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Running Specimens on the TEG 6s Hemostasis Analyzer

PURPOSE

This procedure describes how to run patient blood specimens on the Haemonetics TEG 6s hemostasis analyzer which is used to monitor and analyze the coagulation state of a blood specimen.

POLICY

 Blood specimens may be sent to the laboratory using the pneumatic tube system or walked to the laboratory

SCOPE

· All CLS and MLT staff assigned to the Hematology/Coagulation department

TEG 6S ORDER CODES

- TEGSTD use Citrated Global Hemostasis with CKH Test Cartridge (purple stripe test cartridge)
- TEGRPD use Citrated Global Hemostasis with Lysis 30 Test Cartridge (yellow stripe test cartridge)
- TEGPM use Platelet Mapping Test Cartridge (red stripe test cartridge)

SPECIMEN COLLECTION and REQUIREMENTS

Test Cartridges	Specimen Collection	(Phlebotomy)	Specimen Collection (Line Draw)
Citrated Global Hemostasis Test Cartridges	Not use red top)	with no additive (do discard 3-5 ml prior to completely filled by eversion 5 times ube	 Use no smaller than 21 gauge needle Use 2 syringes, first syringe to discard blood 3 times the dead space Second syringe the required amount of blood Use safe transfer device to allow 3.2 % citrate blood tube to completely fill by vacuum

	 Do not centrifuge Blood tube is allowed to equilibriate at RT for 10 minutes in a horizontal position 	 Mix by gentle inversion 5 times Properly labeled tube Do not refrigerate or freeze Do not centrifuge
Platelet Mapping Test Cartridge	 Use no smaller than 21 gauge needle Draw discard tube with no additive (do Not use red top) If syringe is used, collecting blood Non gel lithium or sodium heparin tube with ≥ 14.5 IU but ≤ 20 IU heparin/ml and completely filled by vacuum Mixed by inversion 5 times Properly labeled tube Do not refrigerate Do not centrifuge Blood tube is allowed to equilibriate at RT for 30 minutes in a horizontal position 	 Refer to line draw specimen collection above Use non gel lithium or sodium heparin tube with ≥ 14.5 IU but ≤ 20 IU

SPECIMEN REJECTION CRITERIA

Test Cartridges	Specimen Interferences
Citrated Global Hemostasis Test Cartridges	 Clotted specimen Hemolysis Incorrect blood container Under filled blood tube Refrigerated or Frozen specimen Centrifuged specimen Unlabeled specimen Protamine contamination Hemodilution Heparin concentration above 1 IU/ml Specimen stability exceeded
Platelet Mapping Test Cartridge	 Clotted specimen Hemolysis Incorrect blood container Under filled blood tube Refrigerated or Frozen specimen Centrifuged specimen Unlabeled specimen Specimen stability exceeded

SPECIMEN STABILITY

Test Cartridge

Specimen stability

Citrated Global HemostasisTest Cartridges	 Specimen can be tested after 10 minutes incubation at room temp in a horizontal position and within 4 hrs of collection In an urgent or emergent situation, citrate specimen may be run immediately
Platelet Mapping Test	Specimen can be tested after 30 minutes incubation at room temp in a
Cartridge	horizontal position and within 2 hrs of collection

EQUIPMENT and SUPPLIES

- Haemonetics TEG 6s hemostasis analyzer
- · Scanner and thermal printer
- · printer paper
- · UPS
- · Functional Fibrinogen kit
- · Citrated Global Hemostasis Test Cartridge
- Citrated Global Hemostasis with Lysis (Trauma)Test Cartridge
- Platelet Mapping Test Cartridge
- · Abnormal QC Vial Test Kit
- · Abnormal & Lysis QC kit
- · Personal Protective Equipment

PROCEDURE A

Follow the procedure below to run TEG 6s patient test cartridge

Step	Action
1.	Log into the TEG 6s analyzer using your unique log-in ID then touch Log In. On the login screen, enter your user name in the username box using the keyboard press tab on the keyboard to move to the password box enter your password then press the return key on the keyboard If TEG 6s analyzer is left unused for a period of time, the display will dim If display is dimmed, touch the screen to remove from stand by
2.	From the home screen, touch new test
3.	Touch the (+) symbol
4.	On the Add Patient Screen, manually enter the patient's MRN
5.	Verify the patient's MRN on the display screen with the patient's MRN on the LIS barcode label
6.	Select OK
7.	At the Confirm Patient Data Screen, Select Confirm to validate the patient's information
8.	At the Preparing Test Screen, insert test cartridge as displayed on the screen Remove the patient test cartridge from its sealed pouch, manually inspect for any damage, and insert with the barcode on the left side The cartridge opening on the instrument will be lit and flashing when the instrument is ready to receive the cartridge

