

Molecular Genetics Request Form

Patient Information:

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

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

1st Phone: _____ 2nd Phone: _____

Date: _____ NCDEG File No.: _____ Geneticist: _____

Clinical Information:

Pregnancy: No Yes Gestation _____ weeks

Last Menstrual Period: _____

Specimen Type: Blood Presymptomatic Test

CVS* Amniotic Fluid*

DNA Saliva

*Back up culture by: inside Lab outside Lab

Date Specimen Drawn: ____/____/____

Drawn by: _____

Date Specimen Received: _____

Sample Appearance: _____

Indications for Test:

Diagnostic: Known Affected

Suspected: Symptoms _____

Carrier: Family History (Attach Page 2)

No Family History

Abnormal Fetal Ultrasound

Known Carrier

Fetal: Family History (Attach Page 2)

Abnormal Fetal Ultrasound

Other: _____

Laboratory Tests Ordered:

DNA Extraction and Storage or/and Shipping

β -Thalassemia mutation Screening

Familial Mediterranean Fever mutation Screening

Hemochromatosis mutation C282Y and H63D analysis

Factor V mutation R506Q analysis

Factor II mutation G20210A analysis

Methylenetetrahydrofolate Reductase mutation C667T analysis

Y-Chromosome Microdeletion analysis

Sex Determining Region testing (SRY)

Fragile-X Syndrome

Cystic Fibrosis mutation Screening

Duchenne Muscular Dystrophy deletion analysis

Spinal Muscular Atrophy deletion test

Sanjad Sakati Syndrome (12pb del.)

Achondroplasia

Alpha-1-Antitrypsin

Aicardi Goutieres Syndrome (A177T)

Glutaricacidemia Type 1 (R402W)

Tyrosinemia type I (Q64H)

Other: _____

Referring M.D.: _____ Hospital Name: _____

