Machaon Diagnostics

-coagulation, platelets, rare disease and genetics

Main Oakland Lab
Medical Director: Brad Lewis, MD
3023 Summit Street, Oakland, CA 94609
Phone: (510) 839-5600 / Fax: (510) 839-6153

Second Lab (**samples can be sent directly)
Medical Director: Gloria Coker, MD
8721 Oak Street, New Orleans, LA 70118
Phone: (504) 866-7090 / Fax: (504) 866-7091

www.MachaonDiagnostics.com
Toll-free: (800) 566-3462

PATIENT HISTORY

Patient's name: (Last, First, M.I.) | Specimen Date: (time)

Sex: (circle one) M / F | Date of birth: | SS# or ID#: | Referring physician:

Platelet Count_____ (K/µL), aPTT_____ (sec.), PT_____ (sec.), INR_____
Hematocrit_____ (%), Bleeding History_____ (Y/N), Clotting History_____ (Y/N)

Is patient anticoagulated? Circle: coumadin / LMWH / UFH / apixaban / rivaroxaban / dabigatran / fondaparinux / other _____. Is the patient on antiplatelet therapy? Circle: aspirin / Plavix / Brilinta / other _____. Is patient on Hemlibra therapy (Y / N)

SUBMITTING FACILITY

Pre-Printed forms available upon request.

Phone:

Fax for results:

☐ STAT ☐ ASAP ☐ ROUTINE ICD-10_____

PANEL TESTING

☐ Antiphospholipid Syndrome Criteria Panel (aPTT-LA, dRVVT, Anticardiolipin and Beta-2 Glycoprotein I Antibodies- IgG, IgM, IgA) ☐ Do not reflex to LA Panel

☐ Heparin Antibody Panel (TAT <24 Hrs)
(Immunologic [ELISA] and Functional [washed-platelet Heparin-induced Platelet Activation])

☐ Hypercoagulability / Thrombophilia Panel (STAT <24 Hrs)
(Please visit our website for complete testing algorithm)

☐ Lupus Anticoagulant Screen (ACL, dRVVT, aPTT-LA) ☐ Do not reflex to LA Panel
☐ Panel (ACL, aPTT Mixing Study, Lupus Anticoagulant Index, dRVVT, Thrombin Time and PT/INR)

☐ Mild Bleeding Work-up (most common tests) (STAT <24 Hrs)
(Platelet Aggregation, vWF Profile, Fibrinogen Activity, Thrombin Time, PT/INR)

☐ Prolonged aPTT / PT Evaluation Work-up (STAT <24 Hrs)
(Please visit our website for complete testing algorithm)

☐ von Willebrand Factor Profile ☐ Do not reflex to vWF:Multimer (STAT <24 Hrs)
(Factor VIII Activity, vWF:Antigen, vWF:RCo, aPTT and if indicated, vWF:Multimer)

SELECT GENETIC PANEL TESTING

☐ Alport Syndrome Genetic Panel (*3 genes*)
☐ aHUS Genetic Panel (STAT <2 Days)
☐ C3 Glomerulopathy Genetic Panel (*5 genes*)
☐ Dysfibrinogenemia Genetic Panel (*FGA, FGB, FGG*)
☐ Hemophagocytic Lymphohistiocytosis (HLH) Genetic Panel
☐ Hemophilia-Complete™ Genetic Panel (*F8, F9, VWF, inversions*)
☐ PlateletGenex™ Functional Defect Panel (*31 genes*)
☐ PlateletGenex™ Thrombocytopenia Panel (*26 genes*)
☐ Polycystic Kidney Disease (PKD) Genetic Panel (*2 genes*)
☐ TMA-Complete™ Genetic Panel (STAT <2 Days)
☐ vWD-Complete™ Genetic Panel (*VWF and GP1BA*)

