

FACTOR IX INHIBITOR EVALUATION #83103

USEFUL FOR: Detection and titering of coagulation inhibitor to factor IX in patients with hemophilia B.

NOTE: This test will replace #7801, Coag Factor IX Inhibitor Screen, Plasma, which will become obsolete on January 26, 2011.

PROFILE INFORMATION:

Unit Code	Reporting Name	Available Separately	Always Performed
9065	Coag Factor IX Assay, P	Yes	Yes

REFLEX TESTS:

Unit Code	Reporting Name	Available Separately	Always Performed
7802	Coag Factor IX Inhibitor Scrn, P	No	No
7288	Bethesda titer	No	No
82539	Coagulation Interpretation	No	No

METHOD: 9065: Activated Partial Thromboplastin Clot-Based Assay
 7289, 7288: Clot-Based Assay

REFERENCE VALUES: Factor IX Activity Assay: Adults: 65-140%
 Factor IX Inhibitor Screen: Negative
 (If positive, quantitated in Bethesda units)
 Bethesda titer: 0

SPECIMEN REQUIREMENTS: Draw blood in a light blue-top (citrate) tube(s). Spin down, remove plasma, spin plasma again, and place 2 mL of platelet- poor plasma into 2 plastic, screw-capped vials each containing 1 mL. (Glass vial is not acceptable.)

Specimen must be drawn prior to factor replacement therapy. Double-centrifuged specimen is critical for accurate results as platelet contamination may cause spurious results. Freeze specimens immediately at < or = -40 degrees C, if possible.

LIST FEE: \$237.60

The following test(s) may be added per the testing algorithm:

\$60.50 for #7802 “Coag Factor IX Inhibitor Screen”

\$123.70 for #7288 “Bethesda titer”

\$138.80 for #82539 “Coagulation interpretation”

CPT CODES:

85250-Factor IX activity

85335-Factor IX inhibitor (if appropriate)

85335-Bethesda titer (if appropriate)

85390-Coagulation Interpretation (if appropriate)

ANALYTIC TIME: 1 day

DAY(S) SET-UP: Monday through Friday

QUESTIONS: Contact your Mayo Medical Laboratories' Regional Manager
Kim J. Baker, Mayo Medical Laboratories' Technologist Support
Telephone: 800-533-1710