СНРУМ	ThinPrep PreservCyt Transport Medium	274
		HPV	Digene Hybrid Capture Technology Only	195
		ISH	In Situ Hybridization	250
Human papillomavirus (cytology) high-risk genotyping		CHPVJ	Mixed Medium	274
		CHPVM	ThinPrep PreservCyt Transport Medium	274
Human parechovirus		IDME	Meningitis/Encephalitis Panel	198
Huntington disease	Х	MGL2	Molecular Genetics	241
Hydrocodone		DFC	Drug-Facilitated Crime	107
		DMPM	Drug Monitoring for Pain Management	106
		FTC	Forensic Toxicology, Criminalistics	104
		OFD	Oral Fluid for Drugs of Abuse	100
		Т	Toxicology	96
		UDC	Forensic Urine Drug	99
			Testing, Confirmatory	
		UT	Urine Toxicology	96
Hydromorphone		DFC	Drug-Facilitated Crime	107
		DMPM	Drug Monitoring for Pain Management	106
		FTC	Forensic Toxicology, Criminalistics	104
		OFD	Oral Fluid for Drugs of Abuse	100

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Hydromorphone (cont.)		T	Toxicology	96
		UDC	Forensic Urine Drug	99
			Testing, Confirmatory	
		UT	Urine Toxicology	96
Hydroxyzine		T	Toxicology	96
		UT	Urine Toxicology	96
Ibuprofen		Т	Toxicology	96
		UT	Urine Toxicology	96
IDH1		GLI	Glioma	253
IDH2		GLI	Glioma	253
IgA	Χ	IG/IGX	Immunology, General	202
		LN7	Immunology Cal Ver/Lin	121
IgA, electrophoresis	Χ	SPE	Protein Electrophoresis	76
IgD		S2, S4	Immunology, Special	203
IgE	Х	IG/IGX	Immunology, General	202
	Χ	K/KK	Ligand Assay, General	82
	Х	SE	Diagnostic Allergy	207
IgE allergen-specific, quantitative		SE	Diagnostic Allergy	207
IgE multi-allergen screen	Х	SE	Diagnostic Allergy	207
IGF-1 (somatomedin C)	Х	BGS	Bone and Growth	85
	Х	Y/YY	Ligand Assay, Special	84
IgG	Х	IG/IGX	Immunology, General	202
		LN7	Immunology Cal Ver/Lin	121
		S2, S4	Immunology, Special	203
IgG, electrophoresis	Х	SPE	Protein Electrophoresis	76
IgG, CSF	Х	M, OLI	CSF Chemistry and	74
			Oligoclonal Bands	
IgG subclass proteins		S2, S4	Immunology, Special	203
IGHV		IGHV	Mutation Analysis	250
IgM	Х	IG/IGX	Immunology, General	202
		LN7	Immunology Cal Ver/Lin	121
IgM, electrophoresis	Χ	SPE	Protein Electrophoresis	76
Ilb/Illa assay		PIA, PIAX	Drug-Specific Platelet Aggregation	165
IL-2		CTKN	Cytokines	206
IL-6		CTKN	Cytokines	206
IL-8		CTKN	Cytokines	206
IL-10		CTKN	Cytokines	206
IL28B		PGX1	Pharmacogenetics	243
Imipramine		DFC	Drug-Facilitated Crime	107
		FTC	Forensic Toxicology,	104
			Criminalistics	
		Т	Toxicology	96
		UT	Urine Toxicology	96
	Χ	ZT	TDM, Special	60
Immature granulocyte		FH9, FH9P	Hematology, Auto Diff	135
parameter		DDAE\/	DDAEVCOOF	207
Immunohistochemistry		BRAFV	BRAF V600E	267
		GHER2	Gastric HER2	269

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Immunohistochemistry	Х	HER2	HER2 by	269
(cont.)			Immunohistochemistry	
		MK	Immunohistochemistry	267
		MMR	DNA Mismatch Repair	267
		PDL1	PDL1	267
		PM1	CD117 by	268
			Immunohistochemistry	
	X	PM2	ER, PR by Immunohistochemistry	269
		PM3	CD20 by	268
			Immunohistochemistry	
		PM5	Immunohistochemistry TMA	268
		PM6	Anaplastic Lymphoma	268
			Kinase IHC	
India ink		IND	India Ink	186
Infectious	Х	IL, IM	Immunology	202
mononucleosis (IM)				
	Х	IMW	Infectious Mononucleosis, Waived	203
Influenza virus		ID2	Nucleic Acid Amp, Resp	197
	Х	ID3	Influenza A, Influenza B, RSV by NAA	197
	Х	IDR	Infectious Disease,	198
			Respiratory Panel	
		POC8	POC Influenza A/B Ag	52
	X	VR1	Virology Culture	191
	X	VR2	Viral Antigen Detection by DFA	191
	Х	VR4	Viral Antigen Detection by EIA and Latex	192
In situ hybridization	Х	ISH	In Situ Hybridization	250
	Х	ISH2	In Situ Hybridization HER2	250
Instrument function		I	Instrumentation	129
Instrument linearity		I	Instrumentation	129
		LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
		LN2BV	Chemistry, Lipid,	118
			Enzyme all Beckman	
			except AU, Vitros Cal	
			Ver/Lin	
		LN3	TDM Cal Ver/Lin	119
		LN5	Ligand Assay Cal Ver/Lin	119-
				120
		LN5S	Ligand Assay, Siemens	119-
		LNC	Cal Ver/Lin	120
		LN6	Urine Chemistry Cal Ver/Lin	120
		LN7	Immunology Cal Ver/Lin	121

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Instrument linearity (cont.)		LN8	Reproductive Endocrinology Cal Ver/ Lin	121
		LN9	Hematology Cal Ver/Lin	121
		LN11	Serum Ethanol Cal Ver/ Lin	122
		LN12, LN12E	C-Reactive Protein Cal Ver/Lin	122
		LN13	Blood Gas Cal Ver/Lin	122- 123
		LN13C	Blood Gas Cal Ver/Lin	122- 123
		LN15	Hemoglobin A _{1c} Cal Ver/Lin	123
		LN16	Homocysteine Cal Ver/ Lin	123
		LN17	Whole Blood Glucose Cal Ver/Lin	123
		LN18, LN19	Reticulocyte Cal Ver/Lin	124
		LN20	Urine Albumin Cal Ver/ Lin	124
		LN21	High-Sensitivity C-Reactive Protein Cal Ver/Lin	124
		LN22	Flow Cytometry Cal Ver/Lin	124
		LN23	PSA Cal Ver/Lin	125
		LN24	Creatinine Accuracy Cal Ver/Lin	125
		LN25	Troponin I Cal Ver/Lin	125
		LN27	Troponin T Cal Ver/Lin	125
		LN30	BNP Cal Ver/Lin	125
		LN31	Immunosuppressive Drugs Cal Ver/Lin	126
		LN32	Ammonia Cal Ver/Lin	126
		LN33	Serum Myoglobin Cal Ver/Lin	126
		LN34	Tumor Markers Cal Ver/ Lin	126
		LN35	Thrombophilia Cal Ver/ Lin	127
		LN36	Heparin Cal Ver/Lin	127
		LN37	von Willebrand Factor Ag Cal Ver/Lin	127
		LN38	CMV viral load Cal Ver/ Lin	127
		LN39	HIV Viral Load Cal Ver/ Lin	127
		LN40	Vitamin D Cal Ver/Lin	127
		LN41	Procalcitonin Cal Ver/ Lin	128
		LN42	D-Dimer Cal Ver/Lin	128

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Instrument linearity (cont.)		LN43	Lamellar Body Count Cal Ver/Lin	128
		LN44	Fibrinogen Cal Ver/Lin	128
		LN45	HCV Viral Load Cal Ver/ Lin	127
Insulin	Х	ING	Insulin, Gastrin, C-Peptide, PTH	86
Interferon (IFN) gamma		CTKN	Cytokines	206
Interleukin (IL)-1 beta		CTKN	Cytokines	206
International normalized ratio (INR)	Х	CGB	Basic Coagulation	158
	Х	CGL	Coagulation, Limited	158
		CGLQ	Quality Cross Check, Coagulation, Limited	49
		CGS1	Coag Special, Series 1	160- 161
		CGS4	Coag Special, Series 4	160- 161
		POC6	POC PT/INR, CoaguChek XS Plus	52
	Х	WP3, WP4, WP6, WP9	Whole Blood Coagulation	166
		WP10	Whole Blood Coagulation	166
Ionized calcium	Х	AQ, AQ2, AQ3, AQ4	Aqueous Blood Gas	92
		AQQ, AQ2Q, AQ3Q, AQ4Q	Quality Cross Check, Critical Care Aqueous Blood Gas Series	46
	Х	C3, CZ, CZX	Chemistry and TDM	56- 58
		P0C10, P0C11	POC Competency Blood Gases	53
Iron	Х	C1, C3, C3X, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		IFS	Interfering Substances	130
		LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
		LN2BV	Chemistry, Lipid, Enzyme all Beckman except AU, Vitros Cal Ver/Lin	118
Isopropanol	Х	AL1	Whole Blood Alcohol/ Ethylene Glycol/ Volatiles	101
	Х	AL2	Serum Alcohol/Ethylene Glycol/Volatiles	101
JC virus		ID1T	Nucleic Acid Amp, JC and BK	196

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Jo-1 (antihistidyl t-RNA		RDS	Rheumatic Disease	207
synthetase)			Special	
Kaolin-activated APTT		CGE/CGEX	Coagluation, Extended	159
Kaolin-activated CT		CGE/CGEX	Coagluation, Extended	159
Kappa/Lambda	Х	ISH	In Situ Hybridization	250
Kappa/Lambda ratio		IG/IGX	Immunology, General	202
		S2, S4	Immunology, Special	203
Free Kappa/Lambda ratio		SFLC	Serum Free Light Chains	208
Karyotype nomenclature	Х	CY, CYBK	Cytogenetics	236
Ketamine		DFC	Drug-Facilitated Crime	107
		FTC	Forensic Toxicology, Criminalistics	104
		T	Toxicology	96
		UT	Urine Toxicology	96
Ketones, serum		KET	Ketones	64
Ketones, urine	Х	CMP, CMP1	Clinical Microscopy	144
		CMQ	Quality Cross Check, Urinalysis	48
	Х	HCC2	Waived Combination	66
		POC3	POC Urine Dipstick	52
			Competency	
Kidney stone assessment		KSA	Kidney Stone	69
-			Assessment	
KIT		KIT	KIT/PDGFRA	252
		MTP	Multigene Tumor Panel	253
KOH prep (skin or	Х	CMMP	Clinical Microscopy,	145
vaginal)		5014	Misc	400
	X	FSM	Fungal Smear	186
KRAS	X	KRAS	Colorectal Cancer	252
	V	MTD	Multigana Tumar Danal	253
	Х	MTP	Multigene Tumor Panel	
Laboratory preparedness exercise		LPX	Laboratory Preparedness Exercise	180
Lacosamide		7	'	60
Lacosamide		ZE	Therapeutic Drug Monitoring, Extended	60
Lactate	X	AQ, AQ2, AQ3,	Aqueous Blood Gas	92
Lactate		AQ4	Aqueous blood das	32
		AQQ, AQ2Q,	Quality Cross Check,	46
		AQ3Q, AQ4Q	Critical Care Aqueous	
			Blood Gas Series	
	Х	C3, C3X, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		FLD	Body Fluid	72
		FLDQ	Quality Cross Check, Body Fluid Chemistry	44
		P0C10, P0C11	POC Competency Blood Gases	53
		LN13C	Blood Gas Cal Ver/Lin	122-
			33.75.25.75.75.75.75.75.75.75.75.75.75.75.75.75	123

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Lactate, CSF	Χ	M, OLI	CSF Chemistry and	74
			Oligoclonal Bands	
Lactate dehydrogenase (LD)	X	C1, C3, C3X, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check,	43
			Chemistry and TDM	
		FLD	Body Fluid	72
		FLDQ	Quality Cross Check, Body Fluid Chemistry	44
		IFS	Interfering Substances	130
		LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
		LN2BV	Chemistry, Lipid, Enzyme all Beckman except AU, Vitros Cal Ver/Lin	118
		SC0	Serum Carryover	131
Lactate dehydrogenase (LD), CSF	Х	M, OLI	CSF Chemistry and Oligclonal Bands	74
Lamellar body count		LBC	Lamellar Body Count	149
		LN43	Lamellar Body Count Cal Ver/Lin	128
Lamotrigine		Т	Toxicology	96
		UT	Urine Toxicology	96
		ZE	Therapeutic Drug Monitoring, Extended	60
Large unclassified cells (LUC)		FH4, FH4P	Hematology, Auto Diff	135
LD isoenzymes	Х	CRTI	Cardiac Markers	62
LD1/LD2 ratio	Х	CRTI	Cardiac Markers	62
LDL cholesterol	Х	ABL	Accuracy-Based Lipid	110
LDL cholesterol, calculated		C1, C3, C3X, C4, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
LDL cholesterol, measured	X	C1, C3, C3X, C4, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
LDL cholesterol, waived	Χ	LCW	Ltd Chem, Waived	64
Lead (blood)	Х	BL	Blood Lead	102
Lead, urine		TMU	Trace Metals, Urine	103
Legionella		LBAS	Legionella Ag	176
Legionella pneumophila		IDN, IDO	Nucleic Acid Amp, Organisms	196
	Х	IDR	Infectious Disease, Respiratory Panel	198
Leukemia/lymphoma immunophenotype		FL3	Flow Cytometry	209
Leukemia/lymphoma		FL5	Flow Cytometry	210
interpretation only			Interpretation Only	

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Leukocyte esterase, urine	Χ	CMP, CMP1	Clinical Microscopy	144
		CMQ	Quality Cross Check,	48
			Urinalysis	
	Χ	HCC2	Waived Combination	66
		POC3	POC Urine Dipstick	52
			Competency	
Leukocyte-reduced platelets		TRC	Transfusion-Related Cell Count	218
Leukocyte-reduced RBC		TRC	Transfusion-Related Cell Count	218
Leukocyte, stool, Wright- Giemsa	Х	CMMP	Clinical Microscopy, Misc	145
Levetiracetam		Т	Toxicology	96
		UT	Urine Toxicology	96
		ZE	Therapeutic Drug	60
			Monitoring, Extended	
Lidocaine	Х	CZ, CZ2X,	Chemistry and TDM	56-
		CZX, Z		58
		CZQ	Quality Cross Check,	43
			Chemistry and TDM	
		LN3	TDM Cal Ver/Lin	119
		Т	Toxicology	96
		UT	Urine Toxicology	96
Lipase	Х	C3, C3X, CZ,	Chemistry and TDM	56-
ļ		CZ2X, CZX	,	58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		FLD2	Body Fluid Chemistry 2	73
		IFS	Interfering Substances	130
		LN2	Chemistry, Lipid,	118
			Enzyme Cal Ver/Lin	
		LN2BV	Chemistry, Lipid,	118
			Enzyme all Beckman	
			except AU, Vitros Cal Ver/Lin	
Lipids		ABL	Accuracy-Based Lipid	110
	Х	C1, C3, C3X, CZ, CZ2X,	Chemistry and TDM	56- 58
		CZX		
		CZQ	Quality Cross Check, Chemistry and TDM	43
		LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
		LN2BV	Chemistry, Lipid, Enzyme all Beckman except AU, Vitros Cal Ver/Lin	118
Lipoprotein (a)	Χ	ABL	Accuracy-Based Lipid	110
	Х	C1, C3, C3X, CZ, CZ2X,	Chemistry and TDM	56- 58
		CZX		

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Lipoprotein (a) (cont.)		CZQ	Quality Cross Check,	43
			Chemistry and TDM	
Lipoprotein-associated phospholipase		PLA	Lp-PLA ₂	75
Lipoprotein electrophoresis		LPE	Lipoprotein Electrophoresis	76
Listeria monocytogenes		IDME	Meningitis/Encephalitis Panel	198
Lithium	X	C1, C3, C3X, CZ, CZ2X, CZX, Z	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		LN3	TDM Cal Ver/Lin	119
Liver-kidney microsomal antibody		LKM	Liver-Kidney Microsomal Antibody	207
Lorazepam		DFC	Drug-Facilitated Crime	107
		DMPM	Drug Monitoring for Pain Management	106
		FTC	Forensic Toxicology, Criminalistics	104
		T	Toxicology	96
		UDC	Forensic Urine Drug Testing, Confirmatory	99
		UT	Urine Toxicology	96
Lorazepam glucuronide		DMPM	Drug Monitoring for Pain Management	106
Lupus anticoagulant (screen, conf)		CGE/CGEX	Coagulation, Extended	159
		CGS1	Coag Special, Series 1	160- 161
Luteinizing hormone (LH)		ABS	Accuracy-Based Testosterone, Estradiol	111
		LN8	Reproductive Endocrinology Cal Ver/ Lin	121
	Χ	Y/YY	Ligand Assay, Special	84
Lyme disease		TTD	Tick-Transmitted Disease	200
Lymphocyte immunophenotyping	Х	FL, FL1	Flow Cytometry	209
Lymphoma by FISH		CYL	Fluorescence In Situ Hybridization, Lymphoma	237
Lysergic acid diethylamide (LSD)		FTC	Forensic Toxicology, Criminalistics	104
		T	Toxicology	96
		UDS, UDS6	Urine Drug Screen	98
		UT	Urine Toxicology	96
Magnesium	X	C1, C3, C3X, CZ, CZ2X, CZX	Chemistry and TDM	56- 58

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Magnesium (cont.)		CZQ	Quality Cross Check, Chemistry and TDM	43
		IFS	Interfering Substances	130
		LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
		LN2BV	Chemistry, Lipid, Enzyme all Beckman except AU, Vitros Cal Ver/Lin	118
Magnesium, ionized	Χ	AQ, AQ2	Aqueous Blood Gas	92
		AQQ, AQ2Q	Quality Cross Check, Critical Care Aqueous Blood Gas Series	46
		POC10, POC11	POC Competency Blood Gases	53
Magnesium, urine	Х	U	Urine Chemistry, General	68
Malaria		RMAL	Rapid Malaria	188
Manganese		R	Trace Metals	78
		TMU	Trace Metals, Urine	103
		TMWB	Trace Metals, Whole Blood	103
MCAD	Х	IMD2	MCAD	242
МСН		FH1-FH4, FH6, FH9, FH10, FH13, FH1P-FH4P, FH6P, FH9P, FH10P, FH13P	Hematology, Auto Diff	135
		FH3Q, FH4Q, FH6Q, FH9Q	Quality Cross Check, Automated Hematology Series	47
		HE, HEP	Basic Hematology	134
мснс		FH1-FH4, FH6, FH9, FH10, FH13, FH1P-FH4P, FH6P, FH9P, FH10P, FH13P	Hematology, Auto Diff	135
		FH3Q, FH4Q, FH6Q, FH9Q	Quality Cross Check, Automated Hematology Series	47
		HE, HEP	Basic Hematology	134
MCV		FH1-FH4, FH6, FH9, FH10, FH13, FH1P-FH4P, FH6P, FH9P, FH10P, FH13P	Hematology, Auto Diff	135

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
MCV (cont.)		FH3Q, FH4Q,	Quality Cross Check,	47
		FH6Q, FH9Q	Automated Hematology	
			Series	
		HE, HEP	Basic Hematology	134
MECP2 deletion/		RETT	RETT Syndrome	243
duplication analysis			Genotyping	
MECP2 genotyping	Х	RETT	RETT Syndrome	243
MEN2	X	MCL2	Genotyping Molecular Genetics	241
	^	MGL3		
Meperidine		DFC	Drug-Facilitated Crime	107
		DMPM	Drug Monitoring for Pain Management	106
		FTC	Forensic Toxicology,	104
			Criminalistics	
		Т	Toxicology	96
		UT	Urine Toxicology	96
Mephedrone		Т	Toxicology	96
		UT	Urine Toxicology	96
Meprobamate		DFC	Drug-Facilitated Crime	107
		DMPM	Drug Monitoring for Pain	106
			Management	100
		FTC	Forensic Toxicology,	104
			Criminalistics	
		Т	Toxicology	96
		UT	Urine Toxicology	96
Mercury, urine		TMU	Trace Metals, Urine	103
- Wichout y, utilitie		TMWB	Trace Metals, Whole	103
			Blood	
Metabolic disease testing		BGL	Biochemical Genetics	239
Metanephrine	Х	N/NX	Urine Chemistry, Special	69
Methadone		DFC	Drug-Facilitated Crime	107
		DMPM	Drug Monitoring for Pain Management	106
		FTC	Forensic Toxicology,	104
		050	Criminalistics	100
		OFD	Oral Fluid for Drugs of Abuse	100
		Т	Toxicology	96
		UDC	Forensic Urine Drug	99
			Testing, Confirmatory	
		UDS, UDS6	Urine Drug Screen	98
		UT	Urine Toxicology	96
Methadone metabolite (EDDP)		DFC	Drug-Facilitated Crime	107
		DMPM	Drug Monitoring for Pain Management	106
		FTC	Forensic Toxicology, Criminalistics	104
		Т	Toxicology	96
		1	IOVICOTORA	30

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Methadone metabolite (EDDP) (cont.)		UDC	Forensic Urine Drug Testing, Confirmatory	99
(LDDF) (COIIC.)		TIDE TIDES	Urine Drug Screen	98
		UDS, UDS6		96
Mathamahatamina		DFC	Urine Toxicology	
Methamphetamine			Drug-Facilitated Crime	107
		DMPM	Drug Monitoring for Pain Management	106
		FTC	Forensic Toxicology,	104
		FIC	Criminalistics	104
		OFD	Oral Fluid for Drugs of	100
		OID	Abuse	100
		Т	Toxicology	96
		UDC	Forensic Urine Drug	99
			Testing, Confirmatory	
		UDS, UDS6	Urine Drug Screen	98
		UT	Urine Toxicology	96
Methanol	Х	AL1	Whole Blood Alcohol/	101
			Ethylene Glycol/	
			Volatiles	
	X	AL2	Serum Alcohol/Ethylene Glycol/Volatiles	101
Methaqualone		UDC	Forensic Urine Drug	99
Methaquatone		ODC	Testing, Confirmatory	33
		UDS, UDS6	Urine Drug Screen	98
Methemoglobin	X	S0	Blood Oximetry	94
- Inochomographi	\(\)	SOQ	Quality Cross Check,	46
		004	Blood Oximetry	
Methicillin-resistant Staphylococcus aureus (MRSA)		BCS1	Blood Culture Staphylococcus aureus	177
		IDN, IDO	Nucleic Acid Amp, Organisms	196
		MRS	Methicillin-Resistant S.	179
		MRS2M	MRSA Screen,	179
			Molecular, 2 Challenge	
	Х	MRS5	Methicillin-Resistant S. aureus	179
	X	MRS5M	MRSA Screen,	179
	^	WIROOW	Molecular, 5 Challenge	173
Methotrexate	Х	CZ, CZ2X,	Chemistry and TDM	56-
		CZX, Z		58
		CZQ	Quality Cross Check, Chemistry and TDM	43
Methylenedioxy-		DFC	Drug-Facilitated Crime	107
amphetamine (MDA)		DI C	Ping-i acilitated cillie	107
		DMPM	Drug Monitoring for Pain Management	106
		FTC	Forensic Toxicology,	104
		1 10	Criminalistics	104
	1		J	<u> </u>

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Methylenedioxy- amphetamine (MDA) (cont.)		OFD	Oral Fluid for Drugs of Abuse	100
		T	Toxicology	96
		UDC	Forensic Urine Drug	99
			Testing, Confirmatory	
		UT	Urine Toxicology	96
Methylenedioxyethyl- amphetamine (MDEA)		OFD	Oral Fluid for Drugs of Abuse	100
		UDC	Forensic Urine Drug Testing, Confirmatory	99
Methylenedioxymeth- amphetamine (MDMA)		DFC	Drug-Facilitated Crime	107
		DMPM	Drug Monitoring for Pain Management	106
		FTC	Forensic Toxicology, Criminalistics	104
		OFD	Oral Fluid for Drugs of Abuse	100
		Т	Toxicology	96
		UDC	Forensic Urine Drug	99
		TIDO TIDOC	Testing, Confirmatory	
		UDS, UDS6	Urine Drug Screen	98
Mothylopodiovy		T	Urine Toxicology	96
Methylenedioxy- pyrovalerone (MDPV)			Toxicology	96
		UT	Urine Toxicology	96
Methylenetetra- hydrofolate reductase (MTHFR)	X	MGL1	Molecular Genetics	241
Methylphenidate		T	Toxicology	96
		UT	Urine Toxicology	96
Metoprolol		Т	Toxicology	96
		UT	Urine Toxicology	96
MGMT		GLI	Glioma	253
Microalbumin, urine		LN20	Urine AlbuminCal Ver/ Lin	124
	X	U	Urine Chemistry	68
	X	UMC	Urine Albumin (Microalbumin)/ Creatinine	151
Microsatellite instability		MSI	Microsatellite Instability	250
Microtiter plate reader linearity		I	Instrumentation	129
Minimal residual disease		BALL	B-ALL Minimal Residual Disease	210
		MRD	Minimal Residual Disease, BCR/ABL1 p210	254
		MRD1	Minimal Residual Disease, <i>BCR/ABL1</i> p190	254

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Minimal residual disease		MRD2	Minimal Residual	254
(cont.)			Disease, PML/RARA	
Mirtazapine		T	Toxicology	96
		UT	Urine Toxicology	96
Mite identification		TMO	Ticks, Mites, and Other	189
			Arthropods	
Mitochondrial	X	IMD3	Mitochondrial	242
cytopathies			Cytopathies	
Mitochondrial DNA	X	IMD1	Mitochondrial DNA	242
deletion syndromes		205/205/	Deletion Syndromes	4=0
Mixing studies, PT		CGE/CGEX	Coagulation, Extended	159
Mixing studies, APTT		CGE/CGEX	Coagulation, Extended	159
		CGS1	Coag Special, Series 1	160- 161
MLH1 promoter		MSI	Defective DNA	250
methylation analysis			Mismatch Repair/	
			Hereditary	
			Nonpolyposis Colorectal	
NA PC L CLES	\ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \	D D0 D/ D5	Cancer (HNPCC)	407
Modified acid-fast stain	X	P, P3, P4, P5	Parasitology	187
Mold identification	Х	F	Mycology and Aerobic Actinomycetes	184
Molecular genetics	X	MGL1, MGL2, MGL3, MGL4, MGL5	Molecular Genetics	241
Molecular HLA typing	Х	DML	HLA Molecular Typing	231
Molecular hematologic oncology		MHO, MHO1, MHO2, MHO3, MHO5	Molecular Hematologic Oncology	251, 254
Molecular typing		IDN, IDO	Nucleic Acid Amp, Organisms	196
Monitoring engraftment	Х	ME	Monitoring Engraftment	232
Mononuclear cell count		CBT	Cord Blood Testing	221
		SCP	Stem Cell Processing	221
Morphine		DFC	Drug-Facilitated Crime	107
		DMPM	Drug Monitoring for Pain Management	106
		FTC	Forensic Toxicology, Criminalistics	104
		OFD	Oral Fluid for Drugs of Abuse	100
		T	Toxicology	96
		UDC	Forensic Urine Drug Testing, Confirmatory	99
		UT	Urine Toxicology	96
M-protein (paraprotein)	X	SPE	Protein Electrophoresis	76
identification				

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
MPV		FH1-FH4, FH6, FH9, FH10, FH13, FH1P-FH4P, FH6P, FH9P, FH10P, FH13P	Hematology, Auto Diff	135
		FH3Q, FH4Q, FH6Q, FH9Q	Quality Cross Check, Automated Hematology Series	47
MRSA		HE, HEP BCS1	Basic Hematology Blood Culture Staphylococcus aureus	134
		IDN, IDO	Nucleic Acid Amp, Organisms	196
		MRS	Methicillin-Resistant S. aureus	179
		MRS2M	MRSA Screen, Molecular, 2 Challenge	179
	X	MRS5	Methicillin-Resistant S. aureus	179
	X	MRS5M	MRSA Screen, Molecular, 5 Challenge	179
Mucolipidosis IV	X	MGL4	Molecular Genetics	241
Mucopolysaccharide (Glycosaminoglycan)	Х	BGL	Biochemical Genetics	239
Multiple endocrine neoplasia type 2 (MEN2)	X	MGL3	Molecular Genetics	241
Mumps-IgG		VR3M	Virology	200
Mycobacterial culture	Х	E1	Mycobacteriology, Ltd	183
Mycobacterial identification	Х	Е	Mycobacteriology	183
Mycobacterium tuberculosis		IDO	Nucleic Acid Amp, Organisms	196
Mycobacterium tuberculosis antibody detection		QF	M. tuberculosis Infection Detection	207
Mycobacterium tuberculosis identification and resistance detection		MTBR	Molecular MTB Identification and Resistance Detection	183
Mycophenolic acid	X	MPA	Mycophenolic Acid	59
Mycoplasma pneumoniae		IDN, IDO	Nucleic Acid Amp, Organisms	196
	Х	IDR	Infectious Disease, Respiratory Panel	198
		VR3	Antibody Detection- Infectious Disease Serology	200
Myoglobin	Х	CRT, CRTI	Cardiac Markers	62
		LN33	Serum Myoglobin Cal Ver/Lin	126

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Myoglobin (cont.)	Χ	PCARM, PCARMX	Plasma Cardiac Markers	65
		POC12	Competency Plasma Cardiac Markers	53
Myoglobin, urine		MYG	Myoglobin, Urine	69
Myotonic dystrophy	Χ	MGL2	Molecular Genetics	241
N-acetylprocainamide	Х	CZ, CZ2X,	Chemistry and TDM	56-
(NAPA)		CZX,Z		58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		LN3	TDM Cal Ver/Lin	119
N-desmethyltramadol		DMPM	Drug Monitoring for Pain Management	106
		FTC	Forensic Toxicology,	104
			Criminalistics	
		T	Toxicology	96
		UT	Urine Toxicology	96
Naproxen		Т	Toxicology	96
<u>.</u>		UT	Urine Toxicology	96
Nasal smears, eosinophil	Х	CMMP	Clinical Microscopy, Misc	145
Neisseria gonorrhoeae	Х	D3	GC Cultures	172
	Χ	D4	Bacteriology, Limited	173
	Х	HC6/HC6X	C. trachomatis/GC by	194
			Nucleic Acid Amp	
	Х	HC7	C. trachomatis/GC DNA by NAA	194
	Х	MC1	Microbiology Combination with GC	173
Neisseria meningitidis		IDME	Meningitis/Encephalitis	198
Neoplastic cellularity		NEO	Neoplastic Cellularity	251
Neoplastic disorder by		CYF	Fluorescence In Situ	236
FISH			Hybridization	
Neuropathology		NP/NP1	Neuropathology Program	271
Neutral fats		FCFS	Fecal Fat	75
Next-generation sequencing		NGS	Next-Generation Sequencing	246
		NGSB1	NGS Bioinformatics for Illumina Platforms	247
		NGSB2	NGS Bioinformations for	247
		NGSBV	NGS Bioinformatics Somatic Validated Materials	249
		NGSE	NGS Undiagnosed Disorders-Exome	248
		NGSST	Next Generation Sequencing, Solid Tumor	246

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Next-generation		NGSHM	Next Generation	246
sequencing (cont.)			Sequencing,	
			Hematologic	
			Malignancies	
Nicotine		NTA	Nicotine and Tobacco Alkaloids	102
		Т	Toxicology	96
		UT	Urine Toxicology	96
Niemann Diek tyne A/P	X	MGL4	Molecular Genetics	241
Niemann-Pick type A/B NIPT	^	NIPT	Noninvasive Prenatal	87
MPI		MIPI	Testing	0/
Nitrite, urine	X	CMP, CMP1	Clinical Microscopy	144
		CMQ	Quality Cross Check,	48
			Urinalysis	
		DAI	Urine Drug Adulterant/	98
			Integrity Testing	
	X	HCC2	Waived Combination	66
		POC3	POC Urine Dipstick	52
			Competency	
Nitrogen, total, urine		U	Urine Chemistry,	68
			General	
Nongynecologic		FNA/FNA1	Fine-Needle Aspiration-	277
cytopathology			Digital	
		FNAG/FNAG1	Fine-Needle Aspiration- Glass	278
		NGC/NGC1	Nongynecologic	276
			Cytopathology	
			Education Program	
Noninvasive prenatal		NIPT	Noninvasive Prenatal	87
testing			Testing	
Norbuprenorphine		DMPM	Drug Monitoring for Pain	106
			Management	
		OFD	Oral Fluid for Drugs of	100
			Abuse	
		Т	Toxicology	96
		UDC	Forensic Urine Drug	99
			Testing, Confirmatory	
		UT	Urine Toxicology	96
Norchlordiazepoxide		Т	Toxicology	96
		UT	Urine Toxicology	96
Norclomipramine		Т	Toxicology	96
		UT	Urine Toxicology	96
Norcodeine		T	Toxicology	96
		UT	Urine Toxicology	96
Norcyclobenzaprine		T	Toxicology	96
		UT	Urine Toxicology	96
Nordiazepam		DMPM	Drug Monitoring for Pain Management	106
		FTC	Forensic Toxicology, Criminalistics	104

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Nordiazepam (cont.)		OFD	Oral Fluid for Drugs of Abuse	100
		T	Toxicology	96
		UDC	Forensic Urine Drug	99
			Testing, Confirmatory	
		UT	Urine Toxicology	96
Nordoxepin		DFC	Drug-Facilitated Crime	107
		FTC	Forensic Toxicology,	104
			Criminalistics	
		Т	Toxicology	96
		UT	Urine Toxicology	96
Norepinephrine	Х	N/NX	Urine Chemistry, Special	69
Norfentanyl		DMPM	Drug Monitoring for Pain	106
			Management	
		FTC	Forensic Toxicology,	104
			Criminalistics	
		Т	Toxicology	96
		UT	Urine Toxicology	96
Norfluoxetine		DFC	Drug-Facilitated Crime	107
		FTC	Forensic Toxicology, Criminalistics	104
		Т	Toxicology	96
		UT	Urine Toxicology	96
Norketamine		DFC	Drug-Facilitated Crime	107
		FTC	Forensic Toxicology,	104
			Criminalistics	
		Т	Toxicology	96
		UT	Urine Toxicology	96
Normeperidine		DFC	Drug-Facilitated Crime	107
		DMPM	Drug Monitoring for Pain Management	106
		T	Toxicology	96
		UT	Urine Toxicology	96
Normetanephrine	Х	N/NX	Urine Chemistry Special	69
Norovirus		GIP	Gastrointestinal Panel	199
		SP1	Stool Pathogens	180
Noroxycodone		DMPM	Drug Monitoring for Pain Management	106
		T	-	06
			Toxicology	96
Navayyymaynhana		UT	Urine Toxicology Drug Monitoring for Pain	96
Noroxymorphone		DMPM	Management Management	106
Norpropoxyphene		DFC	Drug-Facilitated Crime	107
		DMPM	Drug Monitoring for Pain	106
-			Management	
		FTC	Forensic Toxicology,	104
			Criminalistics	
		Т	Toxicology	96
		UDC	Forensic Urine Drug Testing, Confirmatory	99

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Norpropoxyphene (cont.)		UT	Urine Toxicology	96
Norsertraline		DFC	Drug-Facilitated Crime	107
		FTC	Forensic Toxicology,	104
			Criminalistics	
		Т	Toxicology	96
		UT	Urine Toxicology	96
Nortrimipramine		Т	Toxicology	96
		UT	Urine Toxicology	96
Nortriptyline		DFC	Drug-Facilitated Crime	107
		FTC	Forensic Toxicology, Criminalistics	104
		Т	Toxicology	96
		UT	Urine Toxicology	96
	Х	ZT	TDM, Special	60
Norverapamil		Т	Toxicology	96
		UT	Urine Toxicology	96
NRAS		MTP	Multigene Tumor Panel	253
nRBC		FH3, FH3P, FH9, FH9P, FH13, FH13P	Hematology, Auto Diff	135
NT-pro B-type natriuretic peptides	Х	BNP	B-Type Natriuretic Peptides, 2 Chall	61
	Х	BNP5	B-Type Natriuretic Peptides, 5 Chall	61
		BNPQ	Quality Cross Check, B-Type Natriuretic Peptides	43
		LN30	BNP Cal Ver/Lin	125
N-telopeptide (NTX)		BMV6	Bone Markers and Vitamin	86
-	Χ	BU	Bone and Mineral, Urine	85
Nucleated cells, total		CBT	Cord Blood Testing	221
		SCP	Stem Cell Processing	221
Nucleated red cells, total		ABF3	Automated Body Fluid	146
		CBT	Cord Blood Testing	221
		SCP	Stem Cell Processing	221
Nucleated red blood cell count		FH3, FH3P, FH9, FH9P, FH13, FH13P	Hematology, Auto Diff	135
Nucleic acid amplification		BSTS	Bacterial Strain Typing Staphylococcus	195
	Х	HBVL, HBVL5, HCV2	Hepatitis Viral Load	193
	Х	HC6/HC6X	C. trachomatis/GC by Nucleic Acid Amp	194
	Х	НС7	C. trachomatis/GC DNA by NAA	194
	Х	HIVG, HV2	HIV Viral Load	193
		IDN, IDO	Nucleic Acid Amp, Organisms	196

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Nucleic acid		ID1, ID1T	Nucleic Acid Amp,	196
amplification (cont.)			Viruses	
		ID2	Nucleic Acid Amp,	197
			Respiratory	
		ID3	Influenza A, Influenza B,	197
			RSV by NAA	
		MRS2M	MRSA Screen,	179
			Molecular, 2 Challenge	
	Χ	MRS5M	MRSA Screen,	179
			Molecular, 5 Challenge	
		SP, SPN, SP1	Stool Pathogens	180
		VLS, VLS2	Viral Load	193
		VRE	Vancomycin-Resistant	182
			Enterococcus	
Nucleic acid testing	Х	NAT	Nucleic Acid Testing	226
O-desmethyltramadol		DMPM	Drug Monitoring for Pain	106
			Management	
		FTC	Forensic Toxicology,	104
			Criminalistics	
		Т	Toxicology	96
		UT	Urine Toxicology	96
Occult blood		OCB	Occult Blood	149
		OCBQ	Quality Cross Check,	48
			Occult Blood	
		POC9	POC Fecal Occult Blood	52
Occult blood, gastric		GOCB	Gastric Occult Blood	148
Ocular micrometer check		I	Instrumentation	129
Olanzapine		Т	Toxicology	96
		UT	Urine Toxicology	96
Oligoclonal bands		OLI	Oligoclonal Bands	74
Opiate group		DMPM	Drug Monitoring for Pain	106
			Management	
		OFD	Oral Fluid for Drugs of	100
			Abuse	
		Т	Toxicology	96
Delayed until 20	19	TQP	Toxicology Quality	108
		LIDO LIDOS	Program	
		UDS, UDS6	Urine Drug Screen	98
		UT	Urine Toxicology	96
		UTCO	Urine Toxicology	131
		201	Carryover	
Organic acids, urine qualitative	Х	BGL	Biochemical Genetics	239
Organic acids, urine quantitative		BGL	Biochemical Genetics	239
Osmolality, measured	Х	C3, C3X, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check,	43
			Chemistry and TDM	
		IFS	Interfering Substances	130
			-	

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Osmolality, measured		LN2	Chemistry, Lipid,	118
(cont.)			Enzyme Cal Ver/Lin	
		LN2BV	Chemistry, Lipid,	118
			Enzyme all Beckman	
			except AU, Vitros Cal	
0 1 111		0145 01454	Ver/Lin	1//
Osmolality, urine	Х	CMP, CMP1	Clinical Microscopy	144
		CMQ	Quality Cross Check, Urinalysis	48
		LN6	Urine Chemistry Cal	120
		D000	Ver/Lin	
		POC3	POC Urine Dipstick	52
	Х	U	Urine Chemistry,	68
	^		General	
Osmometer check		1	Instrumentation	129
Osteocalcin		BGS	Bone and Growth	85
Oxazepam		DFC	Drug-Facilitated Crime	107
		DMPM	Drug Monitoring for Pain	106
		FT0	Management	10/
		FTC	Forensic Toxicology, Criminalistics	104
		OFD	Oral Fluid for Drugs of	100
		OFD	Abuse	100
		Т	Toxicology	96
		UDC	Forensic Urine Drug	99
		020	Testing, Confirmatory	
		UT	Urine Toxicology	96
Oxcarbazepine		ZE	Therapeutic Drug	60
metabolite			Monitoring, Extended	
Oxidants, urine		DAI	Urine Drug Adulterant/	98
			Integrity Testing	
Oxycodone		DFC	Drug-Facilitated Crime	107
		DMPM	Drug Monitoring for Pain	106
			Management	
		FTC	Forensic Toxicology,	104
			Criminalistics	
		OFD	Oral Fluid for Drugs of	100
			Abuse	
		T	Toxicology	96
		UDC	Forensic Urine Drug	99
		LIDO LIDOS	Testing, Confirmatory	
		UDS, UDS6	Urine Drug Screen Urine Toxicology	98
Oughamadahin	V	-		96
Oxyhemoglobin	Х	S0	Blood Oximetry	94
		SOQ	Quality Cross Check, Blood Oximetry	46
Oxylate		KSA	Kidney Stone Risk	69
ολγιαι ο		NOA	Assessment	08
Oxymorphone		DFC	Drug-Facilitated Crime	107
ON, III OI PITOTIC		21.0	Diag i domitated offile	107

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Oxymorphone (cont.)		DMPM	Drug Monitoring for Pain Management	106
		FTC	Forensic Toxicology, Criminalistics	104
		OFD	Oral Fluid for Drugs of Abuse	100
		Т	Toxicology	96
		UDC	Forensic Urine Drug Testing, Confirmatory	99
		UT	Urine Toxicology	96
p16		PM5	Immunohistochemistry TMA	268
Pancreatic amylase	X	C1, C3, C3X, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
PAPP-A		FP1B	First Trimester Maternal Screening, Free Beta	87
		FP1T	First Trimester Maternal Screening, Total hCG	87
Parainfluenza virus		ID2	Nucleic Acid Amp, Respiratory	197
	Х	IDR	Infectious Disease, Respiratory Panel	198
	Х	VR1	Virology Culture	191
	Х	VR2	Viral Antigen Detection by DFA	191
Paraprotein identification	Х	SPE	Protein Electrophoresis	76
Parasite identification	Х	BP	Blood Parasite	188
	Х	P, P3, P4, P5	Parasitology	187
		PEX	Expanded Parasitology	189
Parathyroid hormone (PTH)	Х	ING	Insulin, Gastrin, C-Peptide, PTH	86
		PTHQ	Quality Cross Check, PTH	45
Parentage/relationship testing	Х	PARF	Parentage/Relationship	227
Paroxetine		DFC	Drug-Facilitated Crime	107
		FTC	Forensic Toxicology, Criminalistics	104
		T	Toxicology	96
		UT	Urine Toxicology	96
Parvovirus B19		ID1	Nucleic Acid Amp, Viruses	196
PC02	Х	AQ, AQ2, AQ3, AQ4	Aqueous Blood Gas	92
		AQQ, AQ2Q, AQ3Q, AQ4Q	Quality Cross Check, Critical Care Aqueous Blood Gas Series	46

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
PCO2 (cont.)		POC10,	POC Competency Blood	53
		POC11	Gases	
		LN13, LN13C	Blood Gas Cal Ver/Lin	122- 123
PDGFRA		KIT	KIT/PDGFRA	252
		MTP	Multigene Tumor Panel	253
PDL1		PDL1	PDL1	267
Pentobarbital		DFC	Drug-Facilitated Crime	107
		Т	Toxicology	96
		UT	Urine Toxicology	96
Performance		PIP/PIP1,	Performance	256-
improvement program in surgical pathology		PIPW/PIPW1	Improvement Program in Surgical Pathology	257
Peripheral blood smear,		VPBS	Virtual Peripheral Blood Smear	141
pH		AFL	Amniotic Fluid Leakage	146
<u>·</u>	Х	AQ, AQ2, AQ3, AQ4	Aqueous Blood Gas	92
		AQQ, AQ2Q,	Quality Cross Check,	46
		AQ3Q, AQ4Q	Critical Care Aqueous	
			Blood Gas Series	
		FLD	Body Fluid	72
		FLDQ	Quality Cross Check, Body Fluid Chemistry	44
		GOCB	Gastric Occult Blood	148
		POC10, POC11	POC Competency Blood Gases	53
		LN13, LN13C	Blood Gas Cal Ver/Lin	122- 123
pH, gastric		GOCB	Gastric Occult Blood	148
pH, urine	Х	CMP, CMP1	Clinical Microscopy	144
<u>'</u>		CMQ	Quality Cross Check, Urinalysis	48
		DAI	Urine Drug Adulterant/ Integrity Testing	98
	Х	HCC2	Waived Combination	66
		POC3	POC Urine Dipstick Competency	52
		UDC	Forensic Urine Drug Testing, Confirmatory	99
pH meters		I	Instrumentation	129
Phencyclidine		DFC	Drug-Facilitated Crime	107
		FTC	Forensic Toxicology,	104
			Criminalistics	
		OFD	Oral Fluid for Drugs of Abuse	100
		T	Toxicology	96
		UDC	Forensic Urine Drug	99
			Testing, Confirmatory	
		UDS, UDS6	Urine Drug Screen	98
		UT	Urine Toxicology	96

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Phenethylamine		FTC	Forensic Toxicology, Criminalistics	104
		Т	Toxicology	96
		UT	Urine Toxicology	96
Pheniramine		T	Toxicology	96
- I i i i i i i i i i i i i i i i i i i		UT	Urine Toxicology	96
Phenobarbital	X	CZ, CZ2X,	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check,	43
		DEO	Chemistry and TDM	107
		DFC	Drug-Facilitated Crime	107
		FTC	Forensic Toxicology, Criminalistics	104
		LN3	TDM Cal Ver/Lin	119
		Т	Toxicology	96
		UDC	Forensic Urine Drug Testing, Confirmatory	99
		UT	Urine Toxicology	96
Phentermine		FTC	Forensic Toxicology, Criminalistics	104
		Т	Toxicology	96
		UT	Urine Toxicology	96
Phenylephrine		T	Toxicology	96
. 7		UT	Urine Toxicology	96
Phenytoin	Х	CZ, CZ2X,	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check,	43
		DEO	Chemistry and TDM	107
		DFC FTC	Drug-Facilitated Crime Forensic Toxicology,	107
		LNO	Criminalistics	110
		LN3	TDM Cal Ver/Lin	119
		SCO	Serum Carryover	131
		T	Toxicology	96
Dhamatain fusa	V	UT	Urine Toxicology	96
Phenytoin, free	X	CZ, CZ2X, CZX, Z	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
Phosphorus	Х	C3, C3X, CZ, CZX, CZ2X	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		IFS	Interfering Substances	130
		LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
		LN2BV	Chemistry, Lipid, Enzyme all Beckman except AU, Vitros Cal Ver/Lin	118

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Phosphorus, urine		LN6	Urine Chemistry Cal	120
			Ver/Lin	
	Х	U	Urine Chemistry, General	68
PIK3CA		MTP	Multigene Tumor Panel	253
Pinworm prep	Х	CMMP	Clinical Microscopy,	145
			Misc	
Pipette calibration- gravimetric		I	Instrumentation	129
Plasma cell neoplasms		PCNEO	Flow Cytometry, Plasma Cell Neoplasms	211
Plasma hemogloblin		PHG	Plasma Hemoglobin	76
Plasminogen antigen		CGE/CGEX	Coagulation, Extended	159
Plasminogen activator		CGE/CGEX	Coagulation, Extended	159
inhibitor			,	
Plasminogen activator inhibitor (PAI)-1		MGL1	Molecular Genetics	241
Platelet aggregation		PF	Platelet Function	164
Platelet antibody	Х	PS	Platelet Serology	219
detection				
Platelet calculator		TRC	Transfusion-Related Cell Count	218
Platelet count	X	FH1-FH4, FH6, FH9, FH10, FH13, FH1P-FH4P, FH6P, FH9P, FH10P, FH13P	Hematology, Auto Diff	135
		FH3Q, FH4Q, FH6Q, FH9Q	Quality Cross Check, Automated Hematology Series	47
	Х	FH15	Centrifugal Hematology	135
	Х	HE, HEP	Basic Hematology	134
		LN9	Hematology Cal Ver/Lin	121
Platelet count (platelet- rich plasma)	Х	TRC	Transfusion-Related Cell Count	218
Platelet crossmatch		PS	Platelet Serology	219
Platelet count		EHE1	Expanded Virtual	142
(estimated)			Peripheral Blood Smear	
		VPBS	Virtual Peripheral Blood	141
			Smear	
Platelet function		PF1	Platelet Function	164
Platelet mapping		PLTM	Platelet Mapping	167
Plesiomonas shigelloides		GIP	Gastrointestinal Panel	199
PML/RARA		MH02, MH03	Molecular Hematologic Oncology	254
		MRD2	Minimal Residual Disease	254
PNA FISH-		PNA1	PNA FISH for	177
Staphylococcus			Staphylococcus	
PNA FISH-yeast		PNA2	PNA FISH for Yeast	177

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Pneumocystis detection		PCP1	Pneumocystis jiroveci, Calcofluor White Stain	186
		PCP2	Pneumocystis jiroveci, DFA Stain	186
		PCP4	Pneumocystis jiroveci, GMS Stain	186
PNH immunophenotype		PNH	Paroxysmal Nocturnal Hemoglobinuria, RBC	211
P02	Х	AQ, AQ2, AQ3, AQ4	Aqueous Blood Gas	92
		AQQ, AQ2Q, AQ3Q, AQ4Q	Quality Cross Check, Critical Care Aqueous Blood Gas Series	46
		LN13, LN13C	Blood Gas Cal Ver/Lin	122- 123
		POC10, POC11	POC Competency Blood Gases	53
Porphobilinogen, urine		UPBG	Porphobilinogen, Urine	70
Postanalytical DNA sequencing		SEC	DNA Sequencing Count	242
Postvasectomy sperm count	Х	PV	Postvasectomy Sperm Count	154
Potassium	Х	AQ, AQ2, AQ3, AQ4	Aqueous Blood Gas	92
		AQQ, AQ2Q, AQ3Q, AQ4Q	Quality Cross Check, Critical Care Aqueous Blood Gas Series	46
	X	C1, C3, C3X, C4, CZ, CZX, CZ2X	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		FLD2	Body Fluid Chemistry 2	73
		IFS	Interfering Substances	130
		LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
		LN2BV	Chemistry, Lipid, Enzyme all Beckman except AU, Vitros Cal Ver/Lin	118
		LN13C	Blood Gas Cal Ver/Lin	122- 123
		POC10, POC11	POC Competency Blood Gases	53
Potassium, urine		LN6	Urine Chemistry Cal Ver/Lin	120
	Х	U	Urine Chemistry, General	68
Potassium, vitreous fluid		VF	Vitreous Fluid, Post- mortem	101

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
PRA		MX1B, MX1C, MX1E, MXB, MXC	HLA Analysis, Class I	230- 231
		MX2B, MX2C, MX2E, MXB, MXC	HLA Analysis, Class II	230- 231
Prader-Willi/Angelman syndrome	Х	MGL1	Molecular Genetics	241
Prealbumin (transthyretin)	Х	C3, C3X, CZ, CZX, CZ2X	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
	X	S2, S4	Immunology, Special	203
Pregabalin		DMPM	Drug Monitoring for Pain Management	106
		Т	Toxicology	96
		UT	Urine Toxicology	96
		ZE	Therapeutic Drug	60
			Monitoring, Extended	
Prekallikrein		CGE/CGEX	Coagulation, Extended	159
Predictive markers by	Х	HER2	HER2 by	269
immunohistochemistry			Immunohistochemistry	
		GHER2	Gastric HER2	269
		PM1	CD117 by Immunohistochemistry	268
	Х	PM2	ER, PgR by Immunohistochemistry	269
		PM3	CD20 by Immunohistochemistry	268
		PM5	Immunohistochemistry TMA	268
Primidone	Х	CZ, CZX, CZ2X, Z	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		LN3	TDM Cal Ver/Lin	119
Pro B-type natriuretic peptides		BNP	B-Type Natriuretic Peptides, 2 Chall	61
	Х	BNP5	B-Type Natriuretic Peptides, 5 Chall	61
		BNPQ	Quality Cross Check, B-Type Natriuretic Peptides	43
Procainamide	Х	CZ, CZX, CZ2X, Z	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		LN3	TDM Cal Ver/Lin	119
Procalcitonin		LN41	Procalcitonin Cal Ver/ Lin	128
	Х	PCT	Procalcitonin	77

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Progesterone		LN8	Reproductive Endocrinology Cal Ver/ Lin	121
	Х	Y/YY	Ligand Assay, Special	84
Progesterone receptors by immunohistochemistry	Х	PM2	ER, PgR by Immunohistochemistry	269
Prolactin		LN8	Reproductive Endocrinology Cal Ver/ Lin	121
	Х	Y/YY	Ligand Assay, Special	84
Propoxyphene		DFC	Drug-Facilitated Crime	107
		DMPM	Drug Monitoring for Pain Management	106
		FTC	Forensic Toxicology, Criminalistics	104
		T	Toxicology	96
		UDC	Forensic Urine Drug Testing, Confirmatory	99
		UDS, UDS6	Urine Drug Screen	98
		UT	Urine Toxicology	96
Propranolol		Т	Toxicology	96
		UT	Urine Toxicology	96
Prostate-specific antigen (PSA)	Х	K, KK, K2	Ligand Assay, General	82
		LN23	PSA Cal Ver/Lin	125
Prostate-specific antigen, complexed (cPSA)	Х	K/KK	Ligand Assay, General	82
Prostate-specific antigen, free (PSA, free)	Х	K/KK	Ligand Assay, General	82
Prostatic acid phosphatase (PAP)	Х	K/KK	Ligand Assay, General	82
Protein electrophoresis, serum, interpretation		SPE	Protein Electrophoresis	76
Protein C		CGE/CGEX	Coagulation, Extended	159
		CGS2	Coag Special, Series 2	160- 161
		LN35	Thrombophilia Cal Ver/ Lin	127
Protein S		CGE/CGEX	Coagulation, Extended	159
		CGS2	Coag Special, Series 2	160- 161
Protein, total	Х	C1, C3, C3X, CZ, CZX, CZ2X	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		FLD	Body Fluid	72
		FLDQ	Quality Cross Check, Body Fluid Chemistry	44
		IFS	Interfering Substances	130

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Protein, total (cont.)		LN2	Chemistry, Lipid,	118
			Enzyme Cal Ver/Lin	
		LN2BV	Chemistry, Lipid,	118
			Enzyme all Beckman	
			except AU, Vitros Cal	
		CDE	Ver/Lin	76
		SPE	Lipoprotein and Protein Electrophoresis	76
Protein, CSF	X	M, OLI	CSF Chemistry and	74
1 1000111, 001		IVI, OLI	Oligoclonal Bands	, ,
Protein, urine	Х	CMP, CMP1	Clinical Microscopy	144
		CMQ	Quality Cross Check,	48
			Urinalysis	
		DSC	Dipstick Confirmatory	147
	Х	HCC2	Waived Combination	66
		LN6	Urine Chemistry Cal	120
			Ver/Lin	
		POC3	POC Urine Dipstick	52
			Competency	
	Х	U	Urine Chemistry,	68
Prothrombin mutation	V	MGL1	General Molecular Genetics	2/1
Prothrombin mutation	X	TPM	Thrombophilia	241
	^	IPIVI	Mutations	244
Prothrombin fragment		CGE/CGEX	Coagulation, Extended	159
1.2		ode, odex	oodgatation, Extended	100
Prothrombin time	Х	CGB	Basic Coagulation	158
	Х	CGL	Coagulation, Limited	158
		CGLQ	Quality Cross Check,	49
			Coagulation, Limited	
		CGS1	Coag Special, Series 1	160-
				161
		CGS4	Coag Special, Series 4	160-
				161
		DBGN	Anticoagulant	161
		END/	Monitoring, Dabigatran	404
		FNPX	Anticoagulant Monitoring,	161
			Fondaparinux	
		POC6	POC PT/INR, CoaguChek	52
			XS Plus	"-
		RVBN	Anticoagulant	161
			Monitoring Rivaroxaban	
	Χ	WP3, WP4,	Whole Blood	166
		WP6, WP9	Coagulation	
Prothrombin time, dilute		CGE/CGEX	Coagulation, Extended	159
Provider-performed		CMMP	Clinical Microscopy,	145
microscopy		DIA SILI	Misc	4.55
PRU test		PIA, PIAX	Drug-Specific Platelet Aggregation	165
Pseudocholinesterase	Х	C7	Pseudocholinesterase	77

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Pseudoephedrine		FTC	Forensic Toxicology, Criminalistics	104
		T	Toxicology	96
		UT	Urine Toxicology	96
PTEN		GLI	Glioma	253
Pyridinoline (PYD)		BU	Bone and Mineral, Urine	85
Q-MONITORS		QM1	Quality Management Tools	39
Q-PROBES		QP181	Quality Management Tools	25
		QP182	Quality Management Tools	26
		QP183	Quality Management Tools	27
		QP184	Quality Management Tools	28
Q-TRACKS		QT1-5	Quality Management	31-
			Tools	32,
				37
		QT7, 8, 10	Quality Management	33-
			Tools	34
		QT15-17	Quality Management	35-
			Tools	36
Quetiapine		Т	Toxicology	96
		UT	Urine Toxicology	96
Quinidine	X	CZ, CZX,	Chemistry and TDM	56-
		CZ2X, Z		58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		LN3	TDM Cal Ver/Lin	119
		T	Toxicology	96
		UT	Urine Toxicology	96
Quinine		T	Toxicology	96
		UT	Urine Toxicology	96
Ranitidine		T	Toxicology	96
		UT	Urine Toxicology	96
Rapamycin (sirolimus)	Х	CS	Immunosuppressive Drugs	59
Rapid group A strep	Х	D	Bacteriology	170
	Х	D4	Bacteriology, Limited	173
	Х	D6	Rapid Group A Strep	175
	Х	D9	Rapid Group A Strep, Waived	175
	Х	MC1	Microbiology Combination with GC	173
	Х	MC2	Microbiology Combination	173
	Х	MC4	Urine Colony Count Combination	174

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Rapid group A strep	Χ	MC5	Throat Culture/Rapid	174
(cont.)			Strep	
RBC count		ABF1, ABF2, ABF3	Automated Body Fluid	146
	X	FH1-FH4, FH6, FH9, FH10, FH13, FH1P-FH4P, FH6P, FH9P, FH10P, FH13P	Hematology, Auto Diff	135
		FH3Q, FH4Q, FH6Q, FH9Q	Quality Cross Check, Automated Hematology Series	47
	X	HE, HEP	Basic Hematology	134
		LN9	Hematology Cal Ver/Lin	121
RBC count, automated, urine (quantitative)		UAA, UAA1	Automated Urinalysis	147
RBC automated count, fluid		ABF1, ABF2, ABF3	Automated Body Fluid	146
RBC manual count, fluid	Х	HFC, HFCI	Hemocytometer Fluid Count	148- 149
RBC folate	Х	FOL	RBC Folate	88
RBC morphology		EHE1	Expanded Virtual Peripheral Blood Smear	142
		VPBS	Virtual Peripheral Blood Smear	141
RDW		FH1-FH4, FH6, FH9, FH10, FH13, FH1P-FH4P, FH6P, FH9P, FH10P, FH13P	Hematology, Auto Diff	135
		FH3Q, FH4Q, FH6Q, FH9Q	Quality Cross Check, Automated Hematology Series	47
		HE, HEP	Basic Hematology	134
Red blood cell antigen detection		J, J1	Transfusion Medicine	214
Red blood cell antigen genotyping		RAG	Red Blood Cell Antigen Genotyping	216
Red blood cell antigen typing		RBCAT	Red Blood Cell Antigen Typing	217
Reducing substance, urine	Х	CMP, CMP1	Clinical Microscopy	144
		CMQ	Quality Cross Check, Urinalysis	48
	Х	HCC2	Waived Combination	66
		POC3	POC Urine Dipstick Competency	52
Refractometer check		I	Instrumentation	129

Analyte/Procedure	LAP	Program	Description	Pg
	ENR	Code		
Renin	Χ	RAP	Renin and Aldosterone	89
Reptilase time		CGE/CGEX	Coagulation, Extended	159
Respiratory syncytial		ID2	Nucleic Acid Amp,	197
virus (RSV)			Respiratory	
	Χ	ID3	Influenza A, Influenza B,	197
			RSV by NAA	
	Χ	IDR	Infectious Disease,	198
			Respiratory Panel	
	Χ	VR1	Virology Culture	191
	Х	VR2	Viral Antigen Detection by DFA	191
	Χ	VR4	Virology Antigen	192
			Detection by EIA and	
			Latex	
Reticulocyte count, absolute	Х	RT, RT2, RT3, RT4	Reticulocyte	139
		RTQ, RT2Q,	Quality Cross Check,	47
		RT3Q, RT4Q	Reticulocyte	
Reticulocyte count, percent		LN18, LN19	Reticulocyte Cal Ver/Lin	124
	Χ	RT, RT2, RT3, RT4	Reticulocyte	139
		RTQ, RT2Q,	Quality Cross Check,	47
		RT3Q, RT4Q	Reticulocyte	
RETT syndrome	Χ	RETT	RETT Syndrome	243
			Genotyping	
RhD	Χ	MGL2	Molecular Genetics	241
RhD typing	Χ	J, J1	Transfusion Medicine	214
	Х	JAT	Transfusion Medicine, Automated	215
		JATE1	Transfusion Medicine,	215
			Automated, Educational	
		TMCA	Transfusion Medicine,	219
			Competency	
			Assessment	
Rheumatoid factor	Χ	IL, RF/RFX	Immunology	202
Rhinovirus		ID2	Nucleic Acid Amp,	197
			Respiratory	
	Χ	IDR	Infectious Disease,	198
			Respiratory Panel	
RNA sequencing		RNA	RNA Sequencing	252
Rotavirus		GIP	Gastrointestinal Panel	199
		SP, SPN	Stool Pathogens	180
	Х	VR4	Viral Antigen Detection	192
		100	by EIA and Latex	
RSV		ID2	Nucleic Acid Amp,	197
		IDO	Respiratory	107
	Х	ID3	Influenza A, Influenza B,	197
	V	IDD	RSV by NAA	100
	Х	IDR	Infectious Disease,	198
	V	VD1	Respiratory Panel	101
	X	VR1	Virology Culture	191

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
RSV (cont.)	Х	VR2	Viral Antigen Detection by DFA	191
	X	VR4	Viral Antigen Detection by EIA and Latex	192
Rubella antibody, IgG	Х	IL, RUB/ RUBX	Immunology	202
Rubeola antibody (English measles)	Х	VR3	Antibody Detection- Infectious Disease Serology	200
Rufinamide		ZE	Therapeutic Drug Monitoring, Extended	60
Rupture of fetal membranes		ROM1	Rupture of Fetal Membranes	150
Russell's viper venom time, dilute		CGE/CGEX	Coagulation, Extended	159
Salicylate	Х	CZ, CZX, CZ2X, Z	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		FTC	Forensic Toxicology, Criminalistics	104
		LN3	TDM Cal Ver/Lin	119
	X	SDS	Serum Drug Screen	101
		T	Toxicology	96
		UT	Urine Toxicology	96
Salmonella		GIP	Gastrointestinal Panel	199
Sapovirus (I, II, IV, V)		GIP	Gastrointestinal Panel	199
Sarcoma by FISH		CYK	Fluorescence In Situ Hybridization	237
Sarcoma translocation		SARC	Sarcoma Translocation	251
Scl-70 (anti-DNA		RDS	Rheumatic Disease	207
topoisomerase)			Special	
Scopolamine		DFC	Drug-Facilitated Crime	107
Secobarbital		DFC	Drug-Facilitated Crime	107
		FTC	Forensic Toxicology, Criminalistics	104
		UDC	Forensic Urine Drug Testing, Confirmatory	99
Selenium	X	R	Trace Metals	78
Selenium, urine		TMU	Trace Metals, Urine	103
Selenium, whole blood		TMWB	Trace Metals, Whole Blood	103
Semen analysis	X	ASA, SC, SV, PV	Semen Analysis	154
		SC1, SM	Semen Analysis	154
		SMCD, SM1CD, SM2CD	Semen Analysis, CD- ROM	154
SERPINA1 genotyping	Х	AAT	Alpah-1 Antitrypsin Genotyping	239
Sertraline		DFC	Drug-Facilitated Crime	107