TEST LIST

☐ ACL (Anticardiolipin - IgG, IgM and IgA)
☐ ADAMTS-13 Activity (reflexes to Inhib and Ab)
☐ ADAMTS-13 Panel (Activity, Inhibitor and Antibody)
☐ ADAMTS-13 Gene Sequencing
☐ Anti-CFH Autoantibody
☐ Antithrombin III Activity and/or ☐ Antigen
☐ Apixaban (Eliquis) Level
☐ Beta-2 Glycoprotein I Antibody
☐ Beta-2 Glycoprotein I - Domain 1 Antibody
☐ Euglobulin Clot Lysis Time
☐ Factor Activity (aPTT-based) ☐ test all factors
☐ VIII (8) ☐ IX (9) ☐ XI (11) ☐ XII (12)
☐ Factor Activity (PT-based) ☐ test all factors
☐ II (2) ☐ V (5) ☐ VII (7) ☐ X (10)
☐ Factor V (5) Leiden Gene Mutation
☐ Factor VIII (8) Chromogenic Activity (bovine)
☐ Factor VIII (8) Gene Sequencing and Inversions Assay
☐ Factor IX (9) Gene Sequencing
☐ Factor XIII (13) Activity and/or ☐ Gene Sequencing
☐ Fibrinogen Activity and/or ☐ Antigen
☐ Hemlibra-specific Factor VIII (8) Activity
☐ Heparin Antibody Confirmation (wp-HIPA)
☐ Heparin Antibody Reflex (ELISA reflex to wp-HIPA)
☐ Heparin Level (anti-Xa method) (indicate type above)
☐ Hexagonal Phospholipid (STACLOT-LA)
☐ Homocysteine
☐ HPP/OI Genetic Panel (Hypophosphatasia)
☐ Inhibitor to Factor(s)_____ (Bethesda Units)
☐ Inhibitor to F8 (Hemlibra-specific Nijmegen Bethesda)
☐ Mixing Study (aPTT) - reflex to incubated mix
☐ Mixing Study (PT)
☐ MTHFR ☐ C677T and/or ☐ A1298C) Mutations
☐ aPTT ☐ Reflex to work-up
☐ aPTT-LA (Lupus Sensitive Reagent)
☐ Plasminogen Activity and/or ☐ Autoantibody
☐ Plasminogen Gene Sequencing
☐ PAI-1 Activity and/or ☐ PAI-1 Gene Sequence
☐ Platelet Antibody ID: Direct (__) and/or Indirect (__)
☐ Platelet Aggregation Study - LTA**
☐ Platelet ATP / Granule Release Study**
☐ Platelet Aggregation - RIPA (Ristocetin-induced)**
☐ Platelet Electron Microscopy Study
☐ Plavix Sensitivity - LTA and/or ☐ Genotype Assay
☐ Protein C Activity and/or ☐ Antigen
☐ Protein S Activity
☐ Protein S Antigen [Free] and/or ☐ [Total]
☐ Prothrombin Fragment 1.2
☐ Prothrombin Gene Mutation
☐ Prothrombin Time ☐ Reflex to work-up
☐ PS Antibody (phosphotidylserine; IgG, IgM)
☐ PS/PT Antibody (IgG, IgM)
☐ Rivaroxaban (Xarelto) Level
☐ dRVVT (dilute Russel Viper Venom Time)
☐ Soluble Complement 5b-9 (sC5b-9)
☐ Thrombin Generation (profile w/ ETP)
☐ Thrombin Time - TCT (confirmed w/ PS)
☐ Thrombin-Antithrombin (TAT)
☐ Thrombophilia Genetic Panel (*11 genes*)
☐ Thromboelastography (TEG)
☐ Thromboelastometry (ROTEM)
☐ TEG - Platelet Mapping
☐ vWF Activity (Ristocetin cofactor)
☐ vWF Activity (GP1BM-based)
☐ vWF Antigen
☐ vWF Multimer (vWF:Multimer)
☐ wp-HIPA (washed-platelet Heparin-induced Platelet Activation Assay)

Misc._____

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ADDITIONAL INFORMATION

Patients with insurance coverage other than Medicare are considered *out-of-network* and will be billed for services not covered by their insurance provider. Medicare patients must sign an ABN, either located on the reverse side of this form or downloaded from the Machaon Diagnostics website. Patient insurance billing services are provided in accordance with the *Machaon Insurance Billing Policy*. HMO or medical group covered patients may need a prior authorization if they seek reimbursement.

MACHAON USE ONLY

Specimen type received_____ Aliquots_____

Specimen type received_____ Aliquots_____

Tech initials_____ Specimen received stamp_____