


# Schedule of Accreditation

issued by

## United Kingdom Accreditation Service

2 Pine Trees, Chertsey Lane, Staines-upon-Thames, TW18 3HR, UK

 <b>Accredited to ISO 15189:2022</b>	<b>Leeds Teaching Hospitals NHS Trust</b>	
	<b>Issue No:</b> 009 <b>Issue date:</b> 15 February 2024	
	<b>Haematological Malignancy Diagnostic Service</b> <b>St. James' University Hospital</b> <b>Leeds</b> <b>LS9 7TF</b> <b>United Kingdom</b>	<b>Contact:</b> Dr Catherine Cargo / Dr Ruth De Tute <b>Tel:</b> +44 (0)1132067963 <b>Fax:</b> +44 (0)1132067883 <b>E-Mail:</b> catherine.cargo@nhs.net / rdetute@nhs.net <b>Website:</b> www.hmds.info
<b>Testing performed at the above address only</b>		

### DETAIL OF ACCREDITATION

Materials/Products tested	Type of test/Properties measured/Range of measurement	Standard specifications/ Equipment/Techniques used
<b>HUMAN BODY TISSUE AND FLUIDS</b>  Tissue          Tissue within paraffin blocks (produced using procedures above or received as a primary sample type)	<u>Histopathology examination activities in order to identify or exclude morphological abnormalities for the purpose of haematological malignancy diagnosis</u>	Macroscopic and Microscopic examination:  Documented in house methods incorporating manufacturers' instructions where relevant:  Tissue dissection: Morphology SOP HP01, HP39  Decalcification SOP HP44  Tissue processing/embedding (paraffin): Thermo Scientific Excelsior AS tissue processor & Leica HistoCore Arcadia H embedding centre SOP HP04  Microtomy (paraffin blocks): Microm HM325 & Leica HistoCore Autocut microtomes, water baths, hot-plates SOP HP06, HP32



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<p><b>HUMAN BODY TISSUE AND FLUIDS (cont'd)</b></p> <p>Paraffin (produced using procedures above or received as a primary sample type) embedded tissue</p>	<p><u>Histopathology examination activities in order to identify or exclude morphological abnormalities for the purpose of haematological malignancy diagnosis (cont'd)</u></p> <p>Identification of:</p> <p>Basophilic and eosinophilic structures</p> <p>Amyloid</p> <p>Differentiation of different haemopoietic cells</p> <p>Glycogen (and other PAS positive substances)</p> <p>Heamosiderin</p> <p>Reticulin fibres</p> <p>Tubercle bacilli</p> <p>Identification of:</p> <p>T-cells, mantle zone B-cells , follicular lymphomas</p> <p>Germinal centre B-cells &amp; related lymphomas</p> <p>Activated B-cells, plasma cells</p>	<p>Macroscopic and Microscopic examination:</p> <p>Documented in house methods incorporating manufacturers' instructions where relevant:</p> <p>Automated Haematoxylin &amp; Eosin and May Grunwald Giemsa staining using Leica ST5020, light microscope SOP HP11, SOP HP13</p> <p>Manual Special tinctorial staining: Microm coverslipper, light microscope SOP HP12, HP14, HP15, HP19, HP21</p> <p>Congo Red (HP21)</p> <p>May Grunwald Giemsa (HP13)</p> <p>PAS (+/- diastase) (HP19)</p> <p>Perls' (HP14)</p> <p>Reticulin (Gordon &amp; Sweets) (HP12)</p> <p>Ziehl-Neelsen (HP15)</p> <p>Immunohistochemistry: Dako Autostainers, Dako Omnis, Manual staining SOP HP27, HP40, HP42 Using the following monoclonal and polyclonal antibodies:</p> <p>Bcl-2 (clones 124 &amp; E17)</p> <p>Bcl-6</p> <p>Blimp-1</p>



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HUMAN BODY TISSUE AND FLUIDS (cont'd)	<p><u>Histopathology examination activities in order to identify or exclude morphological abnormalities for the purpose of haematological malignancy diagnosis (cont'd)</u></p> <p>Identification of: (cont'd)</p> <p>Langerhan's cells, interdigitating dendritic cells, associated disease states</p> <p>Pan T-cell marker &amp; T-cell lymphomas</p> <p>Pan T-cell marker &amp; T-cell lymphomas</p> <p>T-helper cells &amp; related lymphoma</p> <p>T-cells, mantle zone B-cells, neoplastic B-cells ( CLL, MCL), T-cell lymphomas</p> <p>Early T-cells &amp; NK cells, related T-cell lymphomas</p> <p>Mature suppressor / cytotoxic T-cells, associated T-cell lymphomas</p> <p>Germinal centre B cells &amp; related lymphomas</p> <p>Monocytes, macrophages, Hairy Cell Leukaemia cells</p> <p>Myeloid cells, Reed Sternberg cells</p> <p>B-cells &amp; associated B-Cell lymphomas, normal plasma cells</p> <p>B-cells &amp; associated B-Cell lymphomas</p>	<p>Macroscopic and Microscopic examination:</p> <p>Documented in house methods incorporating manufacturers' instructions where relevant:</p> <p>Immunohistochemistry: Dako Autostainers, Dako Omnis, Manual staining SOP HP27, HP40, HP42 Using the following monoclonal and polyclonal antibodies:</p> <p>CD1a</p> <p>CD2</p> <p>CD3</p> <p>CD4</p> <p>CD5</p> <p>CD7</p> <p>CD8</p> <p>CD10</p> <p>CD11c</p> <p>CD15</p> <p>CD19</p> <p>CD20</p>



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HUMAN BODY TISSUE AND FLUIDS (cont'd)	<p><u>Histopathology examination activities in order to identify or exclude morphological abnormalities for the purpose of haematological malignancy diagnosis (cont'd)</u></p> <p>Identification of: (cont'd)</p> <p>Follicular dendritic cells, mature B-cells</p> <p>B-cells, associated B-cell lymphomas</p> <p>Follicular dendritic cells, B-cells, CLL cells</p> <p>Activated T-cells, inter-leukin-2 receptor cells</p> <p>Activated B &amp; T-cells, Reed Sternberg / Hodgkin's cells</p> <p>Endothelial cells, blast cells</p> <p>Pan Leucocyte marker- normal &amp; neoplastic cells</p> <p>NK cells, nerve cells, neuroblastomas, some neoplastic PCs</p> <p>NK cells, related lymphomas</p> <p>Megakaryocytes, megakaryocytes precursors, and platelets</p> <p>Macrophages &amp; myeloid cells</p> <p>B-cells &amp; associated B-Cell lymphomas</p> <p>Ewings sarcoma, lymphocytes</p>	<p>Macroscopic and Microscopic examination:</p> <p>Documented in house methods incorporating manufacturers' instructions where relevant:</p> <p>Immunohistochemistry: Dako Autostainers, Dako Omnis, Manual staining SOP HP27, HP40, HP42 Using the following monoclonal and polyclonal antibodies:</p> <p>CD21</p> <p>CD22</p> <p>CD23</p> <p>CD25</p> <p>CD30</p> <p>CD34</p> <p>CD45</p> <p>CD56</p> <p>CD57</p> <p>CD61</p> <p>CD68</p> <p>CD79a</p> <p>CD99</p>



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HUMAN BODY TISSUE AND FLUIDS (cont'd)	<p><u>Histopathology examination activities in order to identify or exclude morphological abnormalities for the purpose of haematological malignancy diagnosis (cont'd)</u></p> <p>Identification of: (cont'd)</p> <p>Melanocytes, mast cells, AML tumour cells, GIST</p> <p>Kikuchi disease</p> <p>Normal &amp; neoplastic plasma cells, epithelial cells</p> <p>Assist in identification of positive cells in myeloid sarcomas and T-ALL</p> <p>Macrophages, monocytes</p> <p>Alk + tumours (ALCL)</p> <p>Plasma cells, plasmacytic lymphoma vs myeloma</p> <p>Normal &amp; neoplastic epithelial cells (intestinal)</p> <p>Identification of myeloid sarcomas and T-ALL</p> <p>Normal &amp; neoplastic epithelia (breast)</p> <p>Normal &amp; neoplastic epithelia (colon)</p> <p>Cells showing Myc translocation, disease related</p> <p>Epithelial cells, Mantle cell lymphoma</p> <p>Broad spectrum of normal &amp; neoplastic epithelia</p>	<p>Macroscopic and Microscopic examination:</p> <p>Documented in house methods incorporating manufacturers' instructions where relevant:</p> <p>Immunohistochemistry: Dako Autostainers, Dako Omnis, Manual staining SOP HP27, HP40, HP42 Using the following monoclonal and polyclonal antibodies:</p> <p>CD117 C-KIT</p> <p>CD123</p> <p>CD138</p> <p>CD13</p> <p>CD163</p> <p>CD246</p> <p>CD319 (manual staining only)</p> <p>CDX2</p> <p>CD33</p> <p>CK 7</p> <p>CK20</p> <p>C Myc</p> <p>Cyclin D1</p> <p>Cytokeratin - Cam 5.2</p>



