

**FOR ALL TESTS provide the following**

Patient Name	Patient Identification Number
Referring Physician Name	Phone Number (Include area code)

Attach bar-coded patient label here.

**HEMATOPATHOLOGY – Include the following information and send a copy of Bone Marrow and/or Blood Smear reports**

Specimen Submitted	
Patient's Ethnic or Racial Background	
Recent Transfusion History	
Is there a family history? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	
Is there a history of Splenomegaly? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	
<b>CBC RESULTS:</b>	<b>Check appropriate statement:</b>
HB _____ MCV _____	<input type="checkbox"/> Acute Lymphoblastic Leukemia
HCT _____ WBC _____	<input type="checkbox"/> Chronic Lymphocytic Leukemia
RBC _____ PLT _____	<input type="checkbox"/> Acute Myeloid Leukemia
Reticulocyte Count (if available) _____	<input type="checkbox"/> Chronic Myelogenous Leukemia
	<input type="checkbox"/> Chronic Myeloproliferative Disease
	<input type="checkbox"/> Lymphoma
	<input type="checkbox"/> Plasma Cell Proliferative Disease
	<input type="checkbox"/> Other: _____
Pertinent Clinical Information	

**DERMATOLOGY  
8041 Cutaneous Immunofluorescence, Biopsy**

Biopsy Site
Clinical Impression
Check One: <input type="checkbox"/> Lesional (Involved) <input type="checkbox"/> Perilesional <input type="checkbox"/> Uninvolved
Check One: <input type="checkbox"/> Sun Exposed <input type="checkbox"/> Non-Sun Exposed

**SURGICAL CONSULTATION – Include a brief history, pertinent lab results and suspected diagnosis or indicate in space provided below**

Tissue Source	Patient Birth Date (Month DD, YYYY)
Specimen Sent (check all that apply)	
<input type="checkbox"/> Fixed Formalin	<input type="checkbox"/> Paraffin Block(s), Number Sent: _____
<input type="checkbox"/> Frozen Tissue	<input type="checkbox"/> Slide(s), No. Sent: _____
<input type="checkbox"/> Gluteraldehyde	<input type="checkbox"/> Zeus Media
<input type="checkbox"/> Wet Tissue	<input type="checkbox"/> Other: _____
Pathologist/Clinical Diagnosis (or send copy of pathology report)	

**LABORATORY GENETICS – Biochemical Genetics, Cytogenetics\***

\*Denotes the only information required for Cytogenetics testing.

*Reason for Ordering Test(s)	*Relevant Clinical Information				
Is there a family history of a similar condition? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	Has the patient or a family member had this test before? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown				
If yes to either of the above questions, complete the following (if more than two individuals, list on additional sheet of paper):					
<b>Relationship to Patient</b>	<b>Affected</b>	<b>Carrier</b>	<b>Test Result(s)</b>	<b>Check if Tested at Mayo</b>	<b>Name (Optional)</b>
_____	<input type="checkbox"/>	<input type="checkbox"/>	_____	<input type="checkbox"/>	_____
_____	<input type="checkbox"/>	<input type="checkbox"/>	_____	<input type="checkbox"/>	_____
Patient's Ethnic or Racial Background					
Is patient currently pregnant? <input type="checkbox"/> Yes <input type="checkbox"/> No	If yes, complete the following: Estimated Gestational Age _____ Weeks on (Date) _____ By: <input type="checkbox"/> LMP <input type="checkbox"/> Ultrasound <input type="checkbox"/> Physical Exam				
<b>ADDITIONAL INFORMATION for Biochemical Genetics Tests</b>					
Current Medications/Diet	<input type="checkbox"/> Valproic Acid	<input type="checkbox"/> Carbamazepine	<input type="checkbox"/> Carnitine	<input type="checkbox"/> Oral Contraceptives	<input type="checkbox"/> TPN <input type="checkbox"/> Special Diet
	<input type="checkbox"/> Other (specify): _____				
Check if Applicable	<input type="checkbox"/> Acute Illness	<input type="checkbox"/> Asymptomatic	<input type="checkbox"/> Follow-Up (specify disorder above)		
	<input type="checkbox"/> Repeat Specimen (specify previous findings above)		<input type="checkbox"/> Post-Mortem Specimens		

**NEW YORK STATE PATIENTS: INFORMED CONSENT APPLICABLE TO HIV AND GENETIC TESTING**

The client submitting this request has received reasonable assurance from the ordering physician that the above named New York State patient has given informed consent for the HIV and/or genetic testing ordered and that the patient authorizes MML to report such test results directly to the ordering physician.

