

TEST REQUEST FORM FOR TESTING TO INFORM PARP INHIBITOR TREATMENT OPTIONS

Surname:		First Name:	
Date of Birth:	MRN/Hospital Number:	Gender:	
Residential Address:			
Referring Medical Oncologist (first name, surname, and hospital):		Person requesting test:	
Contact Email Address:	Clinical Team Email Address:	Pathology Email Address:	
DETAILS OF TEST REQUESTED: Breast <input type="checkbox"/> Germline BRCA only (EDTA blood) Pancreas <input type="checkbox"/> Germline BRCA only (EDTA blood) Prostate <input type="checkbox"/> Tumour BRCA and MLPA with reflex gBRCA if required (EDTA blood & FFPE tumour block to same testing site)		CURRENT DIAGNOSIS (tick one): Breast <input type="checkbox"/> HER2-negative, high risk early breast cancer previously treated with neoadjuvant or adjuvant chemotherapy <input type="checkbox"/> HER2 negative locally advanced or metastatic breast cancer (germline BRCA test only) Pancreas <input type="checkbox"/> Metastatic adenocarcinoma of the pancreas being considered for platinum treatment Prostate <input type="checkbox"/> metastatic prostate cancer (tBRCA and MLPA with reflex gBRCA if tBRCA positive)	
*For germline and tumour requests complete the form in full and include a copy of the form with the blood sample and send a <u>photocopy</u> to histopathology to include with block referral.			
CLINICAL INFORMATION: <input type="checkbox"/> Patient is being considered for adjuvant treatment of HER2-negative high risk early breast cancer previously treated with neoadjuvant or adjuvant chemotherapy <input type="checkbox"/> Patient is being considered for PARP inhibitor treatment of HER2-negative locally advanced or metastatic breast cancer. <input type="checkbox"/> Patient is being considered for maintenance treatment with olaparib for metastatic adenocarcinoma of the pancreas. <input type="checkbox"/> Patient is being considered for PARP inhibitor treatment of mCRPC			
To be completed by patient: <ul style="list-style-type: none"> I have read the written information given to me, understand the implications and limitations of the test, have discussed it with..... and consent to BRCA gene testing of my blood and/or tissue sample YES / NO (please circle) I consent that DNA from my blood and/or tissue sample will be stored in the.... laboratory as standard practice, unless I request its disposal (YES/NO) (please circle) I consent that my genetic test result can be made available for use in counselling other family members YES/NO (please circle) I consent for this sample to be used for quality assurance and audit purposes YES/NO (please circle) If I am unable to receive the results of the test, I would like the result to be given to the following person(s) Name: Relationship:..... Contact no:..... Signed: Date:			
For completion by referring doctor: <ul style="list-style-type: none"> I have discussed this test with my patient and they understand the implications of the test and the potential need for referral to the cancer genetics service. Signature..... Name (block capitals)..... Contact Number..... Medical Council Registration Number:			

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Sample Details (complete as appropriate)	
Germline or MLPA (blood) samples	Tumour/FFPE samples
<input type="checkbox"/> Blood Sample (>3ml EDTA) <input type="checkbox"/> Tube labelled with patient name, DOB and MRN Sample Taken by (Full Name): Date Taken: Signature:	Prostate <input type="checkbox"/> Pre-chemotherapy biopsy sample (preferred) <input type="checkbox"/> Post-chemotherapy biopsy sample Age of sample: All samples <input type="checkbox"/> Pathology report attached (required) Pathologist name (Full Name): Hospital Name: Case Number: Signature:

For Germline testing only:

- Sample required is 3-5ml of venous blood in EDTA anticoagulant. Send at room temperature by courier to: Beaumont Hospital Molecular Pathology Laboratory, Beaumont Hospital, Dublin 9, D09 V2N0 Refrigerate if there will be more than a 24 hr delay before posting. DO NOT FREEZE.
- Note the minimum identification requirements for genetic testing are
 - a) patient's forename & surname and date of birth or medical record number
 - b) these identifiers must be present on the sample tube and the genetic test request form and must match exactly.
- Queries regarding the sample, sample identification requirements or transport should be directed to molecular@beaumont.ie / 01-809 3726

For tumour testing only:

- Complete form and photocopy.
- Include one copy of the form with the blood sample.
- Forward a second copy of the form to the histopathology laboratory for block selection. A pathologist will review the available material and select the most appropriate block for testing.
- This block will be sent to Beaumont Hospital Molecular Pathology Laboratory and a report issued.
- A copy of the tumour report will also be sent to the histopathology laboratory for their records.

For combined tumour and germline testing:

- Complete form and photocopy.
- Include one copy of the form with the blood sample.
- Forward a second copy of the form to the histopathology laboratory for block selection.
- A copy of the tumour report will be forwarded to the histopathology laboratory however the germline results and integrated report will only be forwarded to the requesting oncologist.

Information for Pathologists:

- Please indicate if it is a pre-chemotherapy or a post-chemotherapy biopsy sample as this may impact testing outcome
- Please select the block with the largest tumour content (ideally >50% high grade serous carcinoma tumour nuclei content, with minimal necrosis for ovarian samples and block with highest tumour cellularity available for prostate samples), however please note this will be re-assessed at the reference lab also)
- Sending of samples should be prioritised.
- Send the sample with a copy of the histopathology report by courier to: Beaumont Hospital Molecular Pathology Laboratory, Beaumont Hospital, Dublin 9, D09 V2N0.