


Schedule of Accreditation

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United Kingdom Accreditation Service

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

 <p>9092</p> <p>Accredited to ISO 15189:2022</p>	<p align="center">Synnovis Analytics LLP</p> <p align="center">Issue No: 020 Issue date: 16 December 2025</p> <table border="1"> <tr> <td data-bbox="403 443 903 712"> <p>Department of Haematological Medicine 2nd Floor Friars Bridge Court 41-43 Blackfriars Road London SE1 8NZ United Kingdom</p> </td><td data-bbox="903 443 1479 712"> <p>Contact: Nura Ibrahim Tel: +44 (0) 203 299 7685 E-Mail: nura.ibrahim@synnovis.co.uk Website: www.synnovis.co.uk</p> </td></tr> </table>	<p>Department of Haematological Medicine 2nd Floor Friars Bridge Court 41-43 Blackfriars Road London SE1 8NZ United Kingdom</p>	<p>Contact: Nura Ibrahim Tel: +44 (0) 203 299 7685 E-Mail: nura.ibrahim@synnovis.co.uk Website: www.synnovis.co.uk</p>
<p>Department of Haematological Medicine 2nd Floor Friars Bridge Court 41-43 Blackfriars Road London SE1 8NZ United Kingdom</p>	<p>Contact: Nura Ibrahim Tel: +44 (0) 203 299 7685 E-Mail: nura.ibrahim@synnovis.co.uk Website: www.synnovis.co.uk</p>		
<p align="center">Testing performed by the Organisation at the locations specified below</p>			

Locations covered by the organisation and their relevant activities

Laboratory locations:

Location details	Activity	Location code
<p>Address Department of Haematological Medicine Ground Floor, Bessemer Wing King's College Hospital Denmark Hill London SE5 9RS</p> <p>Local contact Nura Ibrahim (as above)</p>	<p>Haematology Molecular Haematology Haemoglobinopathies Blood Transfusion Coagulation</p>	KCH
<p>Blackfriars Hub Laboratory 2nd Floor Friars Bridge Court 41 Blackfriars Road London SE1 8NZ</p> <p>Nura Ibrahim (as above)</p>	<p>Blood Sciences Laboratory: Haematology</p>	HUB
<p>Pathology Department South Wing, 1st Floor Princess Royal University Hospital Orpington Kent BR6 8ND</p> <p>Nura Ibrahim (as above)</p>	<p>Blood Sciences Laboratory: Haematology Blood Transfusion Storage & issue of blood & blood products</p>	PRUH
<p>Pathology Department Royal Brompton Hospital Sydney Street London SW3 6NP</p> <p>Blood Sciences and Blood Transfusion</p> <p>Luke Woodford Luke.Woodford@synnovi s.co.uk</p>	<p>Haematology Coagulation Blood Transfusion Blood Transfusion Issue Fridges located within laboratory, Adult ICU, theatres and paediatric ICU Issue of Blood and Blood Products</p>	RBH
<p>Pathology Building Harefield Hospital Hill End Road Harefield UB9 6JH</p> <p>Blood Sciences and Blood Transfusion</p> <p>Jey Visuvanathan J.Visuvanathan@synnovi s.co.uk</p>	<p>Haematology Coagulation Blood Transfusion Blood Transfusion Issue Fridges located within laboratory and ITU/ theatres Issue of Blood and Blood Products</p>	HH



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Site activities performed away from the locations listed above:

Location details	Activity
<p>King's College Hospital (Denmark Hill):</p> <ul style="list-style-type: none">• Emergency Department• Liver Theatre• Main Theatres• Nightingale Birth Centre (NBC)• Liver Unit ITU (LITU)• Centenary Wing• Harris Birthright (Windsor Walk)• Critical Care Unit• Guthrie Ward (X-ray) <p>King's College Hospital (Orpington Site):</p> <ul style="list-style-type: none">• Orthopaedic Theatres	<p>Storage & issue of blood & blood products only</p>



