

Grand River Regional Cancer Centre Cancer Genetics Referral Form

- Please fax this form to 519-749-4382. If you have questions or concerns, please call 519-749-4370, ext. 2832.
- A family history questionnaire must be completed and submitted with this completed referral form. The family history questionnaire is available online at:

https://www.grhosp.on.ca/professionals/chief-of-staffs-office/referral-forms

- Detailed referrals may be processed quicker than less detailed referrals.
- The completed referral and family history will be reviewed by the genetics clinic to determine your patient's eligibility for a genetic counselling appointment.
- Genetic testing may or may not be offered in the course of a genetics consultation, pending eligibility.

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		erral Eligibility						
	Please complet	te the checklist on	page 2.					
Is this assessment urgent?	Reason and timeframe:	eason and timeframe:						
□ No □ Yes								
Does the patient have a	Please include type (i.e. primary site(s), metastases), age at diagnosis, any concerning							
personal history of cancer?	pathologic features (please send pathology report):							
□ No □ Yes								
Does the patient have a	Please include type (i.e. pr	imary site(s), meta	stases), age at diagnosis, living or deceased:					
family history of cancer?								
□ No □ Yes								
		ent Information						
Name (Last, First, Middle Initial):		DOB (yyyy-mm	DOB (yyyy-mm-dd):					
Sex: ☐ Male ☐ Female		Address:						
Phone number:		Does the natie	nt require an interpreter?					
Thore number.		•	·					
Email address:		Patient consents to receive information by email:						
		□ No □ Yes						
Health card number:		Hospital chart number (if applicable):						
	Physi	cian Information						
Referring physician:	Phone number:		Fax number:					
Primary care provider:	Phone number:		Fax number:					
	Information Accor	mpanying Referral	(Required)					
☐ Family Health Questionnai☐ Other:	re (mandatory) 🔲 Referra	al letter 🗌 Patho	logy Family member's genetic test result					



Cancer Genetic Testing Referral Criteria

Referrals *must* meet one or more of the following criteria. Please check all of the boxes that apply.

Personal or family history of MULTPLE CANCERS among close relatives on the SAME SIDE of the family:
☐ 2 or more cancers from one of the following groups:
☐ breast/ovary/prostate/pancreas
☐ colorectal/uterine/ovarian/gastric/GE junction/ureter-renal pelvis/upper GI/pancreas-biliary tree/primary brain/sebaceous
neoplasm
☐ gastric/breast
☐ melanoma/pancreas/primary brain
☐ renal/hemangioma/cutaneous lesions (not including basal cell carcinoma)
\square endocrine tumours
☐ Multiple primary cancers in one person
☐ 3 or more close relatives with the same type of cancer
□ >5 basal cell carcinoma in one individual
☐ Brain tumour <i>plus</i> any other personal history of cancer, or family history of breast, colon, or sarcoma
☐ History suggestive of colorectal polyposis (≥10 adenoma, ≥5 adenoma plus family history of colon/uterine/brain, multiple fundic gland
polyposis, ≥5 serrated polyps, any hamartomatous/other rare polyps)
Personal or family history of a close relative with YOUNG cancer:
☐ Breast, renal, or sarcoma cancer ≤45 years of age
☐ Colon, uterine cancer, gastric or gastroesophageal junction cancer ≤50 years of age
☐ Basal cell carcinoma ≤30 years of age
Personal or family history of a close relative with RARE or STRONG HEREDITARY RISK cancer:
☐ Triple negative breast cancer ≤60 years of age
☐ Male breast cancer
☐ Invasive epithelial ovarian cancer, fallopian tube or peritoneal cancer, serous tubal intraepithelial lesions
☐ Ashkenazi Jewish ancestry with a personal/family history of breast, ovarian, colon cancer or GI polyposis
☐ Pancreatic adenocarcinoma
☐ Prostate cancer (≥T3 staging, Grade Group 4 or 5/Gleason Score 8-10), lymph node involvement, PSA ≥20)
☐ Uveal melanoma
☐ Bilateral or multifocal renal cell carcinoma, or non-clear cell pathology
☐ Diffuse gastric cancer
□ Other rare tumours with suspected hereditary link such as: gastrointestinal stromal tumour, pheochromocytoma, paraganglioma, cribiform-morular variant of papillary thyroid cancer, hepatoblastoma, desmoid <40yrs, hamartomas of retinal pigment epithelium, medullary thyroid cancer, adrenocortical carcinoma, cutaneous leiomyoma. For other rare tumours, please call the genetics clinic prior to referral.
Specific gene mutations (please send supporting documents):
☐ Family member with a known inherited hereditary cancer gene mutation (e.g. BRAC1/2, MLH1/MSH2/MSH6, APC), please specify:
☐ Mismatch repair deficient/MSI high tumours (MLH1 deficient tumours require BRAF and promotor methylation studies complete)
☐ Reproductive partner of this patient is a carrier of a cancer gene that is associated with an autosomal recessive condition (i.e. ataxia telangiectasia, Fanconi anemia, constitutional mismatch repair deficiency, and certain polyposis syndromes). Please provide name of reproductive partner, if tested at GRH, or copy of molecular result if tested elsewhere:

