

Abnormal PT/PTT Analyzer

Order Name: PT PTT AN

Test Number: 1507500

TEST COMPONENTS	REV DATE: 06/04/2012
Test Name:	Methodology:
Abnormal PT/PTT Analyzer	

SPECIMEN REQUI	REMENTS			
	Specimen Volume(min)	Specimen Type	Specimen Container	Transport Environment
Preferred	46 mL	See Instructions	See Instructions	See Instructions
Instructions:	Please indicate anticoagulant therapy and submit with specimen or fax "Coagulopathy Questionnaire Form" to 918-744-3236. Please collect Twelve 3mL Sodium Citrate 3.2% (Blue Top) tubes and One 10mL Clot tube (Tiger Top) and One 5mL EDTA (Lavender Top). Each Sodium Citrate blue top tube must be filled to the proper level, no hemolysis. Improperly filled tubes can give erroneous results. Whole blood must be transported to lab immediately. Citrated plasma must be filtered and frozen in 1 ml aliquots.			

GENERAL INFORMATION

Testing Schedule: Individual Test Dependant

Expected TAT: 5-10 Days

Clinical Use: This analyzer is designed to evaluate patients with an unexplained prolonged PT or PTT in whom there is no clinical history or strong clinical suspicion of either bleeding or thrombolytic tendency. A pathologist interpretation and patient focused report with summation of

test results will be issued with each order.

Notes: For more information on this Analyzer, access our "Specialized Tests" section of this guide for a complete listing of tests and CPT

Cpt Code(s): See the Test Notes Section of this test.



Amino Acid Analysis, Quantitative, Plasma

Order Name: AA QN BL Test Number: 3617225

TEST COMPONENTS		REV DATE: 06/06/2012
Test Name:	Methodology:	
Amino Acid Analysis, Quantitative, Plasma	LC/MS	

SPECIMEN REQUI	REMENTS			
	Specimen Volume(min)	Specimen Type	Specimen Container	Transport Environment
Preferred	2mL (0.3mL)	Plasma	Sodium Heparin (Green Top)	Frozen
Alternate	2mL (0.3mL)	Plasma	Lithium Heparin PST (Light Green Top)	Frozen
Instructions:	Separate plasma within 30min of draw. Freeze immediately after separation from cells. Do not thaw. Provide patient age (required for correct reference range), sex, a brief clinical history, tentative diagnosis, and their therapy over the last three days (drugs, x-ray, infant formula, diet). *Note: Patient age is required for correct interpretation.			

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Testing Schedule: Mon, Wed - Fri, Sat

Expected TAT: 10-12 Days
Cpt Code(s): 82139



Antidiuretic Hormone (ADH) and Osmolality

Order Name: ADH/OSMO

Test Number: 3600235

TEST COMPONENTS

REV DATE: 06/12/2012

Test Name:

Antidiuretic Hormone (ADH, Arginine Vasopressin, AVP)

Osmolality Serum

SPECIMEN REQUIREMENTS

Specimen Type Specimen Container Transport Environment

Preferred See Instructions Plasma and Serum EDTA (lavender top) and Clot Activator SST (Red/Gray or Tiger Top) See Instructions

Special Collect Both Serum and Plasma

Instructions:

ADH: 6mL (2.5) EDTA Plasma, Frozen.

Separate plasma from cells ASAP or within 2 hours of collection.

Osmolality: 1mL (0.5) Serum, Refrigerated or Frozen.

GENERAL INFORMATION

Testing Schedule: Tue, Thr, Sat

Expected TAT: 3-11 Days (assay dependant)

Cpt Code(s): 83930, 84588



Hemoglobin Electrophoresis

Order Name: HGB ELECT

Test Number: 5000775

TEST COMPONENTS	REV DATE: 06/05/2012
Test Name:	Methodology:
Hemoglobin Electrophoresis	AGHEP

SPECIMEN REQUI	REMENTS			
	Specimen Volume(min)	Specimen Type	Specimen Container	Transport Environment
Preferred	5 mL (4.5)	Whole Blood	EDTA (Lavender Top)	Refrigerated
Special Instructions:	Please provide a full tube for best results. Specimen stability: Ambient= 24 hours, refrigerated= 5 days.			

