

DNA LABORATORY TEST REQUEST FORM *(Please send this completed form with the specimen)*

PATIENT LAST NAME: _____ FIRST NAME : _____ DOB: _____

SEX: M / F REF. LAB #: _____ DATE COLLECTED: _____ DATE RECEIVED: _____

ETHNIC BACKGROUND: (Circle all that apply) European Caucasian, Hispanic, Native American, African American, Asian, Other

REFERRING PHYSICIAN/HEALTH CARE PROFESSIONAL:
 Name: _____
 Address: _____
 City, State, Zip: _____
 Telephone:(_____) _____ FAX:(_____) _____

REFERRING INSTITUTION / CLINIC / LABORATORY:
 Name: _____
 ADDITIONAL REPORTS TO:
 Name: _____

PHYSICIAN SIGNATURE: _____ Date: _____
 (Required for Medicare / Medicaid billing)

BILLING INFORMATION:
 REFERRING INSTITUTION

 New Clients please call Laboratory with financial contact information.

INSURANCE
 Name of policy holder: _____
 Policy holder DOB: _____
 SS # (Guarantor): _____
 Address: _____
 Phone #: _____
 Relationship to patient: _____
 Insurance Co. / Policy #: _____
 Insurance Co Contact / Phone : _____

MEDICAID #: _____
State (MT, WY, CO, UT): _____
SS#: _____
 SELF PAY
 INPATIENT

TEST REQUESTED: (Please check one or more; test descriptions with CPT codes and prices faxed on request)

- | | |
|---|--|
| <input type="checkbox"/> Hemochromatosis (HFE) Mutations (C282Y / H63D) | <input type="checkbox"/> Uniparental Disomy Screen (Chromosome _____)
(chromosomes 2, 6, 7, 8, 9, 11, 13, 14, 15, 16, 20, 21)
* Note: Parents required for UPD studies |
| <input type="checkbox"/> Factor V Leiden (FVL) Mutation | <input type="checkbox"/> Angelman Syndrome (AS) Methylation |
| <input type="checkbox"/> Prothrombin (Factor II) Polymorphism | <input type="checkbox"/> Prader- Willi Syndrome (PWS) Methylation |
| <input type="checkbox"/> Factor V Leiden / Prothrombin / MTHFR Variant | <input type="checkbox"/> Fragile X Syndrome (FMR-1): DNA only |
| <input type="checkbox"/> Huntington Disease (HD) Mutation | Beckwith-Wiedemann Syndrome |
| <input type="checkbox"/> X-Chromosome Inactivation | <input type="checkbox"/> KCNQ1OT1 & H19 Methylation <input type="checkbox"/> UPD 11 |
| <input type="checkbox"/> Zygoty (Twin) Studies | Russell-Silver Syndrome |
| <input type="checkbox"/> Maternal Cell Contamination Analysis | <input type="checkbox"/> H19 Methylation <input type="checkbox"/> UPD 7 |
| <input type="checkbox"/> Microarray | |
| <input type="checkbox"/> Other (Specify: _____) | |

CLINICAL INFORMATION / INDICATION FOR TEST:

Reason for Testing: Diagnostic Carrier Family History Testing Prenatal
 ICD-9 Code: _____ Clinical Findings / History : _____

Pregnancy history (gravida/para/ab): _____ Gestational age & method (US, LMP): _____

ADDITIONAL FAMILY MEMBERS TO BE STUDIED AND RELATIONSHIP (provide parents names & dates of birth for UPD studies)

1. _____ 2. _____

SPECIMEN TYPE: (Please check) PLEASE NOTIFY THE LABORATORY WHEN A SPECIMEN IS BEING SENT.

- | | |
|--|---|
| <input type="checkbox"/> Cultured amniocytes (Two confluent T25 flasks, please retain a backup) | <input type="checkbox"/> Blood: EDTA (purple top tube) or ACD (yellow) 5 to 10 ml, minimum 1 ml for infants, room temp |
| <input type="checkbox"/> Cultured chorionic villus (CVS) cells (Two confluent T25 flasks, please retain a backup) | For chromosome analysis, also submit 3 to 5 ml sodium heparin (green top) |
| <input type="checkbox"/> Direct amniotic fluid or CVS (Please call the laboratory to arrange) | <input type="checkbox"/> Tissue, Fixed or Fresh, source: (block or thick sections, please contact lab) |
| | <input type="checkbox"/> OTHER: |

Please submit 3-5 ml maternal blood collected in an EDTA tube for maternal cell contamination study. We strongly recommend this test when chorionic villus samples results indicate a female fetus (we discard the sample if the results are male). If you do not choose to submit this sample, please check the box below which assures that you have discussed this testing with the patient and she declines this additional testing.

DECLINE

Med Rec # _____ Admit # _____ Shire # _____