

Patient Details

Genetic Tumour Test Request Form

North West Genomic Laboratory Hub (MANCHESTER), Manchester Centre for Genomic Medicine (MCGM)

Referring Clinician

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	9865	

Surname:		Consultant (in full):	9865	
Forename:		Hospital (in full):		
DoB :	NHS No:	Department:	Tel:	
Sex:	Hospital No:	Email:		
Address:		Copy report to (if applicable):		
Postcode:				

3. TEST REQUEST (please select options by placing a tick or cross next to each test required)

Payment Status: NHS Private

For sample requirements please see reverse or <u>https://mft.nhs.uk/nwqlh/</u> for further details. For full details of genes covered see national genomic cancer test directory (<u>https://www.england.nhs.uk/publication/national-genomic-test-directories/</u>)

*Requires 2x tubes of sections; ** Requires additional material if also for another test; *** please complete FFPE CRC pro-forma also

Test/Gene	Required	Test/Gene	Required
Non small cell lung cancer*			
(for plasma testing please see alternative form)		 NTRK1,2,3 fusion ** 	
Fusion only (inc ALK, RET, ROS1)			
		Melanoma (BRAF codon 600 only)	
Colorectal BRAF/RAS testing		NGS Panel (BRAF, NRAS, KIT)	
		Ovarian (BRCA1/BRCA2)	
Somatic Lynch syndrome panel***		Ovarian (SMARCA4)	
		MLH1 hypermethylation**	
GIST		plus BRAF codon 600	
Microsatellite instability (MSI)		Sarcoma/Paediatric Tumour fusion panel** Please include any specific targets indicated	
Other test directory panel – please state tumour type (Excluding neurological tumours – see separate referral form)		FISH ALK/ROS1	

4. PATHOLOGY AND CLINICAL DETAILS	5. PATHOLOGY		
Tumour type/organ of origin	Date sections sent to Genetics lab:		
PLEASE INCLUDE A COPY OF THE PATHOLOGY REPORT	Please circle the approximate neoplastic cells (%) in the sample		
Pathologist:	sent for analysis (this information is important and is used to ensure the test carried out is appropriately sensitive)		
Hospital/Trust:			
Pathology block/sample no.	<10%# 10-20% [#] >20%		
	Neoplastic cells in marked area are %		
EGFR clinically urgent referral (targeted EGFR testing only)	[#] Where overall neoplastic cell content <20% and macrodissection would enhance % of neoplastic cells, please send slide mounted sections with corresponding marked H&E stained slide.		
Yes/No			
Please note material received at <20% NCC or which yields insufficient material for wider panel screening will be tested using a targeted test where available			

PLEASE COMPLETE SECTION 1-3 AND EITHER FORWARD TO THE PATHOLOGY LABORATORY HOLDING THE SAMPLE, OR IF YOU REQUIRE THE GENOMIC DIAGNOSTICS LABORATORY TO OBTAIN THE SPECIMEN PLEASE FORWARD TO mft.Pharmaco.GeneticsRequests@nhs.net. SECTIONS 4-5 TO BE COMPLETED BY THE PATHOLOGY LABORATORY.

INFORMATION FOR PATHOLOGY LAB (ALL SAMPLES)

- Formalin fixed paraffin embedded (FFPE) material should be reviewed by a histo/cyto-pathologist to identify areas containing neoplastic cells and determine suitability for testing.
- Sections should be cut under conditions that prevent cross contamination from other specimens.
- Scrolls should be sent in a sterile tube labelled with at least 2 patient identifiers, one of which should be the pathology sample number. Containers and slides should also be labelled with at least 2 patient identifiers one of which should be the pathology sample number.
- For each additional test indicated to need additional material please send an additional tube of scrolls.
- Please avoid baking slides or heating samples
- Please send appropriate corresponding paperwork with the samples
- Please contact the laboratory for additional guidance or if you are unsure whether a sample is suitable

FISH TESTS

- Prepare 4 unstained sections (4uM thick) floated on the surface of a purified water bath set at 40°C (+/-2°C).
- Mount on positively charged slides and allow to air-dry
- Also include 1 H&E slide with regions enriched for neoplastic cells marked by a Pathologist along with an estimate of neoplastic cell content in the marked area(s)

