



Phone: 905-433-2733 Fax: 905-721-6122

Phone: 905-433-2733 Fax: 905-721-6122

****Incomplete or illegible forms will be returned****

****Please include all relevant medical records to allow efficient booking****

PATIENT DEMOGRAPHICS

Name: _____ ☐ M ☐ F

DOB (DDMMYYYY): _____

OHIP# _____ VC: _____

Address: _____

Home Phone: _____

Alternate Phone: _____

For minor patients, please provide name of parent(s)/legal guardian(s): _____

☐ **PRENATAL REFERRAL (MUST INCLUDE DATING ULTRASOUND)**

(Please attach antenatal records, ultrasound reports, prenatal lab results, screening reports)

LMP: DDMMYYYY:

- ☐ Late maternal age/Prenatal screening
- ☐ Positive prenatal screen
- ☐ Fetal ultrasound anomalies
- ☐ Family history of genetic condition or birth defect

☐ **GENERAL REFERRAL**

(Please provide details on the right and attach all relevant records/consult notes)

Is this patient clinically affected? ☐ No ☐ Yes

- ☐ Pediatric assessment
- ☐ Assessment for adult onset disorders
- ☐ Genetic counselling: family history of genetic conditions or birth defects

☐ **HEREDITARY CANCER REFERRAL**

(see page 2 for referral criteria)

Does the patient have a personal history of cancer? ☐No ☐Yes (please attach pathology)

Type	Count	Percentage
1	10	10.0%
2	20	20.0%
3	30	30.0%
4	40	40.0%
5	50	50.0%
6	60	60.0%
7	70	70.0%
8	80	80.0%
9	90	90.0%
10	100	100.0%

Age at Diagnosis: _____

Has a mutation been identified in the family?

- ☐
- No
- ☐
- Yes

Which gene?

REFERRING PHYSICIAN

Name: _____

Address: _____

Telephone #:

Fax#: _____

Physician Billing #:

Signature: _____

If this referral is URGENT, please specify why:

Will an interpreter be required: ☐ YES ☐ NO

If yes, please specify language: _____

Please provide additional details/relevant family history regarding this referral:

[illegible]

LAKERIDGE HEALTH CLINICAL GENETICS: HEREDITARY CANCER REFERRAL CRITERIA

***** Please check all that apply. Cancers must be present on the
same side of the family *****

HEREDITARY BREAST AND OVARIAN CANCER (HBOC):

- ☐ HBOC associated cancers in family (especially if diagnosed under age 50):
 - ≥3 relatives diagnosed with breast, ovarian, pancreatic, prostate cancer or melanoma OR
 - ≥2 relatives with breast cancer OR
 - Personal or family history of serous ovarian cancer
- ☐ Breast cancer diagnosed under 35 years of age
- ☐ Triple negative breast cancer diagnosed under 60 years of age
- ☐ Bilateral breast cancer
- ☐ Breast/Ovarian cancer in Ashkenazi Jewish women
- ☐ Male breast cancer
- ☐ Known mutation in a breast/ovarian cancer related gene (e.g. BRCA1/2, CHEK2) in the family
- ☐ Previously tested negative for BRCA1/2 – for consideration of additional panel/gene testing
- ☐ Assessment for eligibility for OBSP High-Risk Breast Screening

HEREDITARY COLORECTAL CANCER (Lynch Syndrome, Polyposis):

- ☐ Individual with colorectal or uterine cancer diagnosed under 70 years old
- ☐ Individual with 10 or more colorectal polyps
- ☐ Multiple cases of colorectal cancer in the family
- ☐ Family history suggestive of Lynch Syndrome ≥3 relatives with colorectal, endometrial, pancreatic, ovarian, kidney, urinary tract, small bowel, gastric and brain cancers
- ☐ Family history suggestive of hereditary polyposis
- ☐ Known hereditary colorectal cancer (e.g. MLH1, MSH2, MSH6, PMS2, EPCAM) or polyposis (e.g. APC, MutYH) gene mutation in family

OTHER HEREDITARY CANCERS (Exclusions: lung and cervical cancer):

- ☐ A family history of a known hereditary cancer syndrome. Please provide details on page 1
- ☐ Individual diagnosed with more than one primary cancer
- ☐ Prostate cancer: metastatic prostate cancer or prostate cancer ≤ age 55 with Gleason score >7
- ☐ Multiple family members with the same cancer or rare cancers, especially if diagnosed under age 50

*Please note: We will triage the referral and notify your office of the decision regarding eligibility. We suggest informing your patient that their family history will be evaluated to determine if there is a need for an appointment. Genetic testing is offered only to families that are suggestive of a hereditary cancer syndrome, and in most cases will be offered to a family member affected with cancer first **If you are uncertain whether an individual/family history will meet criteria, please refer.***