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HUMAN BODY TISSUE AND FLUIDS (cont'd)	<p><u>Histopathology examination activities in order to identify or exclude morphological abnormalities for the purpose of haematological malignancy diagnosis</u> (cont'd)</p> <p>Identification of: (cont'd)</p> <p>Broad spectrum of normal &amp; neoplastic epithelia</p> <p>Follicular helper T-cells, angioblastic T-cell lymphomas</p> <p>Suggestive of Hairy Cell Leukaemia</p> <p>Smooth &amp; striated muscle</p> <p>Endothelial cells, Megakaryocytes</p> <p>Erythroid cells &amp; precursors</p> <p>Cytotoxic T-cells &amp; NK cells &amp; related lymphomas</p> <p>HLA class II DR antigen expressing cells</p> <p>Karposi sarcoma virus, associated multicentric Castleman's disease</p> <p>IgA expressing plasma cells</p> <p>IgD expressing plasma cells, Mantle zone B-cells</p> <p>IgG expressing plasma cells &amp; lymphocytes</p> <p>Specific subtype of plasma cells, IgG4 disease</p>	<p>Macroscopic and Microscopic examination:</p> <p>Documented in house methods incorporating manufacturers' instructions where relevant:</p> <p>Immunohistochemistry: Dako Autostainers, Dako Omnis, Manual staining SOP HP27, HP40, HP42 Using the following monoclonal and polyclonal antibodies:</p> <p>Cytokeratin - MNF116</p> <p>CXCL13</p> <p>Leukaemia Hairy cell (DBA44)</p> <p>Desmin</p> <p>Factor VIII</p> <p>Glycophorin C</p> <p>Granzyme B</p> <p>HLA-DR</p> <p>HHV8</p> <p>IgA (manual staining only)</p> <p>IgD</p> <p>IgG (manual staining only)</p> <p>IgG4 (manual staining only)</p>



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HUMAN BODY TISSUE AND FLUIDS (cont'd)	<p><u>Histopathology examination activities in order to identify or exclude morphological abnormalities for the purpose of haematological malignancy diagnosis (cont'd)</u></p> <p>Identification of: (cont'd)</p> <p>IgM expressing plasma cells, mantle zone B-cells</p> <p>Plasma cells, activated T-cells, some germinal centre B-cells, R/S cells &amp; associated lymphomas</p> <p>Kappa light chain expressing plasma cells and lymphoid cells (mantle zone B-cells)</p> <p>Lambda light chain expressing plasma cells and lymphoid cells (mantle zone B-cells)</p> <p>Langerhan's cells, IRDC's</p> <p>Diagnosis of follicular lymphoma</p> <p>EBV</p> <p>Histiocytes</p> <p>Mast cells, related disorders</p> <p>Melanoma cells</p> <p>Proliferating cells</p> <p>Myeloid cells</p> <p>Rhabdomyosarcoma cells</p> <p>Rhabdomyosarcoma &amp; Ewings tumour cells</p>	<p>Macroscopic and Microscopic examination:</p> <p>Documented in house methods incorporating manufacturers' instructions where relevant:</p> <p>Immunohistochemistry: Dako Autostainers, Dako Omnis, Manual staining SOP HP27, HP40, HP42 Using the following monoclonal and polyclonal antibodies:</p> <p>IgM (manual staining only)</p> <p>IRF4</p> <p>Kappa (manual staining only)</p> <p>Lambda (manual staining only)</p> <p>Langerin</p> <p>LMO2</p> <p>LMP-1</p> <p>Lysozyme</p> <p>Mast-cell tryptase</p> <p>Melanoma monoclonal</p> <p>MIB1</p> <p>Myeloperoxidase</p> <p>Myo D1</p> <p>Myogenin</p>



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HUMAN BODY TISSUE AND FLUIDS (cont'd)	<p><u>Histopathology examination activities in order to identify or exclude morphological abnormalities for the purpose of haematological malignancy diagnosis (cont'd)</u></p> <p>Identification of: (cont'd)</p> <p>B-cells, Germinal centre cells, plasma cells</p> <p>Embryonic cells &amp; germ cells &amp; related diseases</p> <p>B-cells</p> <p>Follicular Helper T-cells</p> <p>Neural cells, neuroblastoma cells</p> <p>Prostate glandular tissue, related carcinomas</p> <p>Transcription factor, monocytic lineage, early granulocytes</p> <p>Neural cells, melanocytes</p> <p>Mantle cell lymphoma</p> <p>Normal T &amp; B-lymphocyte precursors. T-ALL &amp; B-ALL cells</p>	<p>Macroscopic and Microscopic examination:</p> <p>Documented in house methods incorporating manufacturers' instructions where relevant:</p> <p>Immunohistochemistry: Dako Autostainers, Dako Omnis, Manual staining SOP HP27, HP40, HP42 Using the following monoclonal and polyclonal antibodies:</p> <p>OCT-2</p> <p>OCT-3/4</p> <p>PAX-5</p> <p>PD1</p> <p>PGP 9.5</p> <p>PSMA</p> <p>PU-1</p> <p>S100</p> <p>SOX-11</p> <p>TDT</p>