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Sertraline (cont.)		FTC	Forensic Toxicology, Criminalistics	104
		T	Toxicology	96
		UT	Urine Toxicology	96
Serum free light chains		SFLC	Serum Free Light Chains	208
Sex hormone-binding		ABS	Testosterone and	111
globulin (SHBG)		7120	Estradiol Accuracy	
<u> </u>	Х	DY	Ligand Assay, Special	84
Shiga toxin		SP	Stool Pathogens-Rapid	180
0			and Molecular	
		ST	Shiga Toxin	181
Shiga-like toxin producing <i>E. coli</i> (STEC)		GIP	Gastrointestinal Panel	199
Shigella		GIP	Gastrointestinal Panel	199
Sickle cell screen, qualitative	Х	HG	Hemoglobinopathy	139
4	Х	SCS	Sickle Cell Screen	140
Sirolimus (Rapamycin)	Х	CS	Immunosuppressive Drugs	59
SLC01B1		PGX	Pharmacogenetics	243
Sodium	Х	AQ, AQ2, AQ3, AQ4	Aqueous Blood Gas	92
		AQQ, AQ2Q, AQ3Q, AQ4Q	Quality Cross Check, Critical Care Aqueous Blood Gas Series	46
	X	C1, C3, C3X, C4, CZ, CZ2X, CZX	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		FLD2	Body Fluid Chemistry 2	73
		IFS	Interfering Substances	130
		LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
		LN2BV	Chemistry, Lipid, Enzyme all Beckman except AU, Vitros Cal Ver/Lin	118
		LN13C	Blood Gas Cal Ver/Lin	122- 123
		POC10, POC11	POC Competency Blood Gases	53
Sodium, urine		LN6	Urine Chemistry Cal Ver/Lin	120
	Х	U	Urine Chemistry, General	68
Sodium, vitreous fluid		VF	Vitreous Fluid, Post- mortem	101
Soluble transferrin receptor		STFR	Soluble Transferrin Receptor	80
Somatomedin C (IGF-1)	Х	Y, YY	Ligand Assay, Special	84

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
S0X10		PM5	Immunohistochemistry TMA	268
Specific gravity	Х	CMP, CMP1	Clinical Microscopy	144
		CMQ	Quality Cross Check,	48
			Urinalysis	
		DAI	Urine Drug Adulterant/ Integrity Testing	98
	Х	HCC2	Waived Combination	66
		POC3	POC Urine Dipstick Competency	52
		UDC	Forensic Urine Drug Testing, Confirmatory	99
Spectrophotometer linearity		1	Instrumentation	129
Sperm count	Х	SMCD	Semen Analysis, CD- ROM	154
Sperm count, automated		SC1	Semen Analysis	154
Sperm count, manual	Х	SC	Semen Analysis	154
	Х	PV	Postvasectomy Sperm Count	154
Sperm morphology		SM	Semen Analysis	154
		SM1CD	Semen Analysis, CD- ROM	154
Sperm motility		SMCD	Semen Analysis, CD- ROM	154
Sperm viability		SM2CD	Semen Analysis, CD- ROM	154
	Х	SV	Semen Analysis	154
Spinal fluid meningitis panel	Х	D	Bacteriology	170
Spinal muscular atrophy	Х	MGL2	Molecular Genetics	241
Spinocerebellar ataxia	Х	MGL2	Molecular Genetics	241
Split fats		FCFS	Fecal Fat	75
Staphylococcus aureus-		BCS1	Blood Culture	177
blood culture			Staphylococcus aureus	
STEC (Shiga-like toxin producing <i>E. coli</i>)		GIP	Gastrointestinal Panel	199
Strep screen		POC4	POC/Waived Strep Screen Competency	52
Streptococcus agalactiae	Х	D8	Group B Strep	176
		IDME	Meningitis/Encephalitis Panel	198
Streptococcus		IDME	Meningitis/Encephalitis	198
pneumoniae			Panel	
		SBAS	S. pneumoniae Ag Detection	176
Streptococcus pyogenes	Х	D	Bacteriology	170
	Х	D1, D7	Throat, Urine Cultures	172
	Х	D4	Bacteriology, Ltd	173
	Х	D6	Rapid Group A Strep	175

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Streptococcus pyogenes (cont.)	Х	D9	Rapid Group A Strep, Waived	175
	Х	MC1	Microbiology Combination with GC	173
	Х	MC2	Microbiology Combination	173
	Х	MC4	Urine Colony Count Combination	174
	Х	MC5	Throat Culture/Rapid Strep	174
Strychnine		Т	Toxicology	96
		UT	Urine Toxicology	96
Sulfate		KSA	Kidney Stone Risk Assessment	69
Sulfosalicylic acid (SSA)		DSC	Dipstick Confirmatory	147
Surgical pathology		DPATH/ DPATH1	Online Digital Slide Program	259
		PIP/PIP1, PIPW/PIPW1	Performance Improvement Program in Surgical Pathology	256- 257
		PCSP, PCSP1	Practicum in Cancer Surgical Pathology	262
		VBP/VBP1	Online Virtual Biopsies Program	258
Synthetic cannabinoid/ designer drugs		SCDD	Synthetic Cannabinoid/ Designer Drugs	105
Syphilis	Χ	G	Syphilis Serology	208
T3, free (triiodothyronine)		ABTH	Harmonized Thyroid	112
	X	C1, C3, C3X, CZ, CZX, CZ2X	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
	Χ	K/KK	Ligand Assay, General	82
T3, total (triiodothyronine)		ABTH	Harmonized Thyroid	112
	X	C1, C3, C3X, CZ, CZX, CZ2X	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
	Χ	K/KK	Ligand Assay, General	82
		LN5	Ligand Assay Cal Ver/Lin	119- 120
		LN5S	Ligand Assay, Siemens Cal Ver/Lin	119- 120
T3, uptake and related tests	Х	C1, C3, C3X, CZ, CZX, CZ2X	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
	Χ	K/KK	Ligand Assay, General	82

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
T4, free (thyroxine, free)		ABTH	Harmonized Thyroid	112
	X	C1, C3, C3X, CZ, CZX, CZ2X	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
	Х	K/KK	Ligand Assay, General	82
T4, total (thyroxine, total)		ABTH	Harmonized Thyroid	112
	X	C1, C3, C3X, CZ, CZX, CZ2X	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
	Х	K/KK	Ligand Assay, General	82
		LN5	Ligand Assay Cal Ver/Lin	119- 120
		LN5S	Ligand Assay, Siemens Cal Ver/Lin	119- 120
Tacrolimus	X	CS	Immunosuppressive Drugs	59
		LN31	Immunosuppressive Drugs Cal Ver/Lin	126
Tay Sachs	Χ	MGL4	Molecular Genetics	241
tCO ₂		AQ, AQ2, AQ3, AQ4	Aqueous Blood Gas	92
		AQQ, AQ2Q, AQ3Q, AQ4Q	Quality Cross Check, Critical Care Aqueous Blood Gas Series	46
		POC10, POC11	POC Competency Blood Gases	53
Temazepam		DFC	Drug-Facilitated Crime	107
		DMPM	Drug Monitoring for Pain Management	106
		FTC	Forensic Toxicology, Criminalistics	104
		OFD	Oral Fluid for Drugs of Abuse	100
		T	Toxicology	96
		UDC	Forensic Urine Drug Testing, Confirmatory	99
		UT	Urine Toxicology	96
Teriflunomide		ZE	Therapeutic Drug Monitoring, Extended	60
Testosterone		ABS	Accuracy-Based Testosterone and Estradiol	111
		LN8	Reproductive Endocrinology Cal Ver/ Lin	121
	Χ	Y/YY	Ligand Assay, Special	84
Testosterone, bioavailable		ABS	Testosterone and Estradiol Accuracy	111

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Testosterone,		DY	Ligand Assay, Special	84
bioavailable (cont.)				
Testosterone, free		ABS	Testosterone and	111
	.,		Estradiol Accuracy	
T. I. I. I. I. I. I.	Х	DY	Ligand Assay, Special	84
Tetrahydrozoline		DFC	Drug-Facilitated Crime	107
Thallium, urine		TMU	Trace Metals, Urine	103
Thallium, whole blood		TMWB	Trace Metals, Whole Blood	103
Theophylline	Χ	CZ, CZX, CZ2X, Z	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check,	43
		024	Chemistry and TDM	
		LN3	TDM Cal Ver/Lin	119
Throat culture	Х	D1, D7	Throat, Urine Cultures	172
	Х	D4	Bacteriology, Ltd	173
	Х	MC1	Microbiology	173
			Combination with GC	
	Х	MC2	Microbiology	173
			Combination	
	X	MC4	Urine Colony Count	174
			Combination	
	X	MC5	Throat Culture/Rapid	174
Thrombin time		CGE/CGEX	Strep Coagulation, Extended	159
- Infombin time		CGS4	Coag Special, Series 4	160-
		0004	Coag opecial, ceries 4	161
		DBGN	Dabigatran	161
Thrombin-antithrombin		CGE/CGEX	Coagulation, Extended	159
Thromboelastogram		TEG	Viscoelastometry	164
Thrombophilia mutations	Х	TPM	Thrombophilia Mutations	244
Thyroglobulin	Х	TM/TMX	Tumor Markers	89
Thyroid-stimulating		ABS	Accuracy-Based	111
hormone (TSH)			Testosterone and Estradiol	
		ABTH	Harmonized Thyroid	112
	Х	C1, C3, C3X,	Chemistry and TDM	56-
		CZ, CZX, CZ2X	,	58
		CZQ	Quality Cross Check,	43
	V	K/KK	Chemistry and TDM	02
	X	LN5	Ligand Assay, General Ligand Assay Cal Ver/Lin	82 119–
		LINU	Ligariu Assay Cat Ver/LIII	120
		LN5S	Ligand Assay, Siemens	119-
			Cal Ver/Lin	120
Thyroxine, free		ABTH	Harmonized Thyroid	112
	X	C1, C3, C3X, CZ, CZX, CZ2X	Chemistry and TDM	56- 58

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Thyroxine, free (cont.)		CZQ	Quality Cross Check,	43
			Chemistry and TDM	
	Х	K/KK	Ligand Assay, General	82
Thyroxine, total		ABTH	Harmonized Thyroid	112
	X	C1, C3, C3X, CZ, CZX, CZ2X	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
	Х	K/KK	Ligand Assay, General	82
		LN5	Ligand Assay Cal Ver/Lin	119- 120
		LN5S	Ligand Assay, Siemens	119-
			Cal Ver/Lin	120
Tick identification		TMO	Ticks, Mites, and Other Arthropods	189
Tissue parasite identification	Х	BP	Blood Parasite	188
	Х	Р	Parasitology	187
Tissue plasminogen activator		CGE/CGEX	Coagulation, Extended	159
Tobramycin	Х	CZ, CZX, CZ2X, Z	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		LN3	TDM Cal Ver/Lin	119
Topiramate		Т	Toxicology	96
		UT	Urine Toxicology	96
		ZE	Therapeutic Drug Monitoring, Extended	60
Total bile acids		TBLA	Total Bile Acid	78
Total bilirubin	X	C1, C3, C3X, CZ, CZX, C4, CZ2X	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		FLD2	Body Fluid Chemistry 2	73
		IFS	Interfering Substances	130
		LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
		LN2BV	Chemistry, Lipid, Enzyme all Beckman except AU, Vitros Cal Ver/Lin	118
	Х	NB, NB2	Neonatal Bilirubin	65
Total bilirubin, urine	Х	CMP, CMP1	Clinical Microscopy	144
	Х	HCC2	Waived Combination	66
		DSC	Dipstick Confirmatory	147
Total free fatty acids		FCFS	Fecal Fat	75
Total hCG	Х	FP1T	First Trimester Maternal Screening, Total hCG	87

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Total hemolytic		CH50	Total Hemolytic	208
complement			Complement	
Total iron binding	Χ	C3, C3X, CZ	Chemistry and TDM	56-
capacity, measured and		CZX, CZ2X		58
% saturation				
		CZQ	Quality Cross Check,	43
Total nitragen uring		U	Chemistry and TDM Urine Chemistry,	68
Total nitrogen, urine		0	General	00
Total nucleated cells		CBT	Cord Blood Testing	221
		SCP	Stem Cell Processing	221
Total nucleated cells		HFC/HFCI	Hemocytometer Fluid	148-
manual differential count		6, 6.	Count	149
(body fluid)				
		VBF	Virtual Body Fluid	146
Total nucleated red cells		CBT	Cord Blood Testing	221
		SCP	Stem Cell Processing	221
Total protein	Χ	C1, C3, C3X,	Chemistry and TDM	56-
		CZ, CZX,		58
		CZ2X		
		CZQ	Quality Cross Check,	43
		EL D	Chemistry and TDM	70
		FLDO	Body Fluid	72
		FLDQ	Quality Cross Check,	44
		IFS	Body Fluid Chemistry Interfering Substances	130
		LN2	Chemistry, Lipid,	118
		LINZ	Enzyme Cal Ver/Lin	110
		LN2BV	Chemistry, Lipid,	118
			Enzyme all Beckman	
			except AU, Vitros Cal	
			Ver/Lin	
		SPE	Protein Electrophoresis	76
Total protein, CSF	Χ	M, OLI	CSF Chemistry and	74
			Oligoclonal Bands	
Total protein, urine		CMP, CMP1	Clinical Microscopy	144
		CMQ	Quality Cross Check,	48
	.,		Urinalysis	
	Х	HCC2	Waived Combination	66
		LN6	Urine Chemistry Cal Ver/Lin	120
	Χ	U	Urine Chemistry,	68
			General	
Total tricyclics	Χ	SDS	Serum Drug Screen	101
	Χ	ZT	TDM, Special	60
Touch imprint/crush prep		TICP, TICP1	Touch Imprint/Crush Prep	275
Toxicology, serum,	Χ	SDS	Serum Drug Screen	101
qualitative				
	Χ	Т	Toxicology	96
Toxicology, urine,	Χ	DMPM	Drug Monitoring for Pain	106
qualitative			Management	

