

Guide to referring for Whole Genome Sequencing (WGS) for Solid Tumours

Before referring:

- Check patient is eligible for WGS and select appropriate M code using the National genomic test directory for cancer https://www.england.nhs.uk/publication/national-genomic-test-directories/
- Discuss patient feasibility and eligibility at MDT.

Referral Process:

- 1. Fill in a Record of Discussion with the patient https://southeastgenomics.nhs.uk/wp-content/uploads/2021/04/GMS-WGS-RoD-Pilot-April-2021.pdf
 - o Information sheets available for patients here:
 - WGS https://southeastgenomics.nhs.uk/wp-content/uploads/2020/12/Patient Information Cancer v2.2.pdf
 - National Genomic Research Library: https://southeastgenomics.nhs.uk/wp-content/uploads/2020/12/Patient_Information_Research_V1.3.pdf
- 2. Complete interventional radiology / surgery request with specific request for tissue to be **fresh frozen**, and not to be placed in formalin.
- 3. Request an EDTA blood sample.
- 4. Member of the clinical team to complete sections highlighted in yellow on the Test Order Form (details below).
- 5. Member of the pathology team will sample tumour appropriately for WGS and snap-freeze (OCT, iso-pentane cooled in dry ice/LN2 (store in freezer at -80C)). Frozen section slide reviewed by histopathology team to confirm tumour and tumour cellularity.
- 6. Member of the pathology team to complete sections highlighted in blue on the Test Order Form.
- 7. Completed Test Order Form and Record of Discussion emailed to gst-tr.wgs@nhs.net with CANCER WGS NHS Number: XXX XXXX in the subject.
- 8. Frozen tissue, blood sample and TOF sent to Guy's Hospital (Synnovis Genetics Laboratory, 5th Floor Tower Wing, Guy's Hospital, London SE1 9RT) on dry ice and DNA Duty Tech Lead emailed to inform the sample is on its way: DNADutyTechLead@viapath.co.uk
 - a. The sample should ideally be sent with a morning courier, Monday-Thursday)



NHS Genomic Medicine Service, WGS Test Request Cancer, October 2022 v1.19 to be used for WGS go-live.

This document is subject to version control and is regularly updated. Please confirm you are using the current version by contacting your local Genomic Laboratory Hub

	Genomic Medicine Service Whole Genome Sequencing (WGS) Test Request PLEASE DO NOT USE FOR NON-WGS TESTS	CANCER		NHS
	Requesting organisation: GLH laboratory to receive sample: Test Required Whole Genome Sequencing			
	Patient first name	Ethnicity		
	Patient last name Date of birth (66/mm/yyyy) Hospital number	Other Presentation status First diagnosis Recurrence / Relapse Unknown		
	Gender Male Female Other			
	Postcode NHS number Reason NHS Number not available: Patient not eligible for NHS number (e.g. foreign national) Other (provide reason):	Additional clinical information (if required) E.g. previous tumours, molecular testing, and relevant treatment history with date(s)		
	Solid tumour requests only Primary Metastatic Unknown Lymphoma Mate of this diagnosis (Adfront/YYY)	Additional tumour inform E.g. site of metastasis (if metastatic Tumour topography), or unkn	if relevant) nown primary ur morphology
	Haemato-oncology liquid tumour requests only AML ALL Other (please specify):	SIHMDS Lab ID	Date of this diagnosis (dd/mm/yyy)	
1	Complete for tumour samples (being sent to GLH DNA extraction lab)			
ł	Fresh frozen tumour Bone marrow Blood (EDTA) Other (please specify): % malignant nuclei / blasts or equivalent in this sample (refer to sample handling guidance) must be provided below			
		% Malignant nuclei / blast		If BM/PB provide volume and nucleated cell count
Complete for germline samples (being sent to GLH DNA extraction lab)				
Blood (EDTA) Saliva Fibroblasts Skin biopsy Other (please specify):				iv)-
ŀ	Sample ID Collection date / time	Sample volume if applicab		Comments
Y				
	Responsible consultant Name: Department address:	Main contact (if different from responsible consultant) (Name: (Department address:		
	Phone: Comparison to the Record of Discussion form to follow.			

Requesting organisation: Your hospital

GLH laboratory to receive sample: South East GLH

Providing ethnicity ensures that there is equity of access to genetic testing in the wider population and it also helps us to improve variant analysis and interpretation. Please select from the drop-down options.

Please select the appropriate M code using the drop-down options. If you need further guidance, please refer to the National Genomic Test Directory.

Please provide details of family history of cancer, details of any treatment including radiotherapy and previous chemotherapy, any concurrent or previous tumour

Important to include an NHS number as required for the WGS pipeline. If no NHS number is available a reason will need to be provided.

Please indicate whether the sample is: 'Primary, Metastatic, unknown, Lymphoma'. Please also indicate date of first diagnosis.

Histopathology Lab ID: this is based on patient and surgery

Sample ID: this is based on the particular block or sample. **Collection date / time:** when the tissue was removed. **% Malignant nuclei / blasts:** invasive malignant nuclei must account for at least 30% of the nuclei present in the tissue sample submitted for WGS. There should also be less than 20% necrosis.

Please add details of the germline sample (EDTA) if you have these details.

The 'responsible contact' should be filled in by the consultant who the report should be sent to, 'main contact' should have the histopathologist details.