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<p><b>HUMAN BODY FLUIDS/TISSUE</b></p> <p>Whole Blood Bone Marrow</p> <p>Whole Blood Bone Marrow</p> <p>Whole Blood Bone Marrow Separated cells from blood and bone marrow Fixed Cells Effusions CSF Vitreous biopsies</p>	<p><u>Genomics analysis for the purpose of clinical diagnosis, prognosis, and patient management</u></p> <p>Sample processing, DNA and RNA extraction, quantification and quality check for subsequent in-house analysis (see below), referral to specialist centres and long term storage</p>	<p>In house documented methods incorporating manufacturers' instructions where relevant</p> <p>Manual and Automated separation of cell fractions</p> <p><b>Automated</b> Separation of cell fractions:</p> <p>Using</p> <p>Miltenyi Automacs Pro SOP FC028</p> <p><b>Manual</b> Separation of lineage specific fractions</p> <p>Using:</p> <p>Dynal Magnetic Separator</p> <p>SOP MH05</p> <p>Manual and semi-automated and automated DNA extraction and quantification using:</p> <p><b>Semi-Automated Extraction</b> Qiagen QIAcube with Qiagen QIAamp DNA Blood Mini Kit</p> <p>SOP MH02 SOP MH62</p>



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<p>HUMAN BODY FLUIDS/TISSUE (cont'd)</p> <p>FFPE Tissue Slide Sections Fresh Solid Tissue</p> <p>Genomic DNA extracted in-house from or received as primary sample type from an external source</p> <p>Whole Blood Bone Marrow Separated Cells from Blood and Bone Marrow</p>	<p><u>Genomics analysis for the purpose of clinical diagnosis, prognosis, and patient management (cont'd)</u></p> <p>RNA extraction and preparation of cDNA, for subsequent in-house analysis (see below), referral to specialist centres and long term storage</p>	<p>In house documented methods incorporating manufacturers' instructions where relevant</p> <p><b>Semi-Automated Extraction (cont)</b> Qiagen QIAcube with Qiagen QIAamp DNA FFPE Tissue Kit</p> <p>SOP MH02 SOP MH62</p> <p>DNA Quantification for QC purposes using: Promega Glomax 96 well fluorimeter</p> <p>SOP MH62</p> <p>Manual and semi-automated RNA extraction, conversion to cDNA and quantification using:</p> <p>Manual methods: Qiagen RNeasy Mini Kit</p> <p>SOP MH02</p> <p>Semi-Automated methods: Promega Maxwell RSC and Maxwell RSC simplyRNA blood kit</p> <p>SOP MH02</p>



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HUMAN BODY FLUIDS/TISSUE (cont'd)  Genomic DNA extracted in-house from the sample types listed above or received as primary sample type from an external source	<u>Genomics analysis for the purpose of clinical diagnosis, prognosis, and patient management</u> (cont'd)  Detection of nucleic acid sequence variants - SNVs and small indels  [Definitive list HMDS4406]	In house documented methods incorporating manufacturers' instructions where relevant  <b>Reverse Transcription</b>  Manual method  using  Invitrogen M-MLV reverse transcriptase, thermal cycler and UV cross linker  SOP MH02  <b>Sanger Sequencing</b>  Using:  Standard primer design methodology using SOP MH75  And  PCR using thermal cyclers  Sanger Sequencing by: (Applied Biosystems (ABI) Analysers 3500  Analysis using Mutation Surveyor SOP MH74



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<p>HUMAN BODY FLUIDS/TISSUE (cont'd)</p> <p>Genomic DNA extracted in-house from the sample types listed above or received as primary sample type from an external source</p>	<p><u>Genomics analysis for the purpose of clinical diagnosis, prognosis, and patient management</u> (cont'd)</p> <p>Determination of fragment length size and detection of deletions, known SNVs and indels, gene rearrangements, internal tandem duplications and microsatellites</p> <p>[Definitive list HMDS4407]</p>	<p>In house documented methods incorporating manufacturers' instructions where relevant</p> <p><b>Fragment Analysis</b></p> <p>Fluorescent Based Fragment Analysis</p> <p>Using Commercial Kits:</p> <p>Powerplex 16 – Chimerism Invivoscribe – TCRB only</p> <p>In-house</p> <p>Using:</p> <p>Standard primer design methodology (including TCR) (SOP MH75)</p> <p>And</p> <p>PCR (including multiplex PCR) with Fluorescent fragment analysis using ABI 3500 Genetic Analyzer.</p> <p>Analysis using Genemapper and Chimermarker</p> <p>SOPs</p> <p>MH11 MH14 MH19 MH36 MH56</p>



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<p>HUMAN BODY FLUIDS/TISSUE (cont'd)</p> <p>Genomic DNA extracted in-house from the sample types listed above or received as primary sample type from an external source</p> <p>Genomic DNA extracted in-house from the sample types listed above or received as primary sample type from an external source</p>	<p><u>Genomics analysis for the purpose of clinical diagnosis, prognosis, and patient management</u> (cont'd)</p> <p>Fragments sizing for detection of known SNVs</p> <p>[Definitive list HMDS4407]</p> <p>Detection of known SNVs and small indels</p> <p>[Definitive list HMDS4407]</p>	<p>In house documented methods incorporating manufacturers' instructions where relevant</p> <p>Fluorescent based PCR amplification followed by restriction enzyme digest</p> <p>In house methodology with thermal cyclers and ABI 3500 Genetic Analyzer</p> <p>Analysis using Genemapper</p> <p>SOP: MH11</p> <p>Gel Electrophoresis based Fragment Analysis</p> <p>Using:</p> <p>Standard primer design methodology</p> <p>And</p> <p>PCR or allele specific PCR using Thermal cyclers</p> <p>Resolution with Agarose gel electrophoresis and visualisation using UVP UV transilluminator 312nm, Cannon Powershot digital camera capture system and software.</p> <p>SOP MH08 MH47 MH57 MH58</p>



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<p>HUMAN BODY FLUIDS/TISSUE (cont'd)</p> <p>Genomic DNA extracted in-house from the sample types listed above or received as primary sample type from an external source</p> <p>cDNA derived from RNA extracted and reverse transcribed in-house from blood and bone marrow, or received as primary sample type from an external source</p>	<p><u>Genomics analysis for the purpose of clinical diagnosis, prognosis, and patient management</u> (cont'd)</p> <p>Detection of whole exon deletions/duplications</p> <p>[Definitive list HMDS4408]</p> <p>Detection of fusion gene transcripts and determination of breakpoint regions</p> <p>[Definitive list HMDS4409]</p>	<p><b>In house documented methods incorporating manufacturers' instructions where relevant</b></p> <p><b>Fragment Analysis for Multiplex Ligation-dependent Probe Amplification (MLPA)</b></p> <p>Multiplex Ligation Probe Analysis (MLPA) using: Commercial commercial MRC Holland kits and Thermal cyclers and ABI 3500 Genetic Analyzer</p> <p>Analysis using Coffalyser Analysis Software</p> <p>SOPs MH81 MH82</p> <p><b>Qualitative Real Time PCR</b> In house methodology.</p> <p>Real time PCR using Applied Biosystems Realtime 7500 analyser</p> <p>Analysis using 7500 analyser software</p> <p>SOPs MH35 MH55</p>



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<p>HUMAN BODY FLUIDS/TISSUE (cont'd)</p> <p>cDNA received as primary sample type or derived from RNA extracted and reverse transcribed in-house from blood and bone marrow</p> <p>cDNA received as primary sample type or derived from RNA extracted and reverse transcribed in-house from blood and bone marrow</p> <p>cDNA received as primary sample type or derived from RNA extracted and reverse transcribed in-house from blood and bone marrow</p>	<p><u>Genomics analysis for the purpose of clinical diagnosis, prognosis, and patient management (cont'd)</u></p> <p>Quantification of major breakpoint regions in <i>BCR-ABL1</i> fusion transcripts</p> <p>[Definitive list: HMDS4409]</p> <p>Quantification of breakpoint regions of fusion transcripts</p> <p>[Definitive list HMDS4409]</p> <p>Quantification of SNVs and small indels</p> <p>[Definitive list HMDS4409]</p>	<p>In house documented methods incorporating manufacturers' instructions where relevant</p> <p><b>Quantitative Real Time PCR</b> In house real time PCR methodology with EAC primers and probes.</p> <p>Ipsogen Qiagen plasmids standard curve used for quantitative assay</p> <p>Quantification using Applied Biosystems Realtime 7500 analyser</p> <p>Analysis using 7500 analyser software</p> <p>SOPs</p> <p>MH35 MH55</p> <p>In house real time PCR methodology with EAC primers and probes.</p> <p>Ipsogen Qiagen plasmids standard curve used for quantitative assay</p> <p>Quantification using Applied Biosystems Realtime 7500 analyser and Quantstudio 3</p> <p>Analysis using analyser software</p> <p>SOPs: MH88</p> <p>Real time PCR using ipsogen NPM1 MutaQuant Kits</p> <p>Quantification using Applied Biosystems Realtime 7500 and analysis using 7500 software</p> <p>SOPs: MH88</p>



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<p><b>HUMAN BODY FLUIDS/TISSUE (cont'd)</b></p> <p>Genomic DNA extracted in-house from the sample types listed above or received as primary sample type from an external source</p>	<p><u>Genomics analysis for the purpose of clinical diagnosis, prognosis, and patient management (cont'd)</u></p> <p>Screening of large targeted single or multigene panels for genetic variants</p> <p>[Definitive list HMDS4410]</p> <p>SNVs and small indels</p>	<p>In house documented methods incorporating manufacturers' instructions where relevant</p> <p><b>Next Generation Sequencing</b></p> <p>Targeted amplicon library preparation</p> <p>Fluidigm 48:48 access array with High Throughput Sequencing</p> <p>Using</p> <p>Thermal cyclers and Illumina MiSeq</p> <p>Analysis using in-house Bioinformatics pipeline</p> <p>SOPs</p> <p>MH78 MH83 BI001 BI002 BI003 BI004</p>
<p>Genomic DNA extracted in-house from the sample types listed above or received as primary sample type from an external source</p>	<p>Screening of large targeted single or multigene panels for genetic variants</p> <p>[Definitive list HMDS4410]</p> <p>SNVs and small indels</p>	<p>Library Preparation using Twist enzymatic fragmentation and Custom Twist probes</p> <p>Using</p> <p>Thernak cyclers and NextSeq 550 DX</p> <p>Analysis and interpretation using in-house Bioinformatics pipeline and HaemOncDB v 3 respectively</p> <p>SOPs:</p> <p>MH87 BI010 BI011 BI012 YNEGLH004 YNEGLH002</p>





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<p><b>HUMAN BODY FLUIDS/TISSUE (cont'd)</b></p> <p>Paraffin embedded tissue Bone marrow smears Peripheral Blood smears Tissue imprint/Dab Methanol:acetic acid fixed cell suspensions Ammonium chloride lysed BM and PB CD138+ plasma cell selections</p> <p>FFPE Bone marrow smears, MAA fixed haematologically-derived cell suspensions</p> <p>FFPE Samples</p> <p>FFPE</p>	<p><u>Genomics analysis for the purpose of clinical diagnosis, prognosis, and patient management (cont'd)</u></p> <p>Detection and analysis of genomic rearrangements and imbalances in:</p> <p>Haemato-Oncological disorders Bone Marrow failure Identification/Confirmation of genomic rearrangements, gains and losses – using</p> <p>Locus specific probes:</p> <p>Break apart Dual colour dual fusion Copy Number/Amplification</p> <p>ALK Breakapart IGH/MAFB</p> <p>[Definitive list of probes HMDS4504]</p> <p>Detection of EBV infection in B-Cell lymphoproliferative disorders</p>	<p>In house documented methods incorporating manufacturers' instructions where relevant</p> <p><b>Fluorescent in-situ hybridisation (FISH)</b></p> <p>Separation of cell fractions using Miltenyi Automacs Pro</p> <p>Manual FISH: FISH performed using commercial probes, CytoBrite Slide Incubation System, MicroFISH Hybridization Oven Grant HotPlate, Clifton waterbaths, microwave, pressure cooker, microfuge and pipettes.</p> <p>Automated FISH: FISH performed using Dako OMNIS automated platform using commercial probes.</p> <p>Analysis using: Cytovision analysis software and and Leica fluorescent microscope system</p> <p>SOPs F001 F002 F005 FC028</p> <p><b>In-situ hybridisation</b></p> <p>ISH performed using Dako EBV (EBER) probe (with Dako PNA ISH Detection Kit</p> <p>CytoBrite Slide Incubation System, MicroFISH Hybridization Oven Grant HotPlate, Clifton waterbaths, microwave, microfuge and pipettes.</p> <p>Analysis using Light Microscopy</p> <p>SOP F004</p>



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HUMAN BODY FLUIDS/TISSUE (cont'd)	<u>Genomics analysis for the purpose of clinical diagnosis, prognosis, and patient management</u> (cont'd)	In house documented methods incorporating manufacturers' instructions where relevant
SNP array data files received from an external source within the YNEGLH	Detection of DNA copy number variation and loss of heterozygosity.	<b><u>SNP Array – Data analysis only:</u></b>  Analysis and interpretation of SNP array data using BlueFuse Multi and web-based UCSC genome browser.  SOP MH66
HUMAN BODY TISSUE AND FLUIDS	<u>Haematology examinations for the purpose of haematological malignancy diagnosis</u>	Documented in house methods incorporating manufacturers' instructions where relevant:
Blood, bone marrow aspirate, CSF, Effusions	Morphological assessment of haematopoietic cells	Automated May Grunwald Giemsa staining using Leica ST5020 Light microscopy SOP GEN004
Blood, bone marrow aspirate, CSF, Effusions	Full blood count: White blood cell count (WBC), red cell count (RBC), Haemoglobin (HGB), Haematocrit (HCT), Mean cell volume (MCV), Mean cell haemoglobin (MCH), Mean cell haemoglobin concentration (MCHC), Lymphocyte percentage/ count (LYM%/LYM#), Neutrophil percentage/count (NEUT%/#) and Mixed population percentage/Count (MXD%/#)	Sysmex XP-300 blood analyser, SOP GEN001, GEN 004, GEN005 GEN011, GEN016, GEN021
Blood	Blood film - cell typing and morphology, manual White Blood Cell Differential for enumeration of cell types	Thermo Scientific cytospin 4, Leica stainer using May Grunwald Giemsa Stain Light microscopy SOP GEN001, GEN 004, GEN005 GEN011, GEN016, GEN021



