

Patient Name (last, first):	Sex: <input type="checkbox"/> M <input type="checkbox"/> F	Client Name:
Address:	Medical Record #:	Address:
City/State/Zip:		City/State/Zip:
Home Phone:		Phone: Fax:
Date of Birth (MM/DD/YY):		Referring Physician:
Clinical Indications / Family History (attach pedigree, if appropriate):		DIAGNOSTIC CODE(S) (Required):

<p>MICROARRAY ANALYSIS (Oligo-SNP) (CMA and CMAO)</p> <p><input type="checkbox"/> Microarray Constitutional BLOOD: 4cc sodium heparin (green top) + 4cc EDTA (lavender top)</p> <p><input type="checkbox"/> Microarray Oncology BONE MARROW / PERIPHERAL BLOOD: 3cc sodium heparin (green top) TUMOR: 2mm² in sterile media (RPMI or sterile saline solution)</p> <p>CONSTITUTIONAL CYTOGENETICS (CS) KARYOTYPE / CHROMOSOME ANALYSIS BLOOD: 3cc sodium heparin (green top) or TISSUE: 3x3 mm² in Sterile Media</p> <p><input type="checkbox"/> Routine Chromosomes <input type="checkbox"/> Neonatal Chromosomes <input type="checkbox"/> STAT <input type="checkbox"/> Mosaic Chromosome Study <input type="checkbox"/> Skin <input type="checkbox"/> Autopsy <input type="checkbox"/> Other 3x3 mm² in Sterile Media</p> <p><input type="checkbox"/> Chromosomes <input type="checkbox"/> Cryopreservation <input type="checkbox"/> Send out testing for: _____</p> <p>FISH - CONSTITUTIONAL (CS) FISH ANALYSIS BLOOD: 3cc sodium heparin (green top)</p> <p><input type="checkbox"/> 1p36 deletion Syndrome <input type="checkbox"/> Ambiguous genitalia Sex XX / XY <input type="checkbox"/> Angelman Syndrome 15q <input type="checkbox"/> Cri du Chat Syndrome 5p <input type="checkbox"/> DiGeorge/VCF Syndrome 22q <input type="checkbox"/> Kallmann Syndrome Xp <input type="checkbox"/> Miller-Dieker Syndrome 17p <input type="checkbox"/> Neurofibromatosis (NF1) 17q <input type="checkbox"/> Prader-Willi Syndrome <input type="checkbox"/> Smith-Magenis Syndrome 17p <input type="checkbox"/> Sotos Syndrome 5q <input type="checkbox"/> SRY Yp Male determination <input type="checkbox"/> Steroid Sulfatase (STS) deficiency Xp <input type="checkbox"/> Trisomy 13 <input type="checkbox"/> Trisomy 18 <input type="checkbox"/> Trisomy 21 <input type="checkbox"/> Williams Syndrome 7q <input type="checkbox"/> Wolf-Hirschhorn Syndrome 4p <input type="checkbox"/> FISH Not otherwise specified: _____</p>	<p>ONCOLOGY CYTOGENETICS (CSO) KARYOTYPE / CHROMOSOME ANALYSIS BONE MARROW/BLOOD: 3cc sodium heparin (green top)</p> <p><input type="checkbox"/> Bone Marrow Chromosomes <input type="checkbox"/> Leukemic Blood Chromosomes <input type="checkbox"/> Solid Tumor 3x3 mm² in Sterile Media <input type="checkbox"/> Lymphoma 3x3 mm² in Sterile Media</p> <p>FISH - ONCOLOGY (CSO) FISH ANALYSIS BONE MARROW / PERIPHERAL BLOOD: 3cc sodium heparin (green top) TUMOR touch prep slides: prepare 2 to 3 slides by gently touching tumor surface (to cover ~15mmx15mm area) to middle of slide; immediately transfer slide to 3:1 methanol: acetic acid (Carnoy's fix) for 5 mins., Air dry</p> <p>ONCOLOGY FISH PANELS</p> <p><input type="checkbox"/> ALL FISH Panel (please specify): <input type="checkbox"/> B-CELL <input type="checkbox"/> High Risk B-CELL <input type="checkbox"/> T-CELL</p> <p><input type="checkbox"/> AML FISH Panel <input type="checkbox"/> AML FISH Panel + PML/RARA <input type="checkbox"/> B-CELL Lymphoma FISH Panel <input type="checkbox"/> BRAIN/CNS Tumor Panel <input type="checkbox"/> EOSINOPHILIA FISH Panel <input type="checkbox"/> MDS FISH Panel <input type="checkbox"/> MEDULLOBLASTOMA Panel <input type="checkbox"/> NEUROBLASTOMA Panel <input type="checkbox"/> PH-Like ALL FISH Panel <input type="checkbox"/> SARCOMA – 3 probe Panel <input type="checkbox"/> TUMOR (NOS) – Custom Panel - 5 probes</p> <p>INDIVIDUAL PROBES Please inquire <input type="checkbox"/> FISH Not otherwise specified: _____</p>	<p>BONE MARROW TRANSPLANTATION MONITORING FISH: BONE MARROW / PERIPHERAL BLOOD: 3cc sodium heparin (green top)</p> <p><input type="checkbox"/> Post BMT XX/XY by FISH</p> <p>STR ANALYSIS BONE MARROW / PERIPHERAL BLOOD: 0.5cc EDTA (lavender)</p> <p>BMD <input type="checkbox"/> Donor Specimen STRPRE <input type="checkbox"/> Pre Transplant Recipient STRPOST <input type="checkbox"/> Post-Transplant Recipient</p> <p>LINEAGE SPECIFIC STR ANALYSIS BLOOD: 8.5 cc ACD (yellow top) BONE MARROW: 4cc ACD (yellow top)</p> <p>STRPOST1 <input type="checkbox"/> Post-Transplant, Lineage-Specific (check all that apply) <input type="checkbox"/> Whole <input type="checkbox"/> T-cell <input type="checkbox"/> B-cell <input type="checkbox"/> Myeloid <input type="checkbox"/> NK</p>
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SPECIMEN COLLECTION DATE: _____ TIME: _____ DRAWN BY: _____

FORM COMPLETED BY: _____

PATIENT ID: _____

CLIA ID: 10D0700790, CAP: 153609-01

UNLESS OTHERWISE SPECIFIED - SHIP AT ROOM TEMP TO:
Johns Hopkins All Children's Hospital
601 Fifth Street South
St. Petersburg, FL 33701
ATTN: Pathology and Laboratory Medicine, Cytogenetics Dept. 6500001504

PHYSICIAN SIGNATURE: _____ DATE: _____

WHITE: ORDERING UNIT YELLOW: LAB