GENERAL INFORMATION

Testing Schedule: MON, WED, FRI

Expected TAT: 7 Days

Clinical Use: Alkaline Gel Hemoglobin Electrophoresis is used to identify a large number of hemoglobin variants.

Notes: Additional High Performance Liquid Chromatography (HPLC) testing may be required to completely identify some hemoglobin varients.

See test "HGBOP HPLC" for more information.

Cpt Code(s): 83020; 80500



Herpes Simplex Virus 1 and 2 (HSV), DNA, PCR

Order Name: HSV PCR QL

Test Number: 5586635

TEST COMPONENTS	REV DATE: 06/28/2012
Test Name:	Methodology:
Herpes Simplex Virus, Type 1 DNA	PCR
Herpes Simplex Virus, Type 2 DNA	PCR

SPECIMEN REQUI	REMENTS			
	Specimen Volume(min)	Specimen Type	Specimen Container	Transport Environment
Preferred	2mL(0.3mL)	CSF (Cerebrospinal Fluid)	Sterile Screwtop Container	Refrigerated
Alternate	2mL(0.3mL)	Plasma	EDTA (Lavender Top)	Refrigerated
	2mL(0.3mL)	Serum	Clot Activator SST (Red/Gray or Tiger Top)	Refrigerated
Special Instructions:	Special Indicate Specimen Source on the Specimen Container.			ative, Pleural fluid,

GENERAL INFORMATION

Testing Schedule: Mon - Fri
Expected TAT: 2-3 Days
Cpt Code(s): 87529X2



Hypercoagulation Analyzer

Order Name: HYPRCOAGAN

Test Number: 1506500

TEST COMPONENTS	REV DATE: 06/04/2012
Test Name:	Methodology:
Hypercoagulation Analyzer	

Specimen	Specimen Type	Specimen Container	Transport	
Chariman	Cassimon Tuna	Specimen Container	Transport	
SPECIMEN REQUIREMENTS				

Special Please list the patient's anticoagulant and submit with specimen or fax "Coagulopathy Questionnaire Form" to 918-744-3236. Instructions:

Fifteen (2.7mL) 3.2% Sodium Citrate (Blue Top) tubes,

Two (4.7mL) EDTA (Lavender Top) tubes,

One (7mL) lithium heparin (green top) tube (on ice or frozen pour off aliquot) and

One (10mL) Clot Activator SST (Red/Gray Top) tube.

Tubes must be filled to the proper level, no hemolysis. Improperly filled tubes can give erroneous results. Whole blood must be transported to lab immediately. If submitting only Citrated plasma, plasma must be double spun and frozen in 1.5 ml aliquots. **Do not pool plasma from multiple tubes!**

GENERAL INFORMATION

Testing Schedule: Mon - Fri

Expected TAT: Testing dependent

Clinical Use: A comprehensive algorithm used to assess the cause of hypercoagulability.

Notes: Algorithm begins with an Activated Protein C Resistance, Homocysteine, Lupus sensitive PTT, Prothrombin time (PT), Prothrombin

Gene Mutation, and a Partical Thromboplastin Time (PTT). Further testing is generated based on the results of these tests. A pathology interpretation is included with all orders. For more information on this Analyzer, access our "Specialized Tests" section of this

guide for a complete listing of tests and CPT codes.

Cpt Code(s): See the Test Notes Section of this test.



