



FAMILY NAME

MRN

GIVEN NAME

MALE FEMALE

D.O.B. ____ / ____ / ____

M.O.

Facility:

ADDRESS

**HEREDITARY CANCER CLINIC
REFERRAL**

LOCATION / WARD

COMPLETE ALL DETAILS OR AFFIX PATIENT LABEL HERE

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Prince of Wales & St George Hospitals

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Affected with _____
 Unaffected

URGENT
 Palliative

Interpreter Required
Language _____

Patient to be seen at:

Prince of Wales Hospital St George Hospital The Wollongong Hospital Shoalhaven Hospital

Purpose of Referral Assessment of family history (describe below)

Gastrointestinal / Gynaecological

	Personal Hx	*FHx
Cancer with Abnormal Immunohistochemistry (IHC) Mismatch Repair (MMR) genes	<input type="checkbox"/>	<input type="checkbox"/>
Gastrointestinal Cancer <50yrs	<input type="checkbox"/>	<input type="checkbox"/>
Gastrointestinal Cancer with *FHx	<input type="checkbox"/>	<input type="checkbox"/>
Diffuse Gastric Cancer with *FHx OR with Invasive Lobular Breast Cancer	<input type="checkbox"/>	<input type="checkbox"/>
Lynch Syndrome/ HNPCC	<input type="checkbox"/>	<input type="checkbox"/>
Familial Adenomatous Polyposis (FAP)/ Polyposis _____	<input type="checkbox"/>	<input type="checkbox"/>
Bowel Cancer AND Endometrial Cancer	<input type="checkbox"/>	<input type="checkbox"/>
Endometrial Cancer <60yrs	<input type="checkbox"/>	<input type="checkbox"/>
Ovarian Cancer <70yrs	<input type="checkbox"/>	<input type="checkbox"/>
Rare Gynaecological Cancer	<input type="checkbox"/>	<input type="checkbox"/>

***Family Hx (multiple cases with 1 case <60yrs OR ≥ 1 case with high risk feature)**

Rare / Other

	Personal Hx	*FHx
Medullary Thyroid Cancer, any age	<input type="checkbox"/>	<input type="checkbox"/>
Isolated Retinal Haemangioma <40yrs	<input type="checkbox"/>	<input type="checkbox"/>
Isolated CNS Haemangioblastoma <40yrs	<input type="checkbox"/>	<input type="checkbox"/>
NET <40yrs , or multiple NETs any site, or NET plus a 2nd tumour	<input type="checkbox"/>	<input type="checkbox"/>
Renal Cancer <45yrs or bilateral	<input type="checkbox"/>	<input type="checkbox"/>
Phaeochromocytoma/Paraganglioma <50yrs , or multiple any age	<input type="checkbox"/>	<input type="checkbox"/>
Multiple Primary Cancers ≥ 3 -Discuss with Hereditary Cancer Clinic (HCC); or two <50yr	<input type="checkbox"/>	<input type="checkbox"/>
Cancer Predisposing Conditions (eg. NF2, PTEN, RB, VHL, MEN1, MEN2, TP53, NF1) _____	<input type="checkbox"/>	<input type="checkbox"/>
Familial Clustering of Cancer (type) _____	<input type="checkbox"/>	<input type="checkbox"/>

^Family History (Any FHx)

Breast / Ovary

	Personal Hx	*Family Hx
Known Breast/Ovarian Cancer Gene _____	<input type="checkbox"/>	<input type="checkbox"/>
Breast Cancer, familial	<input type="checkbox"/>	<input type="checkbox"/>
Breast Cancer <40yrs	<input type="checkbox"/>	<input type="checkbox"/>
Triple Negative Breast Cancer <50yrs	<input type="checkbox"/>	<input type="checkbox"/>
Bilateral Breast Cancer <50yrs	<input type="checkbox"/>	<input type="checkbox"/>
Male Breast Cancer	<input type="checkbox"/>	<input type="checkbox"/>
Breast/Ovary Cancer, Jewish Ancestry, any age	<input type="checkbox"/>	<input type="checkbox"/>
Breast and Ovarian Cancer	<input type="checkbox"/>	<input type="checkbox"/>
Ovarian Cancer <70yrs	<input type="checkbox"/>	<input type="checkbox"/>
Ovarian Cancer, familial	<input type="checkbox"/>	<input type="checkbox"/>

***Family History (multiple cases with 1 case <60yo OR ≥ 1 case with high risk feature)**

Other - Management issues / Personal and/or family history / other

Method of contact

- Is the patient aware of referral? Yes No
- Patient to contact clinic Yes Clinic to contact patient

Print Name _____ Provider Number _____

Signature _____ Date _____



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Holes Punched as per AS2828.1: 2012

BINDING MARGIN - NO WRITING

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