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Toxicology, urine, qualitative (cont.)	Χ	Т	Toxicology	96
Delayed until 20	19	TQP	Toxicology Quality Program	108
	Х	UDS, UDS6	Urine Drug Screen	98
	Χ	UT	Urine Toxicology	96
Toxicology, urine, qualitative/quantitative	Х	DMPM	Drug Monitoring for Pain Management	106
	Х	UDC	Forensic Urine Drug Testing, Confirmatory	99
Toxoplasma gondii	Х	VR3	Antibody Detection- Infectious Disease Serology	200
TPMT		PGX3	Pharmacogenetics	243
Tramadol		DFC	Drug-Facilitated Crime	107
		DMPM	Drug Monitoring for Pain Management	106
		FTC	Forensic Toxicology, Criminalistics	104
		T	Toxicology	96
		UT	Urine Toxicology	96
Transferrin	Х	C3, C3X, CZ, CZX, CZ2X	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		LN7	Immunology Cal Ver/Lin	121
	Х	S2, S4	Immunology, Special	203
Transfusion medicine		ETME1	Expanded Transfusion Medicine Exercises	223
		EXM, EXM2	Electronic Crossmatch	215- 216
	Х	J,J1	Transfusion Medicine	214
	Х	JAT	Transfusion Medicine, Automated	215
		JATE1	Transfusion Medicine, Automated	215
		JE1	Transfusion Medicine,	214
		TMCA	Transfusion Medicine, Competency Assessment	219
		TMCAD	Transfusion Medicine, Competency Assessment	219
		TMCAE	Transfusion Medicine, Competency Assessment	220
		TMCAF	Transfusion Medicine, Competency Assessment	220
	Х	TRC	Transfusion-Related Cell Count	218

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Trazodone		FTC	Forensic Toxicology,	104
			Criminalistics	
		Т	Toxicology	96
		UT	Urine Toxicology	96
Treponema pallidum	Χ	G	Syphilis Serology	208
Trichomonas vaginalis		TVAG	Trichomonas vaginalis, Molecular	189
-	Х	VS, VS1	Vaginitis Screen	181
Triovalia group	^	T T	Toxicology	96
Tricyclic group		UDS, UDS6		98
		UT	Urine Drug Screen	
Trianglian Askal	V	-	Urine Toxicology	96
Tricyclics, total	X	SDS	Serum Drug Screen	101
	Х	ZT	TDM, Special	60
Triglycerides		ABL	Accuracy-Based Lipid	110
	X	C1, C3, C3X, C4, CZ, CZX, CZ2X	Chemistry and TDM	56– 58
		CZQ	Quality Cross Check,	43
			Chemistry and TDM	
		FCFS	Fecal Fat	75
		FLD	Body Fluid	72
		FLDQ	Quality Cross Check, Body Fluid Chemistry	44
	Χ	LCW	Ltd Chem, Waived	64
		LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
		LN2BV	Chemistry, Lipid, Enzyme all Beckman except AU, Vitros Cal Ver/Lin	118
Triiodothyronine (T3)		ABTH	Harmonized Thyroid	112
	X	C1, C3, C3X, CZ, CZX, CZ2X	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
	Χ	K/KK	Ligand Assay, General	82
		LN5	Ligand Assay Cal Ver/Lin	119- 120
		LN5S	Ligand Assay, Siemens Cal Ver/Lin	119- 120
Triiodothyronine (T3), free		ABTH	Harmonized Thyroid	112
	Х	C1, C3, C3X, CZ, CZX, CZ2X	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
	Х	K/KK	Ligand Assay, General	82
 Trimipramine		T	Toxicology	96
prammo		UT	Urine Toxicology	96

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Troponin I, plasma	Х	PCARM, PCARMX	Plasma Cardiac Markers	65
		POC12	Competency Plasma Cardiac Markers	53
Troponin I, serum	Х	CRT, CRTI	Cardiac Markers	62
		LN25	Troponin I Cal Ver/Lin	125
Troponin T, serum		LN27	Troponin T Cal Ver/Lin	125
		TNT	Troponin T	62
-	Х	TNT5	Troponin T, 5 Challenge	62
Tumor necrosis factor (TNF)-alpha		CTKN	Cytokines	206
UGT1A1		PGX3	Pharmacogenetics	243
Unsaturated iron binding capacity, measured	Х	C3, C3X, CZ, CZX, CZ2X	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
Urea nitrogen	Х	AQ2, AQ4	Aqueous Blood Gas	92
		AQ2Q, AQ4Q	Quality Cross Check, Critical Care Aqueous Blood Gas Series	46
	X	C1, C3, C3X, C4, CZ, CZX, CZ2X	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		FLDQ	Quality Cross Check, Body Fluid Chemistry	44
		IFS	Interfering Substances	130
		LN2	Chemistry, Lipid, Enzyme Cal Ver/Lin	118
		LN2BV	Chemistry, Lipid, Enzyme all Beckman except AU, Vitros Cal Ver/Lin	118
Urea nitrogen, urine		LN6	Urine Chemistry Cal Ver/Lin	120
	Х	U	Urine Chemistry, General	68
Urea nitrogen, vitreous fluid		VF	Vitreous Fluid, Post- mortem	101
Urease	Х	RUR	Rapid Urease	180
Uric acid	Х	C1, C3, C3X, C4, CZ, CZX, CZ2X	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		FLD2	Body Fluid Chemistry 2	73
		IFS	Interfering Substances	130
		LN2	Chemistry, Lipid,	118
			Enzyme Cal Ver/Lin	

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Uric acid (cont.)		LN2BV	Chemistry, Lipid, Enzyme all Beckman except AU, Vitros Cal Ver/Lin	118
Uric acid, urine		LN6	Urine Chemistry Cal Ver/Lin	120
	X	U	Urine Chemistry, General	68
Urine albumin		LN20	Urine albumin Cal Ver/ Lin	124
	Х	U	Urine Chemistry, General	68
	Х	UMC	Urine Albumin Creatinine	151
Urine albumin: creatinine ratio		ABU	Accuracy-Based Urine	111
		U	Urine Chemistry, General	68
		UMC	Urine Albumin Creatinine	151
Urine colony count		MC3	Urine Colony Count	174
		MC4	Urine Colony Count Combination	174
Urine crystals identification		URC	Crystals	147
Urine crystals, semiquantitative		UAA	Automated Urinalysis	147
Urine culture	Х	D2, D7	Throat, Urine Cultures	172
	Х	D4	Bacteriology, Limited	173
	Х	MC1	Microbiology Combination with GC	173
	Х	MC2	Microbiology Combination	173
		MC3	Urine Colony Count	174
	Х	MC4	Urine Colony Count Combination	174
	Х	MC5	Throat Culture/Rapid Strep	174
Urine dipstick	Х	CMP, CMP1	Clinical Microscopy	144
		CMQ	Quality Cross Check, Urinalysis	48
	Х	HCC2	Waived Combination	66
		POC3	POC/Waived Urine	52
			Dipstick Competency	
Urine drug screen	Х	DMPM	Drug Monitoring for Pain Management	106
	Χ	UDS, UDS6	Urine Drug Screen	98
Urine hCG, qualitative	Χ	UHCG	Urine hCG	150
Urine sediment, color photographs	Х	CMP, CMP1, CMMP	Clinical Microscopy	144- 145
Urobilinogen	X	CMP, CMP1	Clinical Microscopy	144

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Urobilinogen (cont.)		CMQ	Quality Cross Check,	48
			Urinalysis	
	Х			66
		POC3	POC Urine Dipstick	52
Uranarnhyrin	X	N/NX	Competency Urine Chemistry, Special	69
Uroporphyrin Urothelial carcinoma by	X	CYI	Fluorescence In Situ	236
FISH, hybridization and	^	OTT	Hybridization, Urothelial	230
interpretation on-site			Carcinoma	
Vaginal wet preparations	Х	CMMP	Clinical Microscopy, Misc	145
Vaginitis screen		BV	Bacterial Vaginosis	181
	Х	VS	BD Affirm VP III Antigen Detection	181
	Χ	VS1	Genzyme OSOM	181
			Trichomonas	
		VS2	Vaginitis Screen, Virtual Gram Stain	182
Valproic acid	X	CZ, CZX, CZ2X, Z	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
		DFC	Drug-Facilitated Crime	107
		LN3	TDM Cal Ver/Lin	119
		T	Toxicology	96
		UT	Urine Toxicology	96
Valproic acid, free	Х	CZ, CZX, CZ2X, Z	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check, Chemistry and TDM	43
Vancomycin	Х	CZ, CZX, CZ2X, Z	Chemistry and TDM	56- 58
		CZQ	Quality Cross Check,	43
			Chemistry and TDM	
		LN3	TDM Cal Ver/Lin	119
Vancomycin-resistant		IDN, IDO	Nucleic Acid Amp,	196
Enterococcus			Organisms	
		VRE	Vancomycin-resistant Enterococcus	182
Vanillylmandelic acid	Х	N/NX	Urine Chemistry, Special	69
Variant interpretation only		VIP, VIP1	Variant Interpretation Only	244
Varicella-zoster virus		ID1	Nucleic Acid	196
(VZV)			Amplification	
		IDME	Meningitis/Encephalitis Panel	198
	Χ	VR1	Virology Culture	191
	Х	VR2	Viral Antigen Detection by DFA	191
	X	VR3	Antibody Detection- Infectious Disease Serology	200

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Vascular endothelial growth factor (VEGF)		CTKN	Cytokines	206
Venlafaxine		Т	Toxicology	96
		UT	Urine Toxicology	96
Verapamil		T	Toxicology	96
		UT	Urine Toxicology	96
Viability		CBT	Cord Blood Testing	221
		SCP	Stem Cell Processing	221
Viral antigen detection	Х	HC2	HSV by DFA	192
		POC8	POC Influenza A/B Ag	52
	Х	VR2	Viral Antigen Detection by DFA	191
	Х	VR4	Viral Antigen Detection by EIA and Latex	192
Viral isolation/ identification	Х	HC4	HSV Culture	192
	Х	ID3	Influenza A, Influenza B, RSV by NAA	197
	Χ	IDR	Infectious Disease, Respiratory Panel	198
	Χ	VR1	Virology Culture	191
Virtual biopsy program, online		VBP/VBP1	Online Virtual Biopsies Program	258
Virtual gram stain		VGS1	Virtual Gram Stain Basic	175
		VGS2	Virtual Gram Stain	175
			Advanced	
Virtual peripheral blood smear		VPBS	Virtual Peripheral Blood Smear	141
Viscosity		V	Viscosity	208
Vitamin A		BMV3	Bone Markers and Vitamins	86
Vitamin B12	Χ	K, KK, K2	Ligand Assay, General	82
		LN5	Ligand Assay Cal Ver/Lin	119- 120
		LN5S	Ligand Assay, Siemens Cal Ver/Lin	119- 120
Vitamin D, 25-OH	Х	ABVD	Accuracy-Based Vitamin D	85
		LN40	Vitamin D Cal Ver/Lin	127
	Χ	VITD	25-OH Vitamin D	84
Vitamin D, 1, 25 dihydroxy		BMV1	Bone Markers and Vitamins	86
Vitamin E		BMV4	Bone Markers and Vitamins	86
VKORC1		PGX	Pharmacogenetics	243
Volatiles	Х	AL1	Whole Blood Alcohol/ Ethylene Glycol/	101
			Volatiles	
	Х	AL2	Serum Alcohol/Ethylene Glycol/Volatiles	101
von Willebrand factor		CGE/CGEX	Coagulation, Extended	159

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
von Willebrand factor		CGS3	Coag Special, Series 3	160-
(cont.)				161
		LN37	von Willebrand Factor Ag Cal Ver/Lin	127
VZV		ID1	Nucleic Acid	196
			Amplification	
	X	VR1	Virology Culture	191
	Х	VR2	Viral Antigen Detection by DFA	191
	Х	VR3	Antibody Detection-	200
			Infectious Disease Serology	
Wavelength and photometric calibration		I	Instrumentation	129
WBC count		ABF1, ABF2, ABF3	Automated Body Fluid	146
		CBT	Cord Blood Testing	221
	X	FH1-FH4, FH6, FH9, FH10, FH13, FH1P-FH4P, FH6P, FH9P, FH10P, FH13P	Hematology, Auto Diff	135
		FH3Q, FH4Q, FH6Q, FH9Q	Quality Cross Check, Automated Hematology Series	47
	Х	FH15	Centrifugal Hematology	135
		FL4	Flow Cytometry CD34+	209
	Х	HE, HEP	Basic Hematology	134
		LN9	Hematology Cal Ver/Lin	121
	Х	RWBC	Rapid Total White Blood Cell Count	141
		SCP	Stem Cell Processing	221
WBC automated count (fluid)		ABF1, ABF2, ABF3	Automated Body Fluid	146
WBC manual count (fluid)	Х	HFC, HFCI	Hemocytometer Fluid Count	148- 149
WBC count (urine)		UAA, UAA1	Automated Urinalysis	147
WBC count (leukocyte-		TRC	Transfusion-Related	218
reduced platelets)			Cell Count	
WBC count (leukocyte- reduced RBCs)		TRC	Transfusion-Related Cell Count	218
WBC differential		EHE1	Expanded Virtual	142
(manual)			Peripheral Blood Smear	
		VPBS	Virtual Peripheral Blood Smear	141
WBC differental (2-part)	Х	FH15	Centrifugal Hematology	135

