



REFERRAL FOR HEREDITARY GI CANCER CLINIC

Fax to: 416-586-5924

Referral Date: _____

Referring Physician:
Name: _____ Billing Number: _____
Phone: _____ Fax: _____
Institution/Address: _____

URGENCY:
 Yes (specify reason) _____
 No

INTERPRETER NEEDED:
 Yes (specify language) _____
 No

Blood drawn for Sinai Health Genetic Lab
 Yes – mainstream testing ordered
 Yes – banked
 No

PATIENT INFORMATION:
Name: _____ Male Female
Health card #: _____ Version code _____ DOB: _____ (DD/MMM/YEAR)
Phone #: _____ Email: _____
Address: _____

Reason for Referral (see next page for criteria):
 Path included (Please send us all scopes, cancer or polyp, IHC, methylation, BRAF, etc)

Referral reason	Criteria for testing
<input type="checkbox"/> Adenomatous Polyposis	<ul style="list-style-type: none"> • ≥ 20 at any age • ≥ 10 under age 60 (path required) • ≥ 5 with personal or family history (assess by GC) or young w other FAP features
<input type="checkbox"/> FAP features	<ul style="list-style-type: none"> • Desmoid under age 40 • Cribriform moluar variant papillary thyroid • Hepatoblastoma • Retinal pigment epithelial harmartoma
<input type="checkbox"/> Serrated Polyposis	<ul style="list-style-type: none"> • > 20 serrated polyps (at least 5 proximal to rectum) • > 5 serrated polyps (all ≥ 5 mm, at least 2 ≥ 10 mm, all proximal to rectum)
<input type="checkbox"/> Fundic Gland Polyposis	<ul style="list-style-type: none"> • > 100 FGP (including carpeting) • > 30 FGP and FDR with fundic gland polyposis or gastric cancer • Clustering in absence of PPI use and sparing antrum and lesser curvature
<input type="checkbox"/> Hamartomatous Polyps	<ul style="list-style-type: none"> • ≥ 5 Juvenile Polyps • > 2 Peutz-Jeghers hamartoma • Mixed (suggestion of <i>PTEN</i> hamartoma)
<input type="checkbox"/> Gastric or GEJ Adenocarcinoma	<ul style="list-style-type: none"> • Diagnosed w gastric or GEJ adenocarcinoma ≤ 50 • Diffuse gastric cancer + personal or family history of lobular breast cancer, 1 < 70 • Diffuse gastric cancer + cleft lip/palate or of Maori descent • Bilateral lobular breast, one under age 70 • Multiple gastric cancers in family or gastric + lobular in family
<input type="checkbox"/> Colorectal cancer or Endometrial cancer or Other LS cancer (ureter, small bowel, non-serous ovarian, renal pelvis, etc)	<ul style="list-style-type: none"> • Diagnosed < 50 – requiring MMR IHC • Two primary LS cancer, one < 60 – requires IHC • Personal hx of LS cancer + family hx – requires IHC • Diagnosed < 35 yrs old (regardless of IHC) • Amsterdam I/II family history (regardless of IHC) • CRC + polyposis (regardless of IHC) • MMR IHC deficient <ul style="list-style-type: none"> * MSH2 or MSH6 or PMS2 (only) deficiency * MLH1 +/- PMS2 deficient – Negative for <i>BRAF</i> and <i>MLH1</i> methylation * MLH1 +/- PMS2 deficient – Positive for <i>BRAF</i>/methylation <u>but</u> still suspicious (i.e. multiple primaries, young age of onset, strong fam hx) • Unaffected but strong family history
<input type="checkbox"/> Known syndrome in family	Lynch syndrome (confirmed germline MMR mutation), FAP, MAP, Peutz-Jeghers syndrome, Juvenile Polyposis Syndrome, Hereditary Gastric Cancer, GAPPs, <i>GREM1</i> , <i>POLE</i> , <i>POLD1</i> , <i>NTHL1</i> , <i>MSH3</i> , <i>MLH3</i>
<input type="checkbox"/> Ashkenazi Jewish	<ul style="list-style-type: none"> • Personal history of CRC or multiple polyps • Close relative with CRC < 60, breast, ovarian, pancreas or multiple colon polyps