

Coagulation Studies

The Mayo Clinic Coagulation Laboratory has been performing coagulation factor testing on mailed-in specimens for many years. Accurate results can only be obtained on properly prepared specimens. The physician interpreting results may be misled by abnormal results obtained in mishandled specimens.

To ensure the best possible specimen, follow collection requirements as closely as possible.

1. **Patient should be fasting**, if possible; for certain tests, the patient cannot be receiving anticoagulant medication (heparin or warfarin/Coumadin®).
2. **Draw blood from the patient into light blue-top (sodium citrate) vacuum tube(s)** (those used for prothrombin time/activated partial thromboplastin time containing 3.2% sodium citrate). If the patient's hematocrit is $\leq 25\%$ or $\geq 55\%$, the volume of anticoagulant in the tube should be adjusted. Use the following formula to determine the correct anticoagulant volume:

$$\text{anticoagulant volume} = (100 - \text{hematocrit}) / (595 - \text{hematocrit}) \times \text{volume of specimen}$$

The tubes must fill completely. A clean venipuncture is essential to avoid activation of coagulation by tissue thromboplastin. Specimens containing fibrin clots will, in most cases, be rejected.

3. **The specimen must be double-centrifuged to prepare a platelet-free plasma specimen.** Immediately centrifuge specimen at 1,500 x G for 10 minutes, at 4° C, if possible. Carefully remove plasma from cells avoiding the platelet/buffy coat. Dispense specimen into a plastic tube. Centrifuge the plasma in the plastic tube at 1,500 x G for 10 minutes, at 4° C, if possible. Remove the top portion of plasma leaving approximately 250 μ L in the bottom to discard. The double-centrifuged plasma should be aliquoted (0.5-1 mL each) into clearly labeled plastic tubes. **(Glass vial is not acceptable.)** The number of tests ordered will determine the aliquots needed, generally 1 aliquot per test.
4. **Patient specimens should be frozen at $\leq -40^\circ$ C**, if possible, and sent together in the same container with at least 5 lbs of dry ice. They must arrive in a frozen state.
5. **Please include the requested information** (see individual test descriptions) as the testing and interpretations are dependent on clinical history in many of the more complex abnormalities.
6. Careful specimen handling will most often ensure acceptable specimens and valid results.

Pediatric Hemostasis References

1. Hathaway WE, Corrigan J: Report of Scientific and Standardization Subcommittee on Neonatal Hemostasis: normal coagulation data for fetuses and newborn infants. *Thromb Haemost* 1991;65:323-325
2. Andrew M, Paes B, Milner R, et al: Development of the human coagulation system in the full term infant. *Blood* 1987;70:165-172
3. Andrew M, Paes B, Milner R, et al: Development of the human coagulation system in the healthy premature infant. *Blood* 1988;72:1651-1657
4. Andrew M, Vegh P, Johnston M, et al: Maturation of the hemostatic system during childhood. *Blood* 1992;80:1998-2005
5. Andrew M: The hemostatic system in the infant. *In* *Hematology of Infancy and Childhood*. Vol. 1. 4th edition. Edited by DG Nathan, FA Oski. Philadelphia, WB Saunders Company, 1993, pp 115-153
6. Hathaway WE, Bonnar J: *Perinatal Coagulation*. New York, Grune and Stratton, 1978
7. Hathaway WE, Bonnar J: *Hemostatic Disorders of the Pregnant Woman and Newborn Infant*. New York, Elsevier Science Publishing Company, 1987
8. *Hematologic Disorders in Maternal-Fetal Medicine*. Edited by MN Bern, FD Frigoletto Jr. New York, Wiley-Liss, 1990
9. *Perinatal Thrombosis and Hemostasis*. Edited by S Suzuki, WE Hathaway, J Bonnar, AH Sutor. Tokyo, Springer-Verlag, 1991
10. Hathaway WE, Manco-Johnson M: Disorders of coagulation and platelets in the neonate. *In* *Hematology: Basic Principles and Practice*. Edited by R Hoffman, EJ Benz Jr, SJ Shattil, et al. New York, Churchill Livingstone, 1991, pp 1409-1415
11. Corrigan JJ Jr: Normal hemostasis in the fetus and newborn: coagulation. *In* *Fetal and Neonatal Physiology*. Vol. 2. Edited by RA Polin, WW Fox. Philadelphia, WB Saunders Company, 1992, pp 1368-1371