**FOR AFP AND COAGULATION TESTING provide the following**

Patient Name	Patient Identification Number
Referring Physician Name	Phone Number (Include area code)

Attach bar-coded patient label here.

**SECOND TRIMESTER SCREENING (QUAD, MAFP) – The following 10 questions MUST be completed for Second Trimester Screening**

<b>1 Serum Collection Date</b> (Month DD, YYYY)	<b>7 In-Vitro Fertilization (IVF) Pregnancy</b> <input type="checkbox"/> Yes <input type="checkbox"/> No (The age of the egg affects the risk calculations). If egg donor (other than patient), need donor birth date (Month DD, YYYY) _____ If frozen egg or embryo used, how long was egg or embryo frozen _____ (Years) _____ (Months)
<b>2 Birth Date</b> (Month DD, YYYY)	
<b>3 Weight</b> lbs _____ or _____ kg	<b>8 Previous pregnancy with Down Syndrome (Trisomy 21) or other Trisomy?</b> <input type="checkbox"/> Yes <input type="checkbox"/> No
<b>4 Insulin Dependent Diabetic</b> <input type="checkbox"/> Yes <input type="checkbox"/> No	<b>9 Is this a repeat visit?</b> <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, list previous control number _____
<b>5 Race</b> <input type="checkbox"/> Black <input type="checkbox"/> Other/Non-Black/Mixed	<b>10 Gestational Information: EDD</b> (Month DD, YYYY) _____ by <input type="checkbox"/> Ultrasound or <input type="checkbox"/> LMP If LMP is unknown and ultrasound has not been performed, provide other dating, such as physical exam or IVF. Please be specific.
<b>6 Twin Pregnancy</b> (US EDD required) <input type="checkbox"/> Yes <input type="checkbox"/> No	

**COAGULATION**

Identify the coagulation diagnostic concern of other relevant information:

<b>Coagulation-related TESTING RESULTS from referring laboratory</b> PT _____ Normal Range _____ APTT _____ Normal Range _____ Platelet Count _____ Hematocrit _____ Other _____	<b>Does the patient have:</b> Known congenital coagulation factory deficiency? <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, which factor? _____ Known coagulation factor inhibitor? <input type="checkbox"/> Yes <input type="checkbox"/> No (If type of inhibitor is unknown we suggest ordering consult #553) If yes, which factor? _____
Coagulation-related MEDICATION, current or past 7 days? <input type="checkbox"/> Coumadin (Warfarin) <input type="checkbox"/> Vitamin K <input type="checkbox"/> Heparin (Unfractionated) <input type="checkbox"/> Low Molecular Weight Heparin <input type="checkbox"/> Lepirutin (Refludan) <input type="checkbox"/> Thrombolytic (t-PA) <input type="checkbox"/> Fondaparinux (Arixtra) <input type="checkbox"/> Argatroban	<b>For DNA Based Testing, has patient had:</b> Transfusion within the past 3 months? <input type="checkbox"/> Yes <input type="checkbox"/> No Bone marrow transplant? <input type="checkbox"/> Yes <input type="checkbox"/> No Liver transplant? <input type="checkbox"/> Yes <input type="checkbox"/> No
Transfusion of Factor Replacement, past 72 hours? <input type="checkbox"/> Yes <input type="checkbox"/> No Factor Concentrate – Specify product _____ <input type="checkbox"/> DDAVP <input type="checkbox"/> Cryoprecipitate <input type="checkbox"/> Fresh Frozen Plasma	<b>von Willebrand Testing Information</b> Ristocetin Cofactor Activity _____ Normal _____ Factor VIII Activity Results _____ Normal _____ von Willebrand Factor Ag Result _____ Normal _____

**MICROBIOLOGY**

<b>Isolated Organism Referred for Identification</b> – All of the following information must be submitted to obtain identification of any organism submitted	<b>Antimicrobial Susceptibility</b>
Source	Source
Number of times isolated from different specimens (same patient)	Organism Identification – (If not known, add appropriate ID test)
Recovery Medium	Transport Medium
Description (Gram Reaction, Morphology, Tests Performed)	Antibiotic to be Tested (if applicable)
Extent of Identification Request	