

# REQUISITION – Hemoglobinopathy Genetic Testing



**Hamilton Regional  
Laboratory Medicine  
Program**

## Juravinski Hospital

Clinical Genetics Laboratory - Room H2-19A  
711 Concession Street, Hamilton, ON L8V 1C3  
Phone: (905) 521-2100 x76944 | Fax: (905) 521-7913  
Email: [moleculargenetics@hhsc.ca](mailto:moleculargenetics@hhsc.ca)

## Patient Information

\*Name (print):

*Surname, First Name*

\*DOB (DD/MM/YYYY):

\*Sex:  M  F  Other

\*Health Card No.:

*\*Mandatory Information. Specimen cannot be processed without this data.*

**Note: Specimen collection is NOT completed at this lab. Please proceed to any community lab for blood draw.**

## Reports To:

\*Ordering Physician: \_\_\_\_\_

Clinic/Hospital: \_\_\_\_\_

\*Phone: \_\_\_\_\_ Fax: \_\_\_\_\_

Email: \_\_\_\_\_

Physician Signature: \_\_\_\_\_

## Additional Copies To:

\*Name: \_\_\_\_\_

\*Clinic/Hospital: \_\_\_\_\_

\*Phone: \_\_\_\_\_

\*Fax: \_\_\_\_\_

\*Email: \_\_\_\_\_

Please see the HRLMP Laboratory Test Information Guide (LTIG) for complete sample requirements and test information:

<https://ltig.hrlmp.ca/>

## SPECIMEN INFORMATION

*Ship at room temperature. Refrigerate at 4°C if overnight or longer storage is unavoidable. Avoid freezing and exposure to excess heat.*

Collection Date (DD/MM/YYYY): \_\_\_\_\_ Time of collection: \_\_\_\_\_

- Peripheral blood** in EDTA (4ml >1 yr/age, 0.5ml <1 yr/age)
- DNA** (minimum 1 µg). Source: \_\_\_\_\_
- Cord blood** in EDTA (1-4ml)
- Amniotic Fluid** (10-15ml, back up culture required)
- Cleaned Chorionic Villi** (5-15mg, back up culture required)
- Cultured Cells** (1xT25 confluent flask, back up culture required)

## TEST REQUESTED

**Please note: CBC, Hemoglobin electrophoresis, and ferritin results are required for processing all hemoglobinopathy samples. Failure to submit these results may result in testing delays.**

**Hemoglobinopathy:**

Ethnicity: \_\_\_\_\_

Thalassemia

Hemoglobin Variant

Sickle Cell Disease

## CLINICAL INDICATION

- Symptoms of indicated disease**
- Carrier status**
- Newborn Screen Positive**
- Prenatal diagnosis** (please complete information below)  
LMP (DD/MM/YY): \_\_\_\_\_  
Procedure Date (DD/MM/YY): \_\_\_\_\_
- Family history** (please complete information below)
  - Patient is proband/index case
  - Known familial mutation (or HRLMP report #): \_\_\_\_\_Proband name: \_\_\_\_\_ DOB (DD/MM/YY): \_\_\_\_\_ Relationship to patient: \_\_\_\_\_
- Other** (please provide additional details): \_\_\_\_\_

### Urgent/Expedited Cases:

- Prenatal Diagnosis
- Newborn Screen Positive
- Patient Pregnant
- Partner Pregnant (add information below)  
Partner Name: \_\_\_\_\_  
Partner DOB (DD/MM/YY): \_\_\_\_\_