

Client Information (required)

Client Name		
Client Account No.		
Client Phone	Client Order No.	
Address		
City	State	ZIP Code

Submitting Provider Information (required)

Submitting/Referring Provider <i>(Last, First)</i>
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Fill in only if Call Back is required.

Phone (with area code)	Fax (with area code)
Provider's National I.D. (NPI)	

**Fax number given must be from a fax machine that complies with applicable HIPAA regulation.*

Reason for Testing (required)

ICD-10 Diagnosis Code

Note: It is the client's responsibility to maintain documentation of the order.

New York State Patients: Informed Consent for Genetic Testing

"I hereby confirm that informed consent has been signed by an individual legally authorized to do so and is on file with this office or the individual's provider's office."

Signature ▶

Note: It is the client's responsibility to maintain documentation of the order.

Ship specimens to:

Mayo Clinic Laboratories
3050 Superior Drive NW
Rochester, MN 55901

Customer Service: 855-516-8404

Visit www.MayoClinicLabs.com for the most up-to-date test and shipping information.

Patient Information (required)

Patient ID (Medical Record No.)		
Patient Name <i>(Last, First, Middle)</i>		
Sex <input type="checkbox"/> Male <input type="checkbox"/> Female	Birth Date <i>(mm-dd-yyyy)</i>	
Collection Date <i>(mm-dd-yyyy)</i>	Time	<input type="checkbox"/> am <input type="checkbox"/> pm
Street Address		
City	State	ZIP Code
Phone		

Insurance Information

Subscriber's Name (if different than patient)		
Patient Relationship <input type="checkbox"/> Spouse <input type="checkbox"/> Dependent <input type="checkbox"/> Other: _____		
Medicare HIC Number (if applicable)		
Medicaid Number (if applicable)		
Insurance Company's Name (if applicable)		
Insurance Company's Street Address		
City	State	ZIP Code
Policy Number		
Group Number		

MCL Internal Use Only

Billing Information

- An itemized invoice will be sent each month.
- Payment terms are net 30 days.

Call the Business Office with billing related questions:
800-447-6424 (US and Canada)
507-266-5490 (outside the US)

Patient Information (required)

Patient ID (Medical Record No.)	Client Account No.
Patient Name <i>(Last, First, Middle)</i>	Client Order No.
Birth Date <i>(mm-dd-yyyy)</i>	

CONSULTATION/MORPHOLOGY EVALUATION

PATHC Pathology Consultation (submit stained slides and block)

HPWET Hematopathology Consultation, MCL Embed (submit core biopsy, clot section and bone marrow aspirate)

HPCUT Hematopathology Consultation, Client Embed (submit bone marrow aspirate and embedded core biopsy and clot section)

ERYTHROCYTES: ENZYMOPATHIES

AK1 Adenylate Kinase Enzyme Activity, Blood

G6PD1 Glucose 6-Phosphate Dehydrogenase Enzyme Activity, Blood

GPI1 Glucose Phosphate Isomerase Enzyme Activity, Blood

G6PDB Glucose-6-Phosphate Dehydrogenase (G6PD) Full Gene Sequencing, Varies

GSH Glutathione, Blood

HAEV1* Hemolytic Anemia Evaluation, Blood

HK1 Hexokinase Enzyme Activity, Blood

PFK1 Phosphofructokinase Enzyme Activity, Blood

PGK1 Phosphoglycerate Kinase Enzyme Activity, Blood

P5NT Pyrimidine 5' Nucleotidase, Blood

PK1 Pyruvate Kinase Enzyme Activity, Blood

PKLRG Pyruvate Kinase Liver and Red Blood Cell (PKLR), Full Gene Sequencing and Large Deletion Detection, Varies