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DETAIL OF ACCREDITATION

Materials/Products tested	Type of test/Properties measured/Range of measurement	Standard specifications/ Equipment/Techniques used	Location code
<p>HUMAN TISSUES AND FLUIDS (cont'd)</p> <p>Amniotic Fluid Chorionic Villus Sample Cultured Amniocytes Dried Blood Spot Foetal Blood Formalin-Fixed Paraffin-Embedded (FFPE) samples Plasma Saliva Serum Whole blood</p> <p>Genomic DNA extracted in house from whole blood (unless stated otherwise for specified tests) or received as primary sample type</p>	<p><u>Molecular haematology examinations for the purpose of clinical diagnosis</u></p>	<p>In house documented methods incorporating manufacturers' instructions as required</p> <p>DNA Extraction, manual methods and automated using:</p> <p>QIAAsymphony (blood) LP-HAE-PND105</p> <p>Qiagen EZ1 LP-HAE-PND076 (foetal samples) LP-HAE-PND116 (saliva)</p> <p>PCR amplification using any of (unless stated otherwise for specified tests): Applied Biosystems Veriti™ Thermal Cycler and: Veriti™ Thermal Cycler User Guide (PN 4375799) Rotor-Gene Q and: Rotor-Gene Q Operator manual 1.7.87 Eppendorf Mastercycler Nexus and: Eppendorf Mastercycler Nexus operating manual (2012)</p>	KCH
<p>and DNA extracted from Saliva</p>	<p>Thrombophilia genetic analysis:</p> <p>Detection of three genetic variants affecting thrombophilia: Factor V Leiden Prothrombin C Methylene Tetrahydrofolate Reductase</p> <p>Hereditary Haemochromatosis genetic analysis: Detection of the two common genetic variants in the HFE gene associated with hereditary haemochromatosis: HFE p.C282Y HFE p.H63D</p>	<p>RT PCR Allelic discrimination TaqMan assay LP-HAE-PND110</p> <p>RT PCR Allelic discrimination TaqMan assay LP-HAE-PND104</p>	KCH



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<p>HUMAN TISSUES AND FLUIDS (cont'd)</p> <p>Genomic DNA extracted in house from whole blood (unless stated otherwise for specified tests) or received as primary sample type (cont'd)</p>	<p><u>Molecular haematology examinations for the purpose of clinical diagnosis</u> (cont'd)</p> <p>Detection of large deletions and duplications in the HFE gene</p> <p>Alpha-1-antitrypsin deficiency genotyping:</p> <p>Detection of two genetic variants in the SERPINA1 gene which are known to cause Alpha-1-antitrypsin deficiency: Z and S mutations</p> <p>Gilbert's syndrome genetic analysis:</p> <p>Homozygosity for the 7 repeat allele of the dinucleotide repeat region in the promoter of the UGT1A gene</p>	<p>In house documented methods incorporating manufacturers' instructions as required</p> <p>Multiplex ligation-dependent probe amplification (MLPA) by PCR using: Eppendorf Mastercycler Nexus or Applied Biosystems Veriti™ Applied Biosystems 3500XL Genetic Analyser LP-HAE-PND117</p> <p>RT PCR machine allelic discrimination TaqMan assay LP-HAE-PND111</p> <p>Applied Biosystems 3500XL Genetic Analyser LP-HAE-PND071</p>	KCH



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HUMAN TISSUES AND FLUIDS (cont'd)	<u>Molecular haematology examinations for the purpose of clinical diagnosis</u> (cont'd)	In house documented methods incorporating manufacturers' instructions as required	KCH
Genomic DNA extracted in house from whole blood (unless stated otherwise for specified tests) or received as primary sample type (cont'd)			
and DNA extracted from CVS	Pyruvate Kinase deficiency genetic analysis:		
	Detection of large deletions and duplications in the PKLR gene	MLPA by PCR using: Eppendorf Mastercycler Nexus or Applied Biosystems Veriti™ Thermal Cycler- Veriti™ Applied Biosystems 3500XL Genetic Analyser LP-HAE-PND117	KCH
and DNA extracted from: Saliva Dried Blood Spot Amniotic Fluid CVS Cultured Amniocytes Foetal Blood	Haemoglobin variant genetic analysis:		
	Detect of the most common HBB gene mutation known to cause Sick cell disease: HbS and the haemoglobin variant: HbC	RT PCR Allelic discrimination TaqMan assay using: Rotor-Gene Q LP-HAE-PND039	KCH
	Confirmation of HbS and HbC variants	Restriction fragment length polymorphism (RFLP) assay by PCR using: Eppendorf Mastercycler Nexus or Applied Biosystems Veriti™ and Agarose gel electrophoresis LP-HAE-PND013	KCH
	Confirmation of the HbS mutation	Sanger Sequencing using: Applied Biosystems 3500XL Genetic Analyser LP-HAE-PND039	KCH



