

# UCSF Medical Center

## Medical Genomics - Cytogenetics Laboratory

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### Patient Information:

Last Name \_\_\_\_\_ First Name \_\_\_\_\_ MI \_\_\_\_\_

Date of Birth \_\_\_\_\_

Male  Female  Unknown

Ordering Physician / Phone Number \_\_\_\_\_

Patient or Sample ID# \_\_\_\_\_

### Sample Information:

#### Specimen Collected:

Date: \_\_\_\_\_ Time: \_\_\_\_\_ Account #: \_\_\_\_\_

Tissue Biopsy / POC site: \_\_\_\_\_

Gestational Age: (weeks/days) \_\_\_\_\_

#### Sample Type

- |  |   |
|--|---|
| <input type="checkbox"/> Amniotic Fluid            | <input type="checkbox"/> Leukemic Blood |
| <input type="checkbox"/> Chorionic Villus Sampling | <input type="checkbox"/> Bone Core      |
| <input type="checkbox"/> Products of Conception    | <input type="checkbox"/> Solid Tumor    |
| <input type="checkbox"/> Constitutional Blood      | <input type="checkbox"/> Lymph Node     |
| <input type="checkbox"/> Bone Marrow               | <input type="checkbox"/> Other: _____   |

#### For Oncology Studies:

- Transplant:  Yes  No      Type: \_\_\_\_\_      Donor Gender: \_\_\_\_\_
- Staging:  Yes  No      Chemo:  Yes  No
- Site:  Iliac Crest       Left Ant       Left Post       Right Ant       Right Post
- Sternal

### Testing Information:

Indication for Study: (MUST be provided)  Pathology Reports  Patient's Previous Reports

Clinical Data/ Provisional Diagnosis: \_\_\_\_\_

ICD-10 Codes: \_\_\_\_\_

### Test Requested:

#### Cytogenetic Analysis:

- CYAFB Amniotic Fluid, chromosome and AF-AFP analysis
- CYAF Amniotic Fluid, chromosomes only
- CYCV Chorionic Villus Sampling, chromosome analysis
- CYTIS POC / Skin biopsy for routine chromosome analysis
- CYTLL Chromosome analysis; bone marrow, bone core or leukemic blood
- CYHR Peripheral Blood, high resolution chromosome analysis (>550 band level)
- CYSTU Solid Tumor analysis

If results are normal, REFLEX to:

- Cytogenetic Analysis
- Microarray Analysis
- Fluorescence in situ hybridization

#### Microarray Analysis:

- SNPAP SNP Array for Prenatal Analysis
- SNPAT SNP Array for Tissue and POC Analysis  
*\*TCACFCV / TCPOC test codes may be added for tissue culture*
- SNPAB SNP Array for Blood Analysis (Postnatal)
- HSNPA Hold SNP Array
- PSNPA SNP Array, Family Follow Up

#### Constitutional FISH tests:

- CYFDB/CYFD Aneuploidy (AV) Interphase FISH Aneuploidy analysis of Chromosomes 13, 18, 21, X, Y
  - A1321 Aneuploidy analysis of Chromosome 13 and 21 only
  - A18XY Aneuploidy analysis of Chromosome 18, X and Y only
- CYSUB/CYFST FISH Subtelomere analysis, Metaphase analysis of subtelomere regions
- CYFMB/CYFM FISH Microdeletion analysis, Metaphase analysis for regions below (MUST INDICATE SPECIFIC TEST)
  - \_\_\_ Wolf Hirshhorn (WHS, 4p16)      \_\_\_ Prader Willi/Angelman (SNRPN, 15q11-q13)      \_\_\_ DiGeorge/VCF/distal 22q (TUPLE1/ARSA, 22q11.2/22q13)
  - \_\_\_ Cri du Chat (CDCR, 5p15)      <sup>†</sup> Methylation test must be performed first by MDx      \_\_\_ Steroid sulfatase deficiency (STS, Xp22.3)
  - \_\_\_ Williams (ELN, 7q11.23)      \_\_\_ Smith Magenis (SMS, 17p11.2)      \_\_\_ Kallman Syndrome (KAL1, Xp22.3)
  - \_\_\_ Retinoblastoma (RB1, 13q14)      \_\_\_ Miller Dieker (LIS1, 17p13.3)      \_\_\_ SRY (Yp11,3)

#### FISH Oncology (Bill Only Test Codes):

##### CLL Panel (CYCLL)

- DEL11Q ATM
- TRIS12 Trisomy 12
- DEL13Q Deletion 13q
- DEL17P Deletion 17p

##### MDS Panel (CYMDS)

- M5D5Q Monosomy 5/Deletion 5q
- M7D7Q Monosomy 7/Deletion 7q
- TRIS8 Trisomy 8
- DEL20Q Deletion 20q

##### AML Panel (CYAML)

- M5D5Q Monosomy 5/Deletion 5q
- M7D7Q Monosomy 7/Deletion 7q
- TRIS8 Trisomy 8
- TR821 RUNX1/RUNX1T1
- MLLQ23 MLL,11q23
- TR1517 Translocation 15;17 (PML/RARA)
- INV16Q Inversion / Translocation 16
- DEL20Q Deletion 20q

##### Multiple Myeloma Panel (CYMM)

- DUP1Q Duplication 1q
- TR414 Translocation 4;14
- TR1114 Translocation 11;14
- CD138 CD138+ Enrichment
- DEL13Q Deletion 13q
- TR1416 Translocation 14;16
- DEL17P Deletion 17p

##### Lymphoma Panel

- BCL6 3q27 rearrangement
- TR814 Translocation 8;14 (IGH/MYC)
- MYC 8q24 rearrangement Break-apart
- TR1114 Translocation 11;14 (IGH/CCND1)
- TR1418 Translocation 14;18 (IGH/BCL2)

##### Eosinophilia Panel

- PDGFRA 4q12 Rearrangement
- PDGFRB 5q32q33 Break-apart
- FGFR1 8p11 Break-apart

##### Other Oncology FISH:

- TR119 Translocation 1;19
- ALK2P 2p23 rearrangement
- INV3Q RPN1/MECOM
- DEL9P Deletion 9p
- TR922 Translocation 9;22 (BCR/ABL)
- IGHQ32 14q32 IGH Break-apart
- XXXY TX monitoring, sex mismatch

Other (MUST SPECIFY AND PROVIDE SEND-OUT PAPERWORK): \_\_\_\_\_