DIAGNOSTIC CYTOGENETICS
2292 West Commodore Way Ste 100 Seattle, WA 98199
800-328-2026 fax: 206-325-2975 www.dxcyto.com email: dci@dxcyto.com

NEOPLASTIC SPECIMENS REQUEST FORM

Last Name:_____ First Name: _____

Sex:_____ Date of Birth:_____ PT#: _____

SSN: _____ Phone (Day): _____ Phone (Eve): _____

Address:_____

Charge to (please attach admissions face sheet, completed insurance forms, and/or a copy of insurance card. If not available, please complete the following):

Insurance Company:_____ Phone:_____

Address:_____

Policy Holder:_____ ID#:_____ Group#: _____

☐ Inpatient ☐ Outpatient Hospital Name_____ ☐ Nonpatient

SPECIMEN TYPE

DATE SAMPLE DRAWN:_____ TIME SAMPLE DRAWN: _____

☐ Bone Marrow Aspirate ☐ Bone Core Biopsy ☐ Peripheral Blood ☐ Lymph Node

☐ Other (please describe): _____

CLINICAL INFORMATION/ICD-10 (required): _____

☐ New Diagnosis ☐ Relapse ☐ Monitoring

☐ Radiation Therapy_____ ☐ Chemotherapy_____

☐ Bone Marrow Transplant ☐ Autologous ☐ Allogenic ☐ Sex Mismatch

TEST REQUESTED

☐ Chromosome Analysis ☐ Hold Culture for Testing

☐ Flow Cytometry** ☐ Molecular Analysis** _____

☐ FISH (Please mark below)* For paraffin embedded tissue, please see our FFPE request form.

☐ALL ☐AML ☐CLL ☐CML ☐MDS ☐MM ☐MPD ☐Eosinophilia ☐NHL ☐ Other:

☐ REFLEX FISH IF CHROMOSOME RESULTS ARE NORMAL (Please mark below)*

☐ALL ☐AML ☐CLL ☐CML ☐MDS ☐MM ☐MPD ☐Eosinophilia ☐NHL ☐Other:

☐ OTHER _____

*Please complete second page for other testing
**Sent out to reference laboratory

Physician:_____

Referring Hospital/Lab: _____

Address: _____

Phone:_____ Fax: _____

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ADDITIONAL INFORMATION FOR FISH (please see our website for our most current FISH offerings)

Patient Last Name:_____ First Name: _____

FISH

- **MDS**
  - ☐ **MDS Panel**: -5/del(5q), -7/del(7q), +8, del(20q)
  - ☐ Monosomy 5/deletion 5q (EGR1)
  - ☐ Monosomy 7/deletion 7q (D7S522/D7Z1)
  - ☐ Trisomy 8 (D8Z2)
  - ☐ Deletion 20q (D20S108)
  - ☐ Other _____
- **CLL**
  - ☐ **CLL Panel**: del(11q), del(17p), +12, -13/del(13q)
  - ☐ Deletion 11q (ATM)
  - ☐ Deletion 17p (TP53)
  - ☐ Trisomy 12 (D12Z3)
  - ☐ Monosomy 13/Deletion 13q (D13S319/LAMP1)
  - ☐ Translocation 11;14 (IGH/CCND1)
  - ☐ Other _____
- **MM/Plasma cell dyscrasia**
  - ☐ **MM Panel**: 1p/1q, t(4;14), -13/del(13q), del(17p)
  - ☐ Deletion 1p32.3/gain of 1q21 (CKS1B/CDKN2C)
  - ☐ Monosomy 13/Deletion 13q (D13S319/LAMP1)
  - ☐ Deletion 17p (TP53/CEP17)
  - ☐ 14q32 rearrangements (IGH breakapart)
  - ☐ Translocation 4;14 (IGH/FGFR3)
  - ☐ Translocation 11;14 (IGH/CCND1)
  - ☐ Translocation 14;16 (IGH/MAF)
  - ☐ Hyperdiploidy of chromosomes 5, 9, 15
  - ☐ Other _____
- **CML**
  - ☐ Translocation 9;22 (BCR/ABL1)
  - ☐ Deletion 9q34 (ASS)
  - ☐ Other _____
- **AML**
  - ☐ **AML Panel:** -7/del(7q), +8, t(8;21), 11q23 rearrangements, t(15;17), inv(16)/t(16;16)
  - ☐ Monosomy 7/deletion 7q (D7S522/D7Z1)
  - ☐ Trisomy 8 (D8Z2)
  - ☐ Translocation 8;21 (ETO/RUNX1)
  - ☐ 11q23 rearrangements (MLL)
  - ☐ Translocation 15;17 (PML/RARA)
  - ☐ Inversion 16/translocation 16 (CBFB)
  - ☐ Other _____
- **ALL**
  - ☐ **ALL Panel**: t(9;22), 11q23 rearrangements, t(12;21), del(9p),t(8;14)
  - ☐ Translocation 9;22 (BCR/ABL1)
  - ☐ 11q23 rearrangements (MLL)
  - ☐ Translocation 12;21 (ETV6/RUNX1)
  - ☐ Translocation 8;14 (IGH/MYC/D8Z2)
  - ☐ Deletion 9p21 (CDKN2A)
  - ☐ Other _____
- **LYMPHOMA**
  - ☐ **NHL Panel**: 3q27 rearrangements, 8q24 rearrangements, t(11;14), t(11;18), t(14;18)
  - ☐ 3q27 rearrangements (BCL6)
  - ☐ 8q24 rearrangements (MYC)
  - ☐ Translocation 11;14 (IGH/CCND1)
  - ☐ Translocation 11;18 (BIRC3/MALT1)
  - ☐ Translocation 14;18 (IGH/BCL2)
  - ☐ Other _____
- **MPD**
  - ☐ **MPD Panel**: 4q12 rearrangements, 5q33rearrangements,+8, t(9;22), del(13q), del(20q)
  - ☐ 4q12 rearrangements (PDGFRA-FIP1L1)
  - ☐ 5q33 rearrangements (PDGFRB)
  - ☐ Trisomy 8 (D8Z2)
  - ☐ Translocation 9;22 (BCR/ABL1)
  - ☐ Deletion 13q (D13S319/LAMP1)
  - ☐ Deletion 20q (D20S108)
  - ☐ Other _____
- **EOSINOPHILIA**
  - ☐ **Eosinophilia Panel:** 4q12 rearrangements, 5q33 rearrangements, 8p11.23-p11.22 rearrangements
  - ☐ 4q12 rearrangements (PDGFRA-FIP1L1)
  - ☐ 5q33 rearrangements (PDGFRB)
  - ☐ 8p11.23-p11.22 rearrangements (FGFR1)
  - ☐ Other_____
- **OTHER**
  - ☐ 2p23 rearrangements (ALK)
  - ☐ 3q26.2 rearrangements (EVI1)
  - ☐ 6q23 deletion (MYB)
  - ☐ Translocation 8;9 (PCM1/JAK2)
  - ☐ 12p13.1 rearrangements/deletions (ETV6)
  - ☐ 13q14 rearrangements (FOXO1)
  - ☐ 14q11.2 rearrangements (TCR alpha/delta)
  - ☐ 17q21 rearrangements (RARA)
  - ☐ 22q12 rearrangements (EWSR1)