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<p>HUMAN TISSUES AND FLUIDS (cont'd)</p> <p>Genomic DNA extracted in house from whole blood (unless stated otherwise for specified tests) or received as primary sample type (cont'd)</p> <p>and DNA extracted from: Saliva</p>	<p><u>Molecular haematology examinations for the purpose of clinical diagnosis</u> (cont'd)</p> <p>Detection of large deletions and duplications in the HBB, HBA1 and HBA2 genes</p> <p>Thalassaemia syndromes genetic analysis:</p>	<p>In house documented methods incorporating manufacturers' instructions as required</p> <p>MLPA by PCR and: Applied Biosystems 3500XL Genetic Analyser LP-HAE-PND117</p>	KCH
<p>Dried blood spots Amniotic fluid CVS Cultured amniocytes Fetal blood</p>	<p>Detection of common deletions in the HBA1 and HBA2 genes known to cause Alpha Thalassaemia</p> <p>Detection of single base changes and microlesions in the HBB, HBA1 and HBA2 genes that cause Thalassaemia syndromes</p> <p>Detection of large deletions and duplications in the alpha globin loci on chromosome 16 and the beta globin loci on chromosome 11</p>	<p>Gap PCR using: Eppendorf Mastercycler Nexus or Applied Biosystems Veriti™ Thermal Cycler and Agarose gel electrophoresis LP-HAE-PND052</p> <p>Sanger Sequencing using: Applied Biosystems 3500XL Genetic Analyser LP-HAE-PND118</p> <p>MLPA by PCR using: Eppendorf Mastercycler Nexus or Applied Biosystems Veriti™ Thermal Cycler Applied Biosystems 3500XL Genetic Analyser LP-HAE-PND117</p>	



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<p>HUMAN TISSUES AND FLUIDS (cont'd)</p> <p>Genomic DNA extracted in house from whole blood (unless stated otherwise for specified tests) or received as primary sample type (cont'd)</p> <p>and DNA extracted from: Whole blood Saliva Dried Blood Spot</p>	<p><u>Molecular haematology examinations for the purpose of clinical diagnosis</u> (cont'd)</p> <p>Hereditary persistence of fetal haemoglobin (HPFH) genetic analysis:</p> <p>Detection of the most common deletional forms of HPFH caused by large deletions in the beta globin loci</p> <p>Confirmation of the deletional forms of HPFH</p> <p>Detection of single base changes and microlesions in the promoter regions of the A gamma (HBG1) and the G gamma (HBG2) globin genes to that cause non-deletional HPFH</p> <p>Maternal cell contamination:</p>	<p>In house documented methods incorporating manufacturers' instructions as required</p> <p>MLPA by PCR using: Eppendorf Mastercycler Nexus or Applied Biosystems Veriti™ Thermal Cycler Applied Biosystems 3500XL Genetic Analyser LP-HAE-PND072</p> <p>Gap PCR Eppendorf Mastercycler Nexus or Applied Biosystems Veriti™ Thermal Cycler and Agarose gel electrophoresis LP-HAE-PND072</p> <p>Sanger Sequencing using: Applied Biosystems 3500XL Genetic Analyser LP-HAE-PND118</p>	KCH
<p>and DNA extracted from: Whole blood Amniotic Fluid CVS Cultured Amniocytes Dried Blood Spot Fetal Blood Formalin-Fixed Paraffin-Embedded (FFPE) samples Plasma Serum</p>	<p>Analysis of repeat length of 16 microsatellite markers</p>	<p>ABI AmpFLSTR kit multiplexed PCR reaction. Products analysed using Applied Biosystems 3500XL Genetic Analyser LP-HAE-PND067</p>	KCH



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<p>HUMAN TISSUES AND FLUIDS (cont'd)</p> <p>Genomic DNA extracted in house from whole blood (unless stated otherwise for specified tests) or received as primary sample type (cont'd)</p>	<p><u>Molecular haematology examinations for the purpose of clinical diagnosis</u> (cont'd)</p> <p>Red blood cell disorders and mutations:</p> <p>Membranopathy Red Cell Enzyme Haemoglobinopathies Congenital Erythrocytosis Diamond-Blackfan Anaemia Bone Marrow Failure Thrombocytopenia Neutropenia Porphyria Siderblastic Anaemia Iron Regulation HLH Lymphedema Single Genes (HP, Serpina1, UGT1A1) Sex Chromosome Markers</p> <p>Validation of variants from NGS</p>	<p>In house documented methods incorporating manufacturers' instructions as required</p> <p>Beckman Coulter Biomek i7 robot to construct sequencing libraries followed by hybridisation to probe targets and amplification. Tapestation 4150 Amplified library sequenced by Next Generation Sequencing (NGS) on Illumina MiSeq. Data analysed using a NIPY pipeline LP-HAE-PND132</p> <p>Sanger Sequencing using: Applied Biosystems 3500XL Genetic Analyser LP-HAE-PND131</p>	<p>KCH</p> <p>KCH</p>



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HUMAN BODY FLUIDS (cont'd)	<u>Haematology examinations for the purpose of clinical diagnosis</u>	In house documented methods incorporating manufacturers' instructions as required	
Blood	Full Blood Count (FBC): White cell count WBC Red cell count RBC Haemoglobin HGB Haematocrit (HCT) MCV MCH MCHC RDW-CV Platelet count (PLT) MPV #Neutrophils #Lymphocytes #Monocytes #Eosