



## Solid Tumor Oncology

### Anatomic Pathology

#### Morphology

Surgical Pathology Consultation [SPC]

#### Immunohistochemistry & Special Stains

Refer to A-Z listing for IHC Markers

### Cytogenetics

Chromosome Analysis Solid Tumor [CYTO ST]

### Molecular Oncology

#### Next Generation Sequencing (NGS)

*BRAF* Mutation Analysis [M BRAF]

Colorectal Cancer NGS Panel [M COLON NGS]

*BRAF, KRAS, NRAS*

*EGFR* Mutation Analysis [M EGFR]

GIST NGS Panel - *KIT, PDGFRA* [M ST GIST NGS]

*IDH1/IDH2* Mutation Analysis [M IDH1/2]

*KRAS* Mutation Analysis [M KRAS]

*KIT* Mutation Analysis [M KIT]

Lung Cancer NGS Panel [M LUNG NGS]

*BRAF, EGFR, KRAS, NRAS*

Melanoma Cancer NGS Panel [M MELAN NGS]

*BRAF, KIT, NRAS*

Solid Tumor 8 Gene NGS Panel [M 8GENE NGS]

*BRAF, EGFR, IDH1, IDH2, KIT, KRAS, NRAS, PDGFRA*

### Fluorescence in situ Hybridization

*ALK* 2p23 rearrangement [FP ALK]

*BCL6* 3q27 rearrangement [FP BCL6]

Burkitt / "Double Hit" Large B-cell Lymphoma Panel [FP BURKITT]

*C-MYC, BCL2, BCL6*

*C-MYC* 8q24 rearrangement [FP CMYC]

*EGFR* amplification 7p11.2 [FPEGFR]

*HER2/neu* Breast Cancer [FP HER2]

*HER2/neu* Gastric Cancer [FP HER2 GA]

*IGH* 14q32 rearrangement [FP IGH]

*IGH/BCL1 (CCND1)* t(11;14) [FP BCL1]

*IGH/BCL2* t(14;18) [FP BCL2]

*IGH/MALT1* t(14;18) [FP IGH MALT]

*IGH/MYC* t(8;14) [FP IGH MYC]

*MALT1* 18q21 rearrangement [FP MALT]

Oligodendroglioma, 1p-, 19q- [FP GLI]

*ROS1* 6q22.1 rearrangement [FP ROS1]

## Hematology Oncology

### Anatomic Pathology

#### Morphology

Bone Marrow Pathology Evaluation [BMPE]

Surgical Pathology Consultation [SPC]

#### Immunohistochemistry & Special Stains

Refer to A-Z listing for IHC Markers

### Cytogenetics

Chromosome Analysis Bone Marrow [CYTO BM]

Chromosome Analysis Lymphoma (Lymph Node) [CYTO LN]

Chromosome Analysis Leukemic Peripheral Blood [CYTO LPB]

### Flow Cytometry

Flow Cytometry Leukemia / Myeloma / Lymphoma [FLOW]

Flow Cytometry Technical Only [FLOW TC]

Flow Cytometry and Morphology [FLOW M]

Bronchoalveolar Lavage [FLOW BAL]

CD4:CD8 ratio

Leukocyte Adhesion Deficiency [FLOW LAD]

CD11a, CD11b, CD11c, CD18

Paroxysmal Nocturnal Hemoglobinuria - High Sensitivity [FLOW PNH]

FLAER, CD14, CD24, CD59

### Molecular Oncology

#### Next Generation Sequencing (NGS)

Calreticulin type 1 / type 2 Mutation Analysis [M CALR]

*JAK2* Exon 12 Mutation [M JAK2 EX12]

*MPL* Exon 10 Mutation [M MPL]

*MYD88* p.L265P Mutation [M MYD88]

Myeloproliferative Neoplasms Core Mutation Panel [M MPN Core]

*JAK2 V617F, JAK2 EXON 12, CALR, MPL*

Myeloid Neoplasms Extended Mutation Panel [M MPN Extended]

*JAK2 (V617F and Exon 12), CALR (Type 1 and 2), MPL, ASXL1, CBL, CSF3R, ETV6/TEL, EZH2, IDH1, IDH2, KIT, KRAS, NRAS, RUNX1, SETB1, SF3B1, SRSF2, TET2, TP53*

AML Mutation Analysis Panel [M AML]

