

Specialist Integrated Haematological Malignancy Diagnostic Service (SIHMDS)

Pathology Sciences, North Bristol NHS Trust, Southmead Hospital, Westbury-on-Trym, Bristol BS10 5NB

Website: [Haematological Malignancy Diagnostics \(SIHMDS\) | North Bristol NHS Trust \(nbt.nhs.uk\)](http://HaematologicalMalignancyDiagnostics(SIHMDS)|NorthBristolNHSTrust(nbt.nhs.uk))Flow lab: Flowlab@nbt.nhs.uk Tel. 0117 414 8377

Cellular pathology consultants: CellularPathologyHaemPathTeam1@nbt.nhs.uk

PATIENT DETAILS

Surname:	Given Name:	Originating Hospital	Consultant
NHS Number:	Hospital Number:	Postcode	<input type="checkbox"/> NHS <input type="checkbox"/> Private <input type="checkbox"/> Research
Date of Birth: <input type="checkbox"/> Male <input type="checkbox"/> Female	Priority: <input type="checkbox"/> Urgent <input type="checkbox"/> Routine	Previous investigation at BHODS? Yes <input type="checkbox"/> / No <input type="checkbox"/>	Known danger of infection sample? Yes <input type="checkbox"/> / No <input type="checkbox"/>

SPECIMEN DETAILS**Specimen type(s):** *tick all that apply*

- ☐ Marrow aspirate
☐ Marrow trephine
☐ Blood
☐ Lymph Node
☐ Skin (germline)
☐ Saliva
☐ Other fluid / tissue

*Please specify***Specimen Details:**

Biopsy site:

Specimen date:

Specimen time:

☐ Diagnostic☐ Relapse☐ Follow-up / MRD**CLINICAL DETAILS / SUSPECTED DIAGNOSIS:**

Details of relevant current/previous treatment:

Hb:

Lymphocytes:

WBC:

Platelets:

Neutrophils:

Other:

INVESTIGATIONS REQUIRED:*See overleaf for specimen requirements*☐ Aspirate morphology - Further details:☐ Flow cytometry - Further details:☐ Cytology / Cellular Pathology - Further details:☐ Genetic Testing**Please ensure BOTH SIDES of this request are completed and returned with genetics samples.** Genetic testing will be undertaken according to the [National Genomic Test Directory](#) and, unless otherwise indicated, agreed diagnostic algorithms.**REQUESTOR DETAILS:**

Specimen sent by:

Contact details:

In submitting this sample the clinician confirms that consent has been obtained to test for the suspected disorder and for cellular/DNA/RNA storage. May this sample be used for research? Yes ☐ / No ☐

GENOMIC TESTING: Further information on genomic testing is available from the [South West Genomic Laboratory Hub](#) and the [National Genomic Test Directory for Cancer](#).

Indication	Test	Sample	Indication	Test	Sample
ALL[M91]	<input type="checkbox"/> Diagnosis/Relapse	A & B	MDS [M82]	<input type="checkbox"/> Karyotype [M82.2]	A
	<input type="checkbox"/> <i>TPMT/NUDT15</i>	B		<input type="checkbox"/> SNP Array [M82.2]	B
	<input type="checkbox"/> MRD Monitoring [M91.23]	B		<input type="checkbox"/> MDS gene panel [M82.1]	B
	<input type="checkbox"/> Other Monitoring:	B	MDS/MPN [M224]	<input type="checkbox"/> Karyotype [M224.2]	A
AML[M80]	<input type="checkbox"/> Diagnosis/Relapse	A & B		<input type="checkbox"/> <i>PDGFR</i> FISH [M224.6]	A
	<input type="checkbox"/> APL (PML/RARA) [M80.29]	A & B		<input type="checkbox"/> RNA Fusion Panel [M224.6]	A
	<input type="checkbox"/> MLDS panel (<i>GATA1</i>) [M80.19]	A & B		<input type="checkbox"/> MDS/MPN gene panel [M224.1]	B
	<input type="checkbox"/> <i>NPM1</i> Monitoring [M80.9]	B		<input type="checkbox"/> <i>BCR-ABL1</i> [M224.10]	
	<input type="checkbox"/> Monitoring (Specify target):	B	MPN [M85]	<input type="checkbox"/> JMML gene panel [M88.1]	B
CLL [M94]	<input type="checkbox"/> IgVH [M94.5]	B		<input type="checkbox"/> <i>JAK2</i> (V617F) [M85.14]	B
	<input type="checkbox"/> <i>TP53</i> (17p) deletion [M94.4] and mutation [M94.1]	A & B		<input type="checkbox"/> <i>JAK2</i> (ex12), <i>CALR</i> , <i>MPL</i> [M85.1]	B
				<input type="checkbox"/> <i>BCR-ABL1</i> [M85.11]	B
CML [M84]	<input type="checkbox"/> <i>BCR-ABL1</i> diagnosis [M84.1]	B		<input type="checkbox"/> MPN gene panel [M85.2]	B
	<input type="checkbox"/> <i>BCR-ABL1</i> monitoring [M84.2]	B		<input type="checkbox"/> Karyotype [M85.3]	A
	<input type="checkbox"/> <i>BCR-ABL1</i> TKD NGS [M84.8]	B		<input type="checkbox"/> RNA Fusion Panel [M85.7]	A
	<input type="checkbox"/> Karyotype [M84.4]	A		<input type="checkbox"/> <i>PDGFR</i> FISH [M85.7]	A
CNL [M87]	<input type="checkbox"/> <i>CSF3R</i> [M87.1]	B	Myeloma [M92]	<input type="checkbox"/> FISH panel	A
	<input type="checkbox"/> <i>BRAF</i> [M108.1]	B	SM [M86]	<input type="checkbox"/> <i>KIT</i> (D816V) [M86.2]	B
	<input type="checkbox"/> LGL panel (<i>STAT3/STAT5B</i>) [M114.1]	B	Storage	<input type="checkbox"/> SM panel [M224.1]	
	<input type="checkbox"/> LPL panel (<i>MYD88/CXCR4</i>) [M104.1]	B		<input type="checkbox"/> Cell suspension	A
HCL				<input type="checkbox"/> Plasma cell enriched	A
LGL[M114]				<input type="checkbox"/> DNA	B
LPL [M104]				<input type="checkbox"/> RNA	B
			TAM [M81]	<input type="checkbox"/> <i>GATA1</i> [M81.1]	B
			Whole Genome Sequencing (WGS)*:		
			<input type="checkbox"/> Acute Leukaemia		B & C
			<input type="checkbox"/> WGS for TYA (≤ 25 years old)		
			<input type="checkbox"/> WGS for proven or suspected haematological tumour exhausted all standard of care testing/treatment		

Test	Sample Requirements
Morphology	Blood / Marrow aspirate samples; PB slides x2 + BM slides x3, PB 5ml in EDTA +/-or BM Asp 2ml in EDTA
Flow cytometry	Blood or bone marrow in EDTA
Genetics	A) 5ml blood in Lithium Heparin/1-2ml BMA 1-2ml in heparinised tissue culture medium/ 1-2ml BMA in EDTA
	B) 10-20ml blood/1-2ml BMA in EDTAx2
	Samples for molecular monitoring and WGS to reach the lab within 72 hours
	C) Skin biopsy for germline analysis, fresh (NOT fixed) sample in sterile saline to reach the lab within 72 hours.
	Molecular and FISH testing is possible on formalin fixed, paraffin embedded tissue sections. Please contact the laboratory for further details if required.
Tissue/Other samples	*WGS Tumour sample: FRESH (not fixed) sample with >30% tumour, <20% necrosis
	Germline sample: skin (C) or remission (<0.1% MRD) blood/marrow (B)
	Record of Discussion and Test Order form . Contact rde-tr.swgenomicpractitioner@nhs.net for further information.
	Lymph node FNA for cytology: smear slides, needle washings in CytoLyt
	Fresh tissue for flow cytometry and genetics: Saline or cell culture medium (without coagulation or preservative). Please alert laboratory flowlab@nbt.nhs.uk tel: 0117 414 8377 NB Samples placed in Formalin or CytoLyt (an alcohol preservative) cannot be used for flow cytometry
	Lymphoreticular testing on formalin fixed paraffin embedded tissue (FFPE) should be requested using the Cancer Genomics Request Form .