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<p><b>HUMAN BODY TISSUE AND FLUIDS (cont'd)</b></p> <p>Blood, bone marrow aspirate CSF, effusions, histological tissue</p>	<p><u>Haematology examinations for the purpose of haematological malignancy diagnosis (cont'd)</u></p> <p>Immunophenotyping of the following markers/antigens to screen for, diagnose or monitor: Acute leukaemia - AML/B-ALL/T-ALL Chronic myeloid disorders - MDS/CML/CMML B- &amp; T- lymphoproliferative disorders Myeloma/MGUS</p> <p>Stem Cell Screen (SCS): CD64 CD45 CD38 CD10 CD19 CD117 CD34 HLA-DR</p> <p>B-cell screen (BLS): CD19 CD20 CD5 CD10 CD305 (LAIR-1) CD45 Kappa Lambda</p> <p>T-cell screen (TLS): CD3 CD4 CD8 CD16/CD56 CD45 CD7 CD5 HLA-DR</p>	<p>Documented in house methods incorporating manufacturers' instructions where relevant:</p> <p>Flow cytometry on Becton Dickinson FACS Canto II using defined monoclonal antibody panels SOP FC001 - 003, FC022-025, FC027, GEN006 SOP FC004-011, FC026</p> <p>8 colours</p> <p>8 colours</p> <p>8 colours</p>



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<p><b>HUMAN BODY TISSUE AND FLUIDS (cont'd)</b></p> <p>Blood, bone marrow aspirate CSF, effusions, histological tissue (cont'd)</p>	<p><u>Haematology examinations for the purpose of haematological malignancy diagnosis (cont'd)</u></p> <p>Immunophenotyping of the following markers/antigens to screen for, diagnose or monitor: Acute leukaemia - AML/B-ALL/T-ALL Chronic myeloid disorders - MDS/CML/CMML B- &amp; T- lymphoproliferative disorders Myeloma/MGUS (cont'd)</p> <p>Plasma cell screen: CD38 CD138 CD19 CD45 CD56 CD117 CD27 CD81 Outreach monitoring (CMP): CD19 CD38 CD45 CD5 Kappa Lambda Rituximab monitoring: CD3 CD14 CD19 CD38 CD27 CD45</p>	<p>Documented in house methods incorporating manufacturers' instructions where relevant:</p> <p>Flow cytometry on Becton Dickinson FACS Canto II using defined monoclonal antibody panels SOP FC001 - 003, FC022-025, FC027, GEN006 SOP FC004-011, FC026</p> <p>8 colours</p> <p>6 colours</p> <p>6 colours</p>



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<p><b>HUMAN BODY TISSUE AND FLUIDS (cont'd)</b></p> <p>Blood, bone marrow aspirate CSF, effusions, histological tissue</p>	<p><u>Haematology examinations for the purpose of haematological malignancy diagnosis (cont'd)</u></p> <p>Immunophenotyping of the following markers/antigens in an 8-colour panel to diagnose and monitor myeloid disorders (AML, MDS, CMML)</p> <p>TdT MPO CD33 CD117 CD34 CD45 CD79a CD3 CD38 CD56 CD13 CD7 CD11b CD10 CD16 CD15 CD300e CD14 CD64 HLA-DR CD36 CD235a CD105 CD71 CD61 CD2 NG2 CD123 CD25 CD4</p>	<p>Documented in house methods incorporating manufacturers' instructions where relevant:</p> <p>Flow cytometry performed on Becton Dickinson FACS Lyric instrument using defined monoclonal antibody panels SOP FC022-025 &amp; 027, GEN006 SOP FC013</p>



