

TEST SUMMARY

Dysfibrinogenemia Functional Panel

This panel is for the assessment of fibrinogen function. Thrombin time may be prolonged due to afibrinogenemia, hypofibrinogenemia, dysfibrinogenemia, or the presence of heparin and direct thrombin inhibitors. Heparin contamination can be ruled out by adding protamine sulfate if thrombin time is prolonged. The clot-based tests in this panel are designed to be sensitive to fibrinogen disorders including dysfibrinogenemia, hypofibrinogenemia, hypodysfibrinogenemia and afibrinogenemia. In order to confirm the absence of heparins or DTIs, the thrombin time test reflexes to a heparin neutralization test when prolonged.

Please visit Machaondiagnostics.com for further information.

References: 1. Medved L et al. Fibrinogen and Factor XIII Subcommittee of Scientific Standardization Committee of International Society on Thrombosis and Haemostasis. Recommendations for nomenclature on fibrinogen and fibrin. J Thromb Haemost 2009; 7:355. 2. Tennent GA et al. Human plasma fibrinogen is synthesized in the liver. Blood 2007; 109:1971.3. Martinez J. Congenital dysfibrinogenemia. Curr Opin Hematol 1997; 4:357.4. Collen D et al. Metabolism and distribution of fibrinogen. I. Fibrinogen turnover in physiological conditions in humans. Br J Haematol 1972; 22:681.



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WHY CHOOSE US?

- Fastest turnaround in the US
- Weekend testing for STATs
- Oraw kits (includes free shipping)
- 🕥 Clinical consultation
- 🛇 Critical Results called to physician

SPECIMEN REOUIRMENTS

Three aliquots; 1mL each Citrated Plasma

STABILITY

Frozen (-20C): 2 weeks Frozen (-80C): 6 months

CPT CODF

METHODOLOGY

Clot-based

TURNAROUND TIMES

Routine TAT: <1 week STAT TAT: <24 hours (7 days a week)

PANEL COMPONENT TESTS

Fibrinogen Activity Thrombin Time aPTT Prothrombin Time (PT/INR)

ASSOCIATED TESTS

Dysfibrinogenemia Genetic Panel CoagGenex Clotting Genetic Panel

DRAW KITS AVAILABLE

ABOUT US

Machaon Diagnostics is a clinical reference laboratory, specializing in coagulation, platelets, complement, genetics and rare disease



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