

Org ID#: 00 _____

*Organization Name: _____

Organization Address: _____

City: _____ St.: _____ Zip: _____

*Phone: _____ *Fax: _____

Ordering Provider Name (print): _____

Use full provider name. No nicknames or initials
***All information required for valid order**

Provider Signature _____

Additional report to Provider _____ Fax _____

Order Comment _____



SPECTRUM HEALTH

**LABORATORY REQUISITION
ADVANCED TECHNOLOGY**

*DATE ORDERED	DATE COLLECTED	TIME COLLECTED	INITIALS COLLECTED
*PATIENT INFORMATION			
NAME LAST		FIRST	MIDDLE INT.
ADDRESS			PHONE
CITY		STATE	ZIP
SEX	MARITAL STATUS	BIRTH DATE	CELL PHONE
*DIAGNOSIS/ICD CODES			
1st _____			
2nd _____			
Other _____			
INSURANCE INFORMATION			
Providers submitting SPECIMEN orders please attach copy(s) of Insurance cards (front and back). Please indicate Primary and Secondary.			
<input type="checkbox"/> Amniotic Fluid** <input type="checkbox"/> Chorionic Villi** **EDD _____/_____/_____ by <input type="checkbox"/> LMP <input type="checkbox"/> U/S **Gest Age _____ wks Biochemical Markers (Amniotic Fluid Only) 1066 <input type="checkbox"/> Alpha-Fetoprotein 8115 <input type="checkbox"/> Acetylcholinesterase			

SAMPLE TYPE

- Blood Bone Marrow Lymph Node Skin Biopsy
- Solid Tumor, type _____ Fluid, type _____
- Tissue Surgical # _____ Type _____
- Products of Conception* Fetal Autopsy*
- Tissue Type(s) _____ *Gest Age _____ wks

CYTOGENETICS

Fluorescence In-Situ Hybridization (FISH)*

*Diagnostic bone marrow specimens include chromosome study

Hematologic/Neoplastic Disorders

- 9227 Monitor previously identified abnormalities
- 943 ALK rearrangement
- 990 AML panel
- 7145 B-ALL panel
- 946 BCR/ABL1
- 961 Burkitt lymphoma panel
- 949 CHOP (DDIT3) rearrangement
- 992 CLL panel
- 9223 Diffuse Large B-cell lymphoma panel
- 9224 Eosinophilia panel
- 951 EWSR1 rearrangement
- 952 FKHR (FOXO1) rearrangement
- 953 Follicular lymphoma panel
- 954 FUS rearrangement
- 955 Glioma (1p36/19q13)
- 956 Her-2/neu (ERBB2) (Breast Cancer)
- 965 Mantle Cell Lymphoma CCND1/IGH
- 994 MDS panel
- 9225 MDM2 amplification
- 7031 MET amplification
- 966 MLL (11q23) rearrangement
- 967 MYC (8q24) rearrangement
- 993 Myeloma panel
- 968 N-MYC amplification
- 948 PDGFRA (CHIC2)
- 970 PML/RARA
- 7032 RET Gene Arrangement
- 7033 ROS1 Gene Arrangement
- 996 T-ALL panel
- 9222 XX/XY (BMT status)
- Other _____

Constitutional Deletions/Duplications

- 979 DiGeorge, VCFS (22q11.2)
- 974 Gender XX/XY (includes SRY)
- 995 POC aneuploid screen (XY, 13, 16, 18, 21)
- 983 Prader Willi/Angelman
- 991 Prenatal aneuploid screen (XY, 13, 18, 21)
- 984 Smith Magenis
- 980 Trisomy 21
- 705 Turner Syndrome
- 981 Williams/Beuren (7q11.23)
- Other _____

Chromosomal Microarray (aCGH)

- 221 Microarray, chromosomal (aCGH)
- 222 Microarray, confirmatory FISH

CYTOGENETICS (continued)

Chromosome Analysis (Sodium Heparin)

- 926 Hematologic/Neoplastic Study*
- *For bone marrow, provide WBC and Diff
- Lymphoma or other B-cell process suspected?
- Yes No

Constitutional Study

- 923 Routine
- 922 Prenatal
- 925 Mosaic
- 924 High Resolution (Blood only)
- 927 Products of Conception (POC)*
- *Tissue Pathology required

Other

- 928 Breakage Study for Fanconi's Anemia
- 929 Fibroblast Culture for Cryopreservation*
- 930 Fibroblast Culture for Send Out* (includes cryopreservation)
- *Chromosome study not included unless specified above