Include all supporting documentation (e.g. pathology reports, including tumour, polyp and breast biopsies; genetic test results) with the referral



Please complete to the best of your knowledge

Genetics Assessment Questionnaire

Please answer the following questions: 1) What are your main concerns/questions that you would like answered at your genetics appointment? 2) Has anyone in your family ever had genetic counselling? Yes No If Yes; What is their relationship to you? When were they seen? Where were they seen? Yes____No___ Don't Know ____ Was genetic testing offered? Yes_____No____Don't Know____ Was genetic testing completed? Was a genetic mutation found? Yes No Don't Know 3) What is your ethnicity or family country of origin? Mother's side Father's side: Ashkenazi Jewish? No Yes Mother Father



Please complete this to the best of your knowledge

Personal History

YOUR NAME	M/F	Date of Birth (M/D/Y)	If deceased, list year or age of death.	Cancer Type (e.g. breast, or none if never had)	Age at Diagnosis
Mother's initials + details					
Father's initials + details					
Your children- initials 1.					
2.					
3.					
4.					
5.					
6.					
7.					

Family History

YOUR BROTHERS & SISTERS - initials	M/F	Date of Birth (M/D/Y)	If deceased, list	Cancer Type	Age at	# Children	
YOUR BROTHERS & SISTERS - initials Total number of brothers Total number of sisters			year or age of death.	(e.g. breast, or none if never had)	Diagnosis	M	F
1.							
2.							
3.							
4.							
5.							
6.							
7.							



FAMILY HISTORY

GRANDPARENTS	Approximate	If deceased, list	Cancer Type	Age at
	Year of Birth	year or age of death.	(e.g. breast, or none if never had)	Diagnosis
YOUR GRANDMOTHER's initials (on your mother's side)				
YOUR GRANDFATHER's initials (on your mother's side)				
YOUR GRANDMOTHER'S initials (on your fathers side)				
YOUR GRANDFATHER's initials (on your father's side)				

YOUR MOTHER'S BROTHERS & SISTERS-	M/F	Approximate Year of	If deceased, list	Cancer Type	Age at	# Childr	en
initials		Birth	year or age of death.	(e.g. breast, or none if never had)	Diagnosis	M	F
Total number of Aunts on mother's side Total number of Uncles on mother's side							
1.							
2.							
3.							
4.							
5.							
6.							
7.							



FAMILY HISTORY

YOUR FATHER'S BROTHERS & SISTERS- initials	M/F	Approximate Year	If deceased,	Cancer Type	Age at	# Chil	dren
Total number of Aunts on your father's side Total number of Uncles on your father's side		of Birth	list year or age of death.	(e.g. breast, or none if never had)	Diagnosis	M	F
1.							
2.							
3.							
4.							
5.							
6.							
7.							

Name (Please Print) Last, First, (Maiden in Brackets) Initials	M/F	Approximate Year of Birth	Relationship to you and name of parent (eg. cousin, John Doe's daughter)	If deceased, list year or age of death.	Cancer Type (e.g. breast)	Age at Diagnosis
1.			adugnier)	age of aeain.		
2.						
3.						
4.						
5.						
6.						