EEEV1* Red Blood Cell (RBC) Enzyme Evaluation, Blood

TPI1 Triosephosphate Isomerase Enzyme Activity, Blood

ERYTHROCYTES: GENERAL

HGBQ Hemoglobin, Qualitative, Random, Urine

PLHBB Plasma Free Hemoglobin, Plasma

RTIC Reticulocytes, Blood

ERYTHROCYTES: HEMOGLOBIN DISORDERS

WASEQ Alpha Globin Gene Sequencing, Varies

WBDD Beta-Globin Cluster Locus, Deletion/Duplication, Varies

WBSEQ Beta Globin Gene Sequencing, Varies

REVE1* Erythrocytosis Evaluation, Whole Blood

WGSEQ Gamma-Globin Full Gene Sequencing, Varies

HBEL1* Hemoglobin Electrophoresis Evaluation, Blood

HGBCE* Hemoglobin Variant, A2 and F Quantitation, Blood

HAEV1* Hemolytic Anemia Evaluation, Blood

MEV1* Methemoglobinemia Evaluation, Blood

SDEX Sickle Solubility, Blood

THEV1* Thalassemia and Hemoglobinopathy Evaluation, Blood and Serum

ERYTHROCYTES: HEREDITARY ERYTHROCYTOSIS

BPGMM 2,3-Bisphosphoglycerate Mutase, Full Gene Sequencing Analysis, Varies

REVE1* Erythrocytosis Evaluation, Whole Blood

EPO Erythropoietin, Serum

HEMP Hereditary Erythrocytosis Mutations, Whole Blood

ERYTHROCYTES: IMMUNOLOGY

ABYSR Antibody Screen with Reflexed Antibody Identification, RBC

CATR Cold Agglutinin Titer, Serum

BTR Isoagglutinin Titer, Anti-B, Serum

PLINK Paroxysmal Nocturnal Hemoglobinuria, PI-Linked Antigen, Blood

ERYTHROCYTES: MEMBRANE DISORDERS

HAEV1* Hemolytic Anemia Evaluation, Blood

KCNN4 KCNN4 Full Gene Sequencing, Varies

FRAG Osmotic Fragility, Erythrocytes

RBCME* Red Blood Cell Membrane Evaluation, Blood

ERYTHROCYTES: METHEMOGLOBIN

METR1 Cytochrome b5 Reductase Enzyme Activity, Blood

MET Methemoglobin and Sulfhemoglobin, Blood

MEV1* Methemoglobinemia Evaluation, Blood

LEUKOCTYES

MURA Lysozyme (Muramidase), Plasma

LYMPHOCYTES

ALPS Alpha Beta Double-Negative T Cells for Autoimmune Lymphoproliferative Syndrome, Blood

CRGSP Cryoglobulin and Cryofibrinogen Panel, Serum and Plasma

ATR Isoagglutinin Titer, Anti-A, Serum

LCMS Leukemia/Lymphoma Immunophenotyping, Flow Cytometry, Varies

NKSP Natural Killer (NK)/Natural Killer T (NKT) Cell Subset Panel, Blood

SVISC Viscosity, Serum

METABOLISM: MEGALOBLASTIC ANEMIA

FOL Folate, Serum

GAST Gastrin, Serum

MHCZ Methylmalonic Aciduria and Homocystinuria, cblC Type, Full Gene Analysis, Varies

MHDZ Methylmalonic Aciduria and Homocystinuria, cblD Type, Full Gene Analysis, Varies

MMAP Methylmalonic Acid, Quantitative, Plasma

MMAS Methylmalonic Acid, Quantitative, Serum

MMAU Methylmalonic Acid, Quantitative, Urine

ACASM Pernicious Anemia Cascade, Serum

B12 Vitamin B12 Assay, Serum

FB12 Vitamin B12 and Folate, Serum

METABOLISM: METALS

CERS Ceruloplasmin, Serum

FERR Ferritin, Serum

FECHZ Ferrochelatase (FECH) Gene, Full Gene Analysis, Varies

HFE Hemochromatosis HFE Gene Analysis, Blood

FEC Iron and Total Iron-Binding Capacity, Serum

TRSF Transferrin, Serum

NEZPP Zinc Protoporphyrin, Blood

MOLECULAR BENIGN HEMATOLOGY STUDIES

NGHHA Hereditary Hemolytic Anemia Comprehensive Panel, Next-Generation Sequencing, Varies

NGCDA Congenital Dyserythropoietic Anemia Panel, Next-Generation Sequencing, Varies

NGMEM Red Blood Cell Membrane Panel, Next-Generation Sequencing, Varies

NGENZ Red Blood Cell Enzyme Panel, Next-Generation Sequencing, Varies

KCNN4 KCNN4 Full Gene Sequencing

ADDITIONAL TESTS (INDICATE TEST ID AND NAME)

*Metabolic Hematology Patient Information (T810) is strongly recommended for this test. For more information, see corresponding test on MayoClinicLabs.com