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<p><b>HUMAN BODY TISSUE AND FLUIDS (cont'd)</b></p> <p>Blood, bone marrow aspirate CSF, effusions, histological tissue (cont'd)</p>	<p><u>Haematology examinations for the purpose of haematological malignancy diagnosis (cont'd)</u></p> <p>Immunophenotyping of the following markers/antigens to diagnose and monitor Acute lymphoblastic leukaemia (B-ALL &amp; T-ALL)</p> <p>B-ALL: TdT MPO CD33 CD117 CD34 CD45 CD79a CD3 CD66c CD123 CD304 CD73 CD81 CD13 CD33 CD34 CD19 CD10 CD45 CD38 CD79b CD86 CD9 CD58 CD24 CD44 CD22 HLA-DR NG2 CD20 CD15</p>	<p>Documented in house methods incorporating manufacturers' instructions where relevant:</p> <p>Flow cytometry performed on FACS Lyric instrument using defined monoclonal antibody panels SOP FC022-025, 027, GEN006 SOP FC015 &amp; FC017</p> <p>8 colours</p>



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HUMAN BODY TISSUE AND FLUIDS (cont'd)  Blood, bone marrow aspirate CSF, effusions, histological tissue (cont'd)	<u>Haematology examinations for the purpose of haematological malignancy diagnosis</u> (cont'd)  Immunophenotyping of the following markers/antigens to diagnose and monitor Acute lymphoblastic leukaemia (B-ALL & T-ALL) (cont'd)  T-ALL: CD3 CD1a CD2 CD4 CD5 CD7 CD8 CD10 CD16 CD25 CD27 CD45 CD45RA CD56 CD57 HLA-DR TCR $\alpha\beta$ TCR $\gamma\delta$ Tdt MPO CD79a CD33 CD117 CD34	Documented in house methods incorporating manufacturers' instructions where relevant:  Flow cytometry performed on Facs Lyric instrument using defined monoclonal antibody panels SOP FC022-025, 027, GEN006 SOP FC015 & FC017  6 colours



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<p>HUMAN BODY TISSUE AND FLUIDS (cont'd)</p> <p>Blood, bone marrow aspirate CSF, effusions, histological tissue (cont'd)</p>	<p><u>Haematology examinations for the purpose of haematological malignancy diagnosis (cont'd)</u></p> <p>Immunophenotyping of the following markers/antigens to diagnose and monitor mature B- and T-cell lymphoproliferative disorders:</p> <p>Extended B-cell panel: CD19 CD20 CD23 CD43 CD81 CD79b CD5 ROR1 CD95 CD31 CD49d CD305 (LAIR-1) CD38 CD10 CD25 CD11c CD103 CD200 CD39 CD22 CD196 CD185 IgG IgD IgM CD27</p>	<p>Documented in house methods incorporating manufacturers' instructions where relevant:</p> <p>Flow cytometry on Becton Dickinson FACS Canto II using defined monoclonal antibody panels SOP FC001 - 003, FC022-025, FC027, GEN006 FC011, FC019</p> <p>8 colours</p>





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<p>HUMAN BODY TISSUE AND FLUIDS (cont'd)</p> <p>Blood, bone marrow aspirate CSF, effusions, histological tissue (cont'd)</p>	<p><u>Haematology examinations for the purpose of haematological malignancy diagnosis (cont'd)</u></p> <p>Immunophenotyping of the following markers/antigens to diagnose and monitor mature B- and T-cell lymphoproliferative disorders:</p> <p>Extended T-cell panel:</p> <p>CD3 CD1a CD2 CD4 CD5 CD7 CD8 CD10 CD16 CD25 CD27 CD45 CD45RA CD52 CD56 CD57 HLA-DR TCR<math>\alpha\beta</math> TCR<math>\gamma\delta</math></p> <p>CD14</p>	<p>Documented in house methods incorporating manufacturers' instructions where relevant:</p> <p>Flow cytometry on Becton Dickinson FACS Canto II using defined monoclonal antibody panels SOP FC001 - 003, FC022-025, FC027, GEN006 FC011, FC019</p> <p>6 colours</p>



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Blood	Immunophenotyping of the following markers/antigens to assess the deficiency of GPI linked antigens on red blood cells and leucocytes (PNH panel)	Flow cytometry performed on Becton Dickinson FACS Lyric instrument using defined monoclonal antibody panels SOP FC020 SOP FC032
	Red blood cells: CD235a CD59 CD71 CD3d Leucocytes: CD16 CD15 CD14 CD24 CD33 Flaer Alexa-488 CD157	4 Colours  8 Colours
Blood/Bone Marrow/CSF	Identification of normal or aberrant PML protein expression pattern in suspected cases of Acute Promyelocytic Leukaemia	Immunofluorescence; Cytovision analysis software and Leica fluorescent microscope system SOP IMM006
Histology and haematology slides prepared as above	Morphological assessment and interpretation/diagnosis	Microscopy SOP R1, OFF002
END		