FLOW CYTOMETRY

Immunophenotyping

- 910 Leukemia/Lymphoma/Myeloma
- 4061 Complete Immune Competency Panel (CD3, CD4, CD8, CD16/CD56, CD19)
- 4058 CD4 & CD8 (CD3, CD4, CD8, CD19)
- 4059 CD4 Absolute Count (% & cells per mm3)
- 4060 CD20 (CD19, CD20)
- 4190 Fetal Cells by Flow Cytometry
- 4198 HLA B27 Screen
- 913 Neuroblastoma
- 915 Paroxysmal Nocturnal Hemoglobinuria (PNH)
- 707 Severe Combined Immunodeficiency Disorder (SCID) Screen
- 708 Autoimmune Lymphoproliferative Syndrome (ALPS) Screen

Functional Tests

- 917 Oxidative Burst
- 911 Leukocyte Adhesion Deficiency (LAD)

Other - Call 486.6273

- 921 Cell Sort for CD3 _____
- 921 Cell Sort for CD15 _____
- Other _____

MOLECULAR DIAGNOSTICS

Bone Marrow Engraftment Testing

- 901 Pre Bone Marrow Engraftment - Donor
- 902 Pre Bone Marrow Engraftment - Recipient
- 903 Post Bone Marrow Engraftment
- 702 Killer Cell Immunoglobulin-Like Receptors (KIR)

MOLECULAR DIAGNOSTICS (continued)

Provider responsible for documenting informed consent prior to ordering genetic testing.

Infectious Disease

- 7044 APTIMA Chlamydia NAAT
- 7045 APTIMA Gonococcus NAAT
- 7046 APTIMA Chlamydia Gonococcus NAAT
- 7043 Trichomonas NAAT (APTIMA)
- 241 Human Papilloma Virus (HPV) High Risk Screen
- 251 Human Papilloma Virus (HPV) 16/18/45 Genotyping
- 7019 Adenovirus Quant PCR
- 9028 Cytomegalovirus (CMV) Quantitative PCR
- 704 Epstein-Barr Virus (EBV) DNA Quantitative PCR
- 230 Hepatitis B Virus (HBV) DNA Quantitation by PCR
- 4185 Hepatitis C Virus (HCV) RNA Quantitation by RT-PCR
- 4434 Hepatitis C Virus (HCV) Genotype
- 7142 Hepatitis C Virus (HCV) 1a NS5A
- 7125 HIV-1 Quantitative RNA, PCR
- 4189 Herpes Simplex Virus (HSV) PCR (CSF and Plasma)

Inherited Disease

- 900 Cystic Fibrosis Carrier Screen
- 79 Hemochromatosis (HFE C282Y & H63D)
- 80 PraderWilli/Angelman mPCR
- 77 Factor V Leiden
- 76 MTHFR C677T
- 78 Prothrombin (Factor II) G20210A

Oncology Testing

- 7146 Epi proColon
- 7094 Colon Mutation Analysis
- 7030 Lung Cancer Mutation Analysis
- 709 BRAF Mutation Analysis
- 554 EGFR Mutation Analysis
- 710 KRAS Mutation Analysis
- 711 NRAS Mutation Analysis
- 904 Microsatellite Instability (MSI) PCR
- 7091 MLH1 Promoter Hypermethylation
- 706 MGMT Methylation Analysis
- 703 Cancer Hotspot Analysis
- 7060 ODC1 Genotype Analysis
- 9011 BCR-ABL1 t[9;22] RT-PCR
- 7135 JAK2 V617F mutation analysis, do MPN if negative
- 7090 MPN Expanded Panel
- 7014 IDH 1 & 2 Mutation Analysis
- 7130 CEBPA Mutation Analysis
- 7128 KIT Mutation Analysis
- 7129 NPM1 Mutation Analysis
- 7134 MYD88 Mutation Analysis
- 7120 Heme Molecular Sequence Analysis
- 107 IgH PCR for B-Cell Clonality
- 108 TCR PCR for T-Cell Clonality

NOTE: FOR MEDICARE PATIENTS, PLEASE ATTACH COMPLETED ABN FORM CMS-R 131 AS NEEDED

MEDICARE GENERALLY DOES NOT COVER ROUTINE SCREENING TESTS, CERTAIN EXCEPTIONS MAY APPLY

***INDICATES INFORMATION REQUIRED FROM OFFICE FOR VALID WRITTEN ORDER SPECTRUM HEALTH LABORATORY SERVICES 616.774.7721**

White - Laboratory Yellow - Laboratory Pink - Ordering Provider

LAB480 (2/17)