



Genetics Diagnostic Laboratory

Molecular Genetics Test Requisition

**DELIVER TO:**

CHEO Genetics Diagnostic Laboratory
Room w3401
401 Smyth Road Ottawa, ON
Canada K1H 8L1
Phone: (613) 738-3230 Fax: (613) 738-4814
www.cheo.on.ca/en/genetics

PATIENT NAME _____ (LAST) _____ (FIRST)

ADDRESS _____

CITY _____ PROV. _____ POSTAL CODE _____

 DATE OF BIRTH _____ SEX: M F
DD/MM/YYYYY

FACILITY PATIENT ID NUMBER _____

PROVINCIAL HEALTH NUMBER _____

FOR LAB USE ONLY:

COLLECTED DATE: _____

COLLECTION CENTRE: _____

SPECIMEN COLLECTED BY: _____

HEALTH CARE PROVIDER(S) REQUESTING TEST:

NAME _____

REGISTRATION NUMBER _____

ADDRESS _____

CITY _____ PROV. _____ POSTAL CODE _____

CONTACT: _____

PHONE NO. _____ FAX NO. _____

AUTHORIZED SIGNATURE:

IF AN ADDITIONAL REPORT IS BEING REQUESTED, PLEASE COMPLETE THE FOLLOWING:

PROVIDER NAME _____ FAX NO. _____

ADDRESS _____ CITY _____ PROV. _____ POSTAL CODE _____

Sample Required

- | | |
|--|--|
| <input type="checkbox"/> Blood 2x5 mL EDTA | <input type="checkbox"/> Cultured Amniocytes |
| <input type="checkbox"/> Blood 3 mL EDTA (infant only) | <input type="checkbox"/> Amniotic Fluid 5 mL |
| <input type="checkbox"/> DNA _____ug | <input type="checkbox"/> Cultured CVS |
| <input type="checkbox"/> Other _____ | <input type="checkbox"/> CVS |

Sample Information

-
- Routine**
-
-
- Expedited**
-
-
- Patient/Partner Pregnant
-
-
- Prenatal Diagnosis
-
-
- Newborn (less than 3 months of age)

Reason for Test

-
- Symptoms of Indicated Disease
-
-
- Carrier Testing
-
-
- Predictive Testing
-
-
- Prenatal Diagnosis (
- parental samples are required*
-)
-
-
- Other: _____

Other family member(s) tested previously:

-
- No
-
-
- Yes Name: _____
-
- CHEO Pedigree number: _____
-
- Relationship to patient: _____

Test Requested

- | | |
|---|--|
| <input type="checkbox"/> Angelman Syndrome
<input type="checkbox"/> Facioscapulohumeral Muscular Dystrophy (<i>20 mL in EDTA required</i>)
<input type="checkbox"/> Factor V Leiden and Factor II Prothrombin
<input type="checkbox"/> Fragile X (FMR1)
<input type="checkbox"/> HFE-related Hemochromatosis
<input type="checkbox"/> Maternal Cell Contamination Studies
<input type="checkbox"/> Myotonic Dystrophy Type I (<i>ethnic background must be specified below</i>)*
<input type="checkbox"/> Myotonic Dystrophy Type II | <input type="checkbox"/> Oculopharyngeal Muscular Dystrophy
<input type="checkbox"/> Prader-Willi Syndrome
<input type="checkbox"/> Spinal Muscular Atrophy

<input type="checkbox"/> DNA Storage (<i>DNA will be stored for 2 years and then discarded; contact the laboratory directly if a longer storage term is required</i>) |
|---|--|

 * **Ethnic Background** (e.g. Ashkenazi Jewish, Asian, French Canadian, Northern European): _____
Additional relevant clinical and/or family history information: