

PATIENT Surname	Given Name (Including Middle Initial)	SEX	DOB: DD / MM / YYYY	UMRN
		<input type="checkbox"/> M	Telephone (Home)	Telephone (Business)
PATIENT Address		<input type="checkbox"/> F		
		<input type="checkbox"/> Other	Is Patient of Aboriginal Descent?	<input type="checkbox"/> Yes <input type="checkbox"/> No

Homologous Recombination Deficiency (HRD) testing (validated assay, not yet NATA accredited)

PLEASE TICK

- ☐ **(please confirm):** for high grade serous or other high grade ovarian, fallopian tube or primary peritoneal carcinoma ONLY
- ☐ **(please acknowledge):** This assay may detect a variant of germline significance for both the patient and family members. Please ensure the patient is made aware of this.

CLINICAL INFORMATION:

SPECIMEN DETAILS

(Please include PDF copy of external histopathology/cytopathology)

- ☐ Histopathology ☐ Cytopathology

Specimen Number

Laboratory

- ☐ PDF histopathology / cytopathology report included

Note for external pathology: This testing requires the review of all H/E slides and use of relevant paraffin tissue blocks related to the case. The slides and blocks will be returned upon completion of testing. A small number of laboratories require payment for this request. In this instance, we will seek assistance from you, as the patient's treating oncologist, to gain access to the tissue.

SEND THIS REQUEST FORM TO

QEmolecularap.Pathwest@health.wa.gov.au

BILLING

- ☐ PUBLIC PATIENT
- ☐ PRIVATE PATIENT

(Note: The cost of the HRD test is currently covered. When the test receives NATA accreditation, it will be eligible for the Medicare Benefit Scheme rebate)

REQUESTING CONSULTANT

Name: _____ **Provider Number:** _____

Institution: _____ **Fax (private consultants only):** _____

X..... /...../.....

Requesting Doctor Signature

Patient: I consent for my results to be stored in the iCM Signature: X.....

COPY DOCTOR(S):

LABORATORY USE