Lupus Anticoagulant Analyzer

Order Name: LUP ANT AN

Test Number: 1506300

TEST COMPONENTS		REV DATE: 06/04/2012
Test Name:	Methodology:	
Cardiolipin Antibodies, IgM and IgG	EIA	
Prothrombin Time (PT) and INR	CLOT	
Activated Partial Thromboplastin Time (aPTT)	CLOT	
Lupus Anticoagulant PTT	CLOT	
Beta 2 Glycoprotein IgG and IgM Antibody	EIA	
Pathology Report		

SPECIMEN REQUIREMENTS						
	Specimen Volume(min)	Specimen Type	Specimen Container	Transport Environment		
Preferred	See Instructions	See Instructions	Sodium Citrate 3.2% (Blue Top) and Clot Activator SST (Red/Gray or Tiger Top)	See Instructions		
	Please list anticoagulant therapy and submit with specimen or fax "Coagulopathy Questionnaire Form" to 918-744-3236. Collect: Twelve 2.7mL Sodium Citrate Blue top tubes and One 10mL Tiger top clot tube. Each blue top tube must be filled to the proper level, no hemolysis. Improperly filled tubes can give erroneous results. Whole blood must be transported to lab immediately. If sending citrated plasma aliquots, they must be double spun then aliquot 1.5 ml plasma from each tube into individual plastic aliquot tubes and freeze. Do not pool aliquots together!					

GENERAL INFORMATION

Testing Schedule: Individual Test Dependant

Expected TAT: 5 Days

Clinical Use: This analyzer is designed to evaluate patients in whom there is a clinical suspicion of a lupus anticoagulant or clinical features of the

anti-phospholipid syndrome (e.g. thrombocytopenia, thrombosis, recurrent abortion).

Notes: The algorithm begins with a Prothrombin Time (PT/INR), Partial Thromboplastin time (PTT), Lupus Sensitive PTT, Beta 2 Glycoprotein IgG/IgM Antibodies and Cardiolipin IgG/IgM testing. Subsequent tests are generated based on the results of this first level of testing. A

pathology interpretation is included with all orders. For more information on this Analyzer, access our "Specialized Tests" section of this

guide for a complete listing of tests and CPT codes.

Cpt Code(s): See the Test Notes Section of this test.



Phenobarbital, Free, Serum or Plasma

Order Name: PHENOB FR

Test Number: 3804075

TEST COMPONENTS		REV DATE: 07/02/2012
Test Name:	Methodology:	
Phenobarbital, Free, Serum or Plasma	HPLC	

SPECIMEN REQUIREMENTS					
	Specimen Volume(min)	Specimen Type	Specimen Container	Transport Environment	
Preferred	3mL(1.2)	Serum	No Additive Clot (Red Top, No-Gel, Plastic)	Refrigerated	
Alternate	3mL(1.2)	Plasma	EDTA (Lavender Top)	Refrigerated	
	3mL(1.2)	Plasma	Lithium Heparin (Dark Green Top / No-GEL)	Refrigerated	
Special Instructions:	Promptly centrifuge and separate Serum or Plasma into a plastic screw capped vial using approved guidelines.				

GENERAL INFORMATION

Testing Schedule: Mon - Fri
Expected TAT: 3-4 Days
Cpt Code(s): 80184



Platelet Autoantibody

Order Name: PLT AUTOAB

Test Number: 5577375

TEST COMPONENTS	REV DATE: 06/19/2012
Test Name:	Methodology:
Platelet Autoantibody	ELISA

SPECIMEN REQUIREMENTS

Specimen Volume(min)

Specimen Type Specimen Container

Specimen Container

Transport
Environment

Preferred 5 mL (3)

Whole Blood

EDTA (Lavender Top)

On Ice

Special Instructions: All Collections Must Be Scheduled for Collection! Contact: Immunology (918)744-2553 x15511 or Immunology Manager x15788.

Due to specimen integrity and stability reasons, it is highly suggested this should be collected at the RML Main Laboratory - Monday through Thursday..!

Testing must begin within 24hrs.

Please make this a Separate Specimen, Do not share this specimen with other testing.

Keep specimen as Whole Blood and On-ICE..! Do Not Centrifuge!

Transport specimen directly to Lab Section ASAP On-ICE but Do Not Freeze!

(Note: Special processing of the sample will be performed within the performing laboratory section. Hemolyzed, Icteric and Lipemic specimens will yield false results and will not be tested.)

