



Confidential

Cytogenetics Request Form

Patient Information:

Name (4 parts): _____ Male Female Unknown/Date of Birth: ___/___/___

Ethnic Origin /City: _____ Country _____ Address/City: _____ Country: _____

1st Phone: _____ 2nd Phone: _____

Hospital No.: _____ NCDEG File No.: _____ Geneticist: _____

Specimen Type:

Date Specimen Drawn: ___/___/_____

Drawn by: _____

Date Specimen Received: _____

Pregnancy: No Yes Gestation _____ weeks

Last Menstrual Period: _____

Sample Appearance: _____

Specimen Type: Blood Amniotic Fluid

CVS Fetal/Cord Blood

Bone Marrow

Bone Core Biopsy

Blood-hematological disorder

WBC: _____ % Blasts: _____

Pretreatment Pre BMT Remission

Relapse/Blast Crisis

Post BMT-donor sex Male Female Auto

Treatment w/cytotoxic agents No Yes

Tumor

Other: _____

Indication for Study:

To avoid delay in sample processing, provide an accurate list of indications (must check one or more items below):

Ambiguous Genitalia

Autism Spectrum

Autosomal Trisomy

Dysmorphic Features

Fetal Demise

Developmental Delay

Mild Moderate Severe

Infertility

Klinefelter/Turner

Multiple Congenital Anomalies

Multiple Miscarriages

Seizure Disorder

Failure to Thrive

Still Birth

Other: _____

Previous Genetic Reports: No Yes: _____

Laboratory Tests Ordered:

Karyotype

Special Culture for Fragile-X Syndrome

Breakage Study (Control Requested)

Chromosomal Microarray

FISH for Philadelphia chromosome

FISH for X, Y, 13, 18, 21 chromosomes

FISH for X, Y, 21 chromosomes

FISH for SRY Region

FISH for Microdeletion Syndromes:

Miller-Dicker/Lissencephaly Region

Prader-Willi/Angelman Region

William's Syndrome

FISH for 21 chromosome

FISH for 13, 18, 21 chromosomes

Smith Magenis Syndrome

Digeorge Syndrome

Other: _____

Referring M.D.: _____ Hospital Name: _____



Pedigree Information

History Taken by: _____ Date: ____/____/____

Patients Pregnancy History/Gravida: _____ Para: _____ Miscarriages: _____

Maternal Age: _____ Paternal Age: _____