- DO NOT remove the cartridge until prompted to do so, either after normal completion of the test or as a result of choosing to stop the test
- Once test cartridge is inserted, verify that you have selected the correct test cartridge and have the correct sample type then touch Next to confirm
 - When cartridge is inserted the analyzer performs the Pre Test internal QC to verify the integrity of the cartridge-analyzer interface
 - · Any failure will result in an error
- 10. On the Test Information Screen, enter the patient name (last name then first name)
- 11. At the Preparing Test-Load sample screen display:
 - Ensure blood sample has met minimum equilibrium time in a horizontal position at RT dependent on test cartridge
 - Ensure the blood sample is mixed by gentle inversion 5-7 times
 - Use the transfer pipette included in the test kit to load blood into the cartridge sample port, filling up to or above the line marked on the side of the test cartridge
 - · DO NOT under fill the test reservoir
 - When loading blood into the cartridge sample port, Do Not release the transfer pipette bulb prior to removing the pipette tip from the cartridge reservoir to avoid inadvertently drawing the blood sample out
 - NOTE: Precise measurement is not necessary as any excess blood is moved to a sealed waste area within the cartridge during the test
- 12. Touch Next to start the test and display the Results Screen.
- The touch tracings button is used to view a graphic representation of the result and to cycle
 through superimposed, offset, and the single-tracing views until the desired view is
 displayed
 - The touch results button is used to return to the results screen
 - When the test is complete, the following changes occur on the results screen: The test timer
 is replaced by the date and time that the test started, the stop button is replaced by a done
 button, the print button is enabled
- 14. Log into the TEG manager:
 - At the TEG manager patient viewer screen, enter the patient's MRN or click Today then click search to view the patient's TEG tracings
 - In the patient's tracings screen, click the clipboard icon at the upper left hand corner to display the charting notes screen
 - In the Accession ID prompt, scan the LIS barcode to document the CID then scan the barcode again to confirm (CID should now display in the Accession ID prompt)
 - · Click the Done button
- 15. When the analyzer displays the "Remove cartridge" prompt and the cartridge port flashes, pull the used cartridge out of the slot and immediately dispose of it in a marked, biohazard waste receptacle
- 16. In the TEG manager and patient tracing display screen, print 2 copies of the patient's test report
 - · In the toolbar at the top of screen, click the Export button
 - · Click the Create report button

- · Click the open button
- · Click File at the upper left hand then click print
- · Retrieve the patient's test report
- Using the pneumatic tube system, send one copy of the patient's test report to the appropriate nursing unit
- Place the other copy of the patient's test report in the bin labeled "Epic to be scanned" located by the Epic scanner in OP processing

PROCEDURE B

Follow the procedure below to startup and shutdown the TEG 6s analyzer when necessary

Step Action

- 1. To power on the analyzer:
 - Ensure that the TEG 6s analyzer is connected to a power outlet
 - Turn on the analyzer by moving the power switch at the rear of the analyzer from the "O" to the "I" position
 - analyzer will startup and perform a power on self test
 - when power on self test is successful, the login screen displays
 - Log into the analyzer (refer to procedure A step 1)
- 2. To power off the analyzer:
 - Press logout in the lower left corner of the screen to log out of the home screen
 - Make sure to remove any existing test cartridge from the cartridge slot prior to turning off the analyzer
 - Turn off the analyzer by moving the power switch at the rear of the analyzer to the off position ("O")

REFERENCE RANGES AND REPORTABLE TESTS

Citrated Global He	mostasis Test Cartridge	
Test Assay	Test Parameter	Reference Range
СК	R (min)	4.6 - 9.1
	K (min)	0.8 - 2.1
	Angle (deg)	63 - 78
	MA (mm)	52 - 69
CRT	MA (mm)	52 - 70
СКН	R (min)	4.3 - 8.3
CFF	MA (mm)	15 - 32
	FLEV (mg/dl)	278 - 581
Citrated Global He	mostasis with Lysis Test Cartrid	lge (Trauma)
Test Assay	Test Parameter	Reference Range

CK	R (min)	4.6 - 9.1	
	LY30 (%)	0.0 -0 2.6	
CRT	MA (mm)	52 - 70	
CFF	MA (mm)	15 - 32	
Platelet Mapping 1	Test Cartridge		
Test Assay	Test Parameter	Reference Range	
нкн	MA (mm)	53 - 68	
ActF	MA (mm)	2 - 19	
ADP	MA (mm)	45 - 69	
	% Inhibition	0 - 17	
	% Aggregation	83 -100	
AA	MA (mm)	51 - 71	
	% Inhibition	0 - 11	
	% Aggregation	89 - 100	

REPORTING RESULTS

When the TEG test is complete, the TEG order code is reported by following the steps below

- · Sunquest manual entry function MEM
- Worksheet RVKM
- Under the ACC number, result with ETC "ROTEMI" (translated to "Thromboelastograph scanned to patient record")

RELATED DOCUMENTS

· TEG 6s Citrated and Platelet Mapping test cartridge package inserts

REFERENCES

· Haemonetics TEG 6s User Manual

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Attachments		
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