GENERAL INFORMATION

Testing Schedule: Mon-Thr
Expected TAT: 2-3 Days

Clinical Use: The platelet autoantibody study is designed to detect platelet autoantibodies eluted from the patient's platelets or circulating in the

patient's serum or plasma directed against GPIIb/IIIa, GPIb/IX, and GPIa/IIa. These antibodies can be detected in patients with autoimmune thrombocytopenic purpura (ITP or AITP). This test is intended to help identify patients who present with unexplained thrombocytopenia that is secondary to immune destruction. A positive test is considered diagnostic, while a negative test does not rule

out the diagnosis. Repeat testing can sometimes be of benefit.

Cpt Code(s): 86022



Pre Albumin

Order Name: PRE ALB Test Number: 3603830

TEST COMPONENTS		REV DATE: 06/19/2012
Test Name:	Methodology:	
Pre Albumin	Turb	

SPECIMEN REQUIREMENTS					
	Specimen Volume(min)	Specimen Type	Specimen Container	Transport Environment	
Preferred	1 mL (0.5)	Serum	Clot Activator SST (Red/Gray or Tiger Top)	Refrigerated	
Special Instructions:	Stability: Refrigerated 7 days. Freeze for greater than 7 days.				

GENERAL INFORMATION

Testing Schedule: Mon - Fri Expected TAT: 1-2 days

Clinical Use: Use to evaluate protein malnutrition, total parenteral nutrition, and liver dysfunction. Serum level decreased in inflammatory processes, malignancy. Serum level increased in Hodgkin's disease.

Cpt Code(s): 84134



Prograf (FK506)

Order Name: PROGRAF
Test Number: 4503275

TEST COMPONENTS	REV DATE: 06/19/2012
Test Name:	Methodology:
Prograf (FK506)	MEIA

SPECIMEN REQUI	REMENTS			
	Specimen Volume(min)	Specimen Type	Specimen Container	Transport Environment
Preferred	1 mL (0.5)			Room Temperature
	To be drawn 12 hours aft to 14 days. Frozen great		4 hour after dose for 24 hour trough. St	ability: Ambient 3 days. Refrigerated up

GENERAL INFORMATION

Testing Schedule: Daily

Expected TAT: 24hrs (same day if specimen is in lab by 11am)

Clinical Use: Useful for assessing the adequacy of systemic drug delivery since metabolism can exhibit significant variability.

Notes: Also known as Tacrolimus

Cpt Code(s): 80197



Vaginosis Profile from Swab (with Trichomonas Antigen)

Order Name: V PROF SWB

Test Number: 2915445

TEST COMPONENTS		REV DATE: 06/25/2012
Test Name:	Methodology:	
Whiff test	Amine	
Gram Stain	MC	
Trichomonas Antigen	EIA	
Vaginal Yeast Examination	MC	
Clue Cell Examination	MC	

SPECIMEN REQUIREMENTS					
	Specimen Volume(min)	Specimen Type	Specimen Container	Transport Environment	
Preferred	See Instructions	Swab	BBL Red top culturette in Amies media (double swab)	See Instructions	
Alternate	See Instructions	Swab	BBL White top culturette swab (double swab)	See Instructions	
	Collect BBL Red top culturette in Amies media (double swab) or BBL White top culturette (double swab preferred) Specimen Stability: 24hrs Room Temperature or 36hrs Refrigerated (Do Not Freeze). Samples contaminated with preparations containing iodine or by the immediate prior use of vaginal lubricants are not recommended. BBL Blue top swabs are Not Acceptable.				

	. INFC	

Testing Schedule: Sun-Sat
Expected TAT: 1 Day

Clinical Use: This vaginosis profile provides an interpretation of the types of vaginal pathology present: Yeast infections, Trichomonas vaginalis,

Bacterial vaginosis and even Mixed Flora infections. The Trichomonas antigen along with gram stain and evaluation for yeast,

clue cells, white blood cells and all bacterial types present. Whiff test (amine test) is reported as positive or negative.

Notes: Created to handle extended transportation times seen with vaginosis profile specimens

Cpt Code(s): 87205, 87808, 82120