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
WBC differential, automated	X	FH1-FH4, FH6, FH9, FH10, FH13, FH1P-FH4P, FH6P, FH9P, FH10P, FH13P	Hematology, Auto Diff	135
		FH3Q, FH4Q,	Quality Cross Check,	47
		FH6Q, FH9Q	Automated Hematology	
			Series	
WBC differential, body fluid		VBF	Virtual Body Fluid	146
WBC morphology		EHE1	Expanded Virtual	142
			Peripheral Blood Smear	
		VPBS	Virtual Peripheral Blood Smear	141
West Nile virus	Χ	NAT	Nucleic Acid Testing	226
Worm identification		WID	Worm Identification	189
Yeast identification	Х	F	Mycology and Aerobic Actinomycetes	184
	Х	F1	Yeast	184

Analyte/Procedure	LAP ENR	Program Code	Description	Pg
Yeast identification (cont.)	Х	F3	Candida Culture	185
Yersinia enterocolitica		GIP	Gastrointestinal Panel	199
Zaleplon		DFC	Drug-Facilitated Crime	107
ZAP-70		ZAP70	ZAP-70 Analysis by Flow Cytometry	212
Zika virus		VBDM	Vector-Borne Disease- Molecular	195
Zinc	Х	R	Trace Metals	78
Zinc, urine		TMU	Trace Metals, Urine	103
Zinc, whole blood		TMWB	Trace Metals, Whole Blood	103
Ziprasidone		DFC	Drug-Facilitated Crime	107
Zolpidem		DFC	Drug-Facilitated Crime	107
		FTC	Forensic Toxicology, Criminalistics	104
		Т	Toxicology	96
		UT	Urine Toxicology	96
Zopiclone/Eszopiclone		DFC	Drug-Facilitated Crime	107
Zonisamide		ZE	Therapeutic Drug Monitoring, Extended	60

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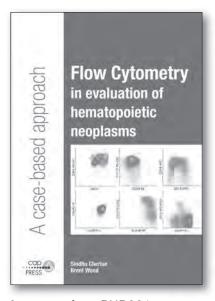
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23 Program Code Page Index



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Program Code Page Index

Program Code	Pg	Program Code	Pg	Program Code	Pg	Program Code	Pg	Program Code	Pg
A2MG	204	AQ3Q	46	C4	56-58	CRYP	185	DMPM	106
AAT	239	AQ4	92	C7	77	CS	59	DPATH	259
ABF1	146	AQ4Q	46	CAMP	177	СТ	162	DPATH1	259
ABF2	146	AQQ	46	CBT	221	CT1	162	DSC	147
ABF3	146	ARP	205	CCP	206	CT1Q	50	DY	84
ABL	110	ASA	154	CD	102	CT2	162	E	183
ABOSG	216	ASC	205	CDF2	178	CT2Q	50	E1	183
ABS	111	ASO	202	CDF5	178	CT3	162	EGFR	252
ABT	217	AUP	270	CES	206	CT3Q	50	EHE1	142
ABT1	217	AUP1	270	CESX	206	CT5	162	ELU	218
ABT2	217	B27	231	CFDNA	252	CT5Q	50	EMB	155
ABT3	217	BALL	210	CGB	158	CTKN	206	EP0	86
ABTH	112	BCP	134	CGDF	158	CTQ	50	ESR	134
ABU	111	BCP2	134	CGE	159	CY	236	ESR1	134
ABVD	85	BCR	147	CGEX	159	CYBK	236	ESR2	134
ACA	204	BCS	176	CGL	158	CYCGH	238	ESR3	134
ACE	71	BCS1	177	CGLQ	49	CYCMA	238	ETB	102
ACL	205	BDP	222	CGM	167	CYF	236	ETME1	223
ADL	71	BDP5	222	CGS1	160-	CYH	237	EV	59
AFL	146	BDPV	222		161	CYI	236	EXM	215
AG	71	BDPV5	222	CGS2	160-	CYJ	237	EXM2	216
AHIV	225	BFC	147		161	CYK	237	F	184
AHIVW	225	BGL	239	CGS3	160-	CYL	237	F1	184
AHT	204	BGL1	239		161	CYS	74	F3	185
AL1	101	BGS	85	CGS4	160-	CZ	56-58	FCFS	75
AL2	101	BL	102	l	161	CZ2X	56-58	FCN	204
AMH	84	BMD	138	CGS5	160-	CZQ	43	FF	86
ANA	202	BMV1	86		161	CZVM	67	FGAL	185
APAPCE	273	BMV2	86	CGS6	160- 161	CZX	56-58	FH1	135
APAPCPT	272	BMV3	86	CGS7	160-	D	170	FH1P	135
APAPJE	273	BMV4	86	Cus/	161	D1	172	FH2	135
APAPJPT	272	BMV5	86	CGS8	160-	D2	172	FH2P	135
APAPKE	273	BMV6	86	0030	161	D3	172	FH3	135
APAPKPT	272	BNP	61	CH50	208	D4	173	FH3P	135
APAPLE	273	BNP5	61	CHPVD	274	D5	174	FH3Q	47
APAPLPT	272	BNPQ	43	CHPVJ	274	D6	175	FH4	135
APAPME	273	BOR	177	CHPVK	274	D7	172	FH4P	135
APAPMPT	272	BP	188	CHPVM	274	D8	176	FH4Q	47
APC	204	BRAF	252	CMMP	145	D9	175	FH5	135
APOE	239	BRAFV	267	CMP	144	DADR1	233	FH5P	135
APS	205	BRCA	240	CMP1	144	DADR2	233	FH6	135
APT	148	BSTS	195	CMQ	48	DAI	98	FH6P	135
APXBN	161	BU	85	CPIP	14	DAT	218	FH6Q	47
AQ	92	BV	181	CPIP1	14	DBGN	161	FH7	135
AQ2	92	C1	56-58	CRP	202	DEX	171	FH7P	135
AQ2Q	46	C3	56-58	CRT	62	DFC	107	FH8	135
AQ3	92	C3X	56-58			DML	231	FH8P	135
	1		1	CRTI	62				

Program Code	Pg	Program Code	Pg	Program Code	Pg	Program Code	Pg	Program Code	Pg
FH9	135	HBVL	193	IGX	202	LN19	124	MRD2	254
FH9P	135	HBVL5	193	IL	202	LN20	124	MRS	179
FH9Q	47	HC1	178	IM	202	LN21	124	MRS2M	179
FH10	135	HC2	192	IMD1	242	LN22	124	MRS5	179
FH10P	135	HC3	178	IMD2	242	LN23	125	MRS5M	179
FH13	135	HC4	192	IMD3	242	LN24	125	MSI	250
FH13P	135	HC6	194	IMW	203	LN25	125	MTBR	183
FH15	135	HC6X	194	IND	186	LN27	125	MTP	253
FID	280	HC7	194	ING	86	LN30	125	MVM	80
FL	209	HCC	66	ISH	250	LN31	126	MX1B	230
FL1	209	HCC2	66	ISH2	250	LN32	126	MX1C	230
FL2	209	HCG	202	J	214	LN33	126	MX1E	230
FL3	209	HCV2	193	J1	214	LN34	126	MX2B	230
FL4	209	HE	134	JAT	215	LN35	127	MX2C	230
FL5	210	HEP	134	JATE1	215	LN36	127	MX2E	230
FLAC	178	HER2	269	JE1	214	LN37	127	MXB	231
FLD	72	HFC	148	K	82	LN38	127	MXC	231
FLD2	73	HFCI	149	K2	82	LN39	127	MYG	69
FLDQ	44	HG	139	KET	64	LN40	127	N	69
FNA	277	HGM	240	KIT	252	LN41	128	NAT	226
FNA1	277	HIVG	193	KK	82	LN42	128	NB	65
FNAG	278	HMS	64	KRAS	252	LN43	128	NB2	65
FNAG1	278	HPATH	143	KSA	69	LN44	128	NEO	251
FNPX	161	HPATH1	143	KVM	90	LN45	127	NGC	276
FOL	88	HPS	178	LBAS	176	LPE	76	NGC1	276
FP	87	HPV	195	LBC	149	LPX	180	NGS	246
FP1B	87	HQBX1	266	LCW	64	M	74	NGSB1	247
FP1T	87	HQBX2	266	LKM	207	MBT	171	NGSB2	247
FPX	87	HQBX3	266	LN2	118	MC1	173	NGSBV	249
FR	281	HQBX4	266	LN2BV	118	MC2	173	NGSE	248
FR1	281	HQIHC	264	LN3	119	MC3	174	NGSHM	246
FSER	185	HQIP	263	LN5	119-	MC4	174	NGSST	246
FSM	186	HQIPBX	265		120	MC5	174	NIPT	87
FT	75	HQMMR	264	LN5S	119-	ME	232	NP	271
FTC	104	HQNSC	265		120	MGL1	241	NP1	271
G	208	HSCRP	64	LN6	120	MGL2	241	NTA	102
G6PDS	75	HUEP	89	LN7	121	MGL3	241	NX	69
GH2	63	HV2	193	LN8	121	MGL4	241	OCB	149
GH5	63	1	129	LN9	121	MGL5	241	OCBQ	48
GH5I	63	ID1	196	LN11	122	MHO	254	OFD	100
GHER2	269	ID1T	196	LN12	122	MH01	254	OLI	74
GHQ	44	ID2	197	LN12E	122	MHO2	254	Р	187
GIP	199	ID3	197	LN13	122-	MHO3	254	P3	187
GLI	253	IDME	198	111400	123	MH05	251,	P4	187
GNBC	199	IDN	196	LN13C	122-		254	P5	187
GOCB	148	IDO	196	LNIAE	123	MK	267	PAPCE	273
GPBC	199	IDR	198	LN15	123	MMR	267	PAPCPT	272
GSA	64	IFS	130	LN16	123	MPA	59	PAPJE	273
Н	204	IG	202	LN17	123	MRD	254	PAPJPT	272
HBF	218	IGHV	250	LN18	124	MRD1	254	PAPKE	273
				I		I			

Program Code	Pg	Program Code	Pg	Program Code	Pg	Program Code	Pg
PAPKPT	272	POC12	53	SALC	77	TPM	244
PAPLE	273	POC14	54	SARC	251	TQPDelayed until	2019
PAPLPT	272	POC15	54	SBAS	176	TRC	218
PAPME	273	POC16	54	SC	154	TTD	200
PAPMPT	272	PS	219	SC1	154	TVAG	189
PARF	227	PTHQ	45	SCDD	105	U	68
PCARM	65	PV	154	SCM1	150	UAA	147
PCARMX	65	QF	207	SCM2	150	UAA1	147
PCNEO	211	QM1	39	SC0	131	UBJP	76
PCP1	186	QT1	31	SCP	221	UDC	99
PCP2	186	QT2	31	SCS	140	UDS	98
PCP4	186	QT3	32	SDS	101	UDS6	98
PCSP	262	QT4	32	SE	207	UDSM	108
PCSP1	262	QT5	37	SEC	242	UHCG	150
PCT	77	QT7	33	SEC1	242	UMC	151
PDL1	267	QT8	33	SFLC	208	UPBG	70
PEX	189	QT10	34	SM	154	URC	147
PF	164	QT15	35	SM1CD	154	UT	96
PF1	164	QT16	35	SM2CD	154	UTCO	131
PGX	243	QT17	36	SMCD	154	UVM	70
PGX1	243	R	78	SO	94	V	208
PGX2	243	RAG	216	SOQ	46	VBDM	195
PGX3	243	RAP	89	SP	180	VBF	146
PHG	76	RBCAT	217	SP1	180	VBP	258
PIA	165		207	SPE	76	VBP1	258
PIAX	165	RDS	243	SPN		VF	
		RETT			180		101
PIP	256	RF	202	ST	181	VGS1	175
PIPM	256	RFAV1	212	STFR	80	VGS2	175
PIPW	257	RFAV2	212	SV	154	VIP	244
PIPW1	257	RFX	202	SW1	79	VIP1	244
PLA	75	RHCVW	225	SW2	79	VITD	84
PLTM	167	RMAL	188	SW3	79	VLS	193
PM1	268	RNA	252	SW4	79	VLS2	193
PM2	269	ROM1	150	T	96	VM1	224
PM3	268	RT	139	TBLA	78	VM2	224
PM5	268	RT2	139	TBX	166	VM3	224
PM6	268	RT2Q	47	TEG	164	VM4	224
PNA1	177	RT3	139	TICP	275	VM5	225
PNA2	177	RT3Q	47	TICP1	275	VM6	225
PNH	211	RT4	139	TM	89	VPBS	141
POC1	52	RT4Q	47	TMCA	219	VR1	191
POC2	52	RTQ	47	TMCAD	219	VR2	191
POC3	52	RUB	202	TMCAE	220	VR3	200
POC4	52	RUBX	202	TMCAF	220	VR3M	200
POC6	52	RUR	180	TMO	189	VR4	192
POC7	52	RVBN	161	TMU	103	VRE	182
POC8	52	RWBC	141	TMWB	103	VS	181
POC9	52	S2	203	TMX	89	VS1	181
POC10	53	S4	203	TNT	62	VS2	182
POC11	53	S5	203	TNT5	62	WBCR	66

Program Code

WBGQ

WID

WP3

WP4

WP6

WP9

WP10

YVM

ΥY

Z

ZE

ZT

ZAP70

Pg

43

189

166

166

166

166

166 84

90

84

56-58

212

60

60

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