*ASXL1, DNMT3A, FLT3, IDH1, IDH2, KIT, NPM1, RUNX1, TET2, TP53, WT1*

*IgVH* Somatic Hypermutation [M IgVH]

B-Cell *IGH* Gene Rearrangement [M BCELL]

B-Cell *IGK* Gene Rearrangement [M IGK]

T-Cell Gene Rearrangement [M TCR]

BCR/ABL qRT PCR Major p210 & Minor p190 transcripts [M BCR ABL]

*JAK2 V617F* Mutation [M JAK2]

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## Fluorescence in situ Hybridization (FISH)

### Hematology Panels

Adult Acute Lymphoblastic Leukemia Panel [F AALL]

t(9;22), t(12;21), *MLL*(KMT2A)

Burkitt "Double Hit" Large B-cell Lymphoma Panel [F BURKITT]

*C-MYC*, *BCL2*, *BCL6*

Chronic Lymphocytic Leukemia Panel [F CLL]

*MYB* (6q23), *ATM* (11q22.3), +12, *DLEU1* (13q14.3), *TP53* (17p13)

Myelodysplasia Panel [F MDS]

-5/5q-, -7/7q-, +8, 20q-

Myeloproliferative Neoplasms Panel [F MPD]

+8, t(9;22), 13q-, 20q-

Plasma Cell Neoplasm Panel [F MM]

1p-, 1q+, +5, +9, t(11;14), 13q-, +15, 17p-, reflex t(4;14) / t(14;16)

### Hematology Probes

*AML1*(*RUNX1*)/*ETO*(*RUNX1*) t(8;21) [F AML ETO]

*API2*/*MALT1* t(11;18) [F API MALT]

*ATM*/*CEP11* deletion 11q22.3 [F ATM]

*BCL6* 3q27 rearrangement [F BCL6]

*BCR*/*ABL1* t(9;22) [F BCR ABL]

*CBFB* t(16;16), inv(16) [F CBFB]

*CEP 7 -7/7q31* [F CEP7]

*C-MYC* 8q24 rearrangement [F CMYC]

Deletion 7q22 [D7S796, D7S658]/7q31.2 [D7S486] [F 7q]

Deletion 13q14.3 [F D1314]

Deletion 20q12 [F D20]

*EGR1* 5q deletion, monosomy 5 [F EGR1]

*FIPIL1*/*PDGFRA* 4q12 gene rearrangement [F 4q12]

*IGH* 14q32 rearrangement [F IGH]

*IGH*/*BCL1* (*CCND1*) t(11;14) [F BCL1]

*IGH*/*BCL2* t(14;18) [F BCL2]

*IGH*/*FGFR3* t(4;14) [F FGFR3]

*IGH*/*MAF* t(14;16) [F IGH MAF]

*IGH*/*MALT1* t(14;18) [F IGH MALT]

*IGH*/*MYC* t(8;14) [F IGH MYC]

*MALT1* 18q21 rearrangement [F MALT]

*MLL* (KMT2A) 11q23 Gene Rearrangement [F MLL]

*MYB* 6q deletion [F MYB]

*PDGFRB* Gene Rearrangement [F PDGFRB]

*PML*/*RARA* t(15;17) [F PML RARA]

*TEL* (*ETV6*)/*AML1* (*RUNX1*) t(12;21) [F TEL AML1]

*TP53* 17p13 deletion [F P53]

Trisomy 5, 9, 15 [F T5T9T15]

Trisomy 8 [F T8]

## Women's Health

### STD and Infectious Disease

Chlamydia Trachomatis, Qualitative by Aptima COMBO® 2 TMA [CT]

Human Papillomavirus Genotyping [HPV G]

Human Papillomavirus High Risk by Aptima HPV [HPV HR]

Herpes Simplex Virus Type 1 and 2 Qualitative by PCR [HSV]

Neisseria Gonorrhoeae Qualitative by Aptima COMBO® 2 TMA [NG]

Trichomonas Vaginalis by Aptima [TV]

## Other

### Fluorescence in situ Hybridization (FISH)

Bladder Cancer Panel [F URO]

+3,+7,+17,9p21-

Molar Pregnancy Panel [FP ANEU]

+13,+18,+21, X, Y

### Cytogenetics

Chromosome Analysis Peripheral Blood (Constitutional) [CYTO PB]

Chromosome Analysis Products of Conception [CYTO POC]

