



HEALTH HOLDING

HAFER ALBATIN HEALTH
CLUSTER
MATERNITY AND
CHILDREN HOSPITAL

Department:	Laboratory and Blood Bank (Hematology)		
Document:	Internal Policy and Procedure		
Title:	Deficient Factors VIII,IX,XI,XII Assay		
Applies To:	All Hematology Staff		
Preparation Date:	January 07, 2025	Index No:	LB-IPP-066
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1. PURPOSE:

- 1.1 To provide Hematology technologists with a standard methodology for performing determination of the activity of coagulation factor in human plasma by coagulometric methods.

2. DEFINITIONS:

N/A

3. POLICY:

- 3.1 The assay consists of the measurement of the clotting time, in the presence of cephalin and activator, of a system in which all the factors are present, constant and in excess (supplied by STA® - Deficient VIII, IX, XI, XII) except factor VIII, IX, XI, XII which is derived from the sample being tested.

4. PROCEDURE:

4.1 Specimen:

- 4.1.1 Collect the blood in a blue stopper vacutainer tube using a ratio of 9 parts of whole blood and 1 part of 3.2% buffered sodium citrate.
- 4.1.2 Centrifuge the specimen at 3500 rpm for 10 minutes
- 4.1.3 Remove the platelet poor plasma and transfer to a 12 x 75 glass test tube. For the best results, test must be performed immediately.
- 4.1.4 Freeze the plasma at -20°C or lower until ready to test (MAX 2 weeks).
- 4.1.5 Frozen plasma must be thawed directly at 37°C for 15 minutes before testing.
- 4.1.6 Specimens are stable for 4 hours at +20°C and 2 weeks at -20°C.
- 4.1.7 Samples that have an abnormally high hematocrit, i.e. >55%, must be re-drawn into specially modified tubes that have had the volume of anticoagulant adjusted to ensure a correct ratio of blood to anticoagulant.

4.2 Procedural steps:

- 4.2.1 When the reagents are ready for use, scan the vial label bar-code across the bar-code reader.
- 4.2.2 Validate the reagent volume indicated by the analyzer. Then, place the vial in drawers as indicated.
- 4.2.3 When the beep emitted by machine this indicates to the operator that identification and position of the reagent are validated.
- 4.2.4 Factor VIII, IX, XI, XII assays of the plasmas to be tested are loaded automatically as soon as the samples have been loaded in accordance with the test definition indicated.
- 4.2.5 If any of the patient results falls outside the working range of the assay, the machine automatically retests the sample in question at an appropriate dilution.

4.3 Results:

- 4.3.1 The factor level VIII, IX, XI, XII (%) of the plasmas being tested is displayed in the "Test Panel/Test
- 4.3.2 If the control values are outside the stated ranges, check all components of the test system to ensure that all are functioning correctly, i.e., assay conditions, reagents, calibration, integrity of the plasmas being tested, etc. If necessary, repeat the assays.

5. MATERIALS AND EQUIPMENT:

- 5.1 Néoplastine® CI or - Néoplastine® CI Plus
- 5.2 Owren-Koller.
- 5.3 Unicalibrator
- 5.4 System Control + Stirring-bar
- 5.5 Kit containing control plasmas, normal and abnormal levels.

6. RESPONSIBILITIES:

- 6.1 This policy applies to all Hematology technologists involved in this special Hematology test.







7. APPENDICES:

- 7.1 Reagent Preparation And Storage: Factor VIII, IX, XI

8. REFERENCES:

- 8.1 Factor II, V, VII and X inserts Kit Manual.

9. APPROVALS:

	Name	Title	Signature	Date
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APPENDIX 7.1: REAGENT PREPARATION AND STORAGE: FACTOR VIII, IX, XI

Reagents	Preparation	Stability after reconstitution/ opening on board STA Compact®	Storage position on STA Compact®
STA® - Deficient XI	Add 1 ml of distilled water. Allow the reconstituted reagent to stand at room temperature (18-25 °C) for 30 minutes. Then, gently homogenize.	4 hours	Product drawer
STA® - Deficient VIII	Add 1 ml of distilled water. Allow the reconstituted reagent to stand at room temperature (18-25 °C) for 30 minutes. Then, gently homogenize.	4 hours	Product drawer
STA® - Deficient IX	Add 1 ml of distilled water. Allow the reconstituted reagent to stand at room temperature (18-25 °C) for 30 minutes. Then, gently homogenize.	4 hours	Product drawer
STA® - Néoplastine® CI or STA® - Néoplastine® CI Plus	Transfer the contents of a vial of Reagent 2 (Solvent) of the same kit. Allow the reconstituted reagent to stand at room temperature (18-25 °C) for 30 minutes. Gently homogenize. Then, place a stirring-bar, a new STA® - Reducer and the perforated cap.*	48 hours	Area with magnetic stirring-motor Product drawer
STA® - Owren-Koller	15-ml vial. Allow the solution to stand at room temperature (18-25 °C) for 30 minutes before use.	3 days	Sample drawer
STA® - Unicalibrator	Add exactly 1 ml of distilled water. Allow the solution to stand at room temperature (18-25 °C) for 30 minutes. Then, homogenize.	4 hours	Product drawer
STA® - System Control N STA® - System Control P	Add exactly 1 ml of distilled water. Allow the solution to stand at room temperature (18-25 °C) for 30 minutes. Then, homogenize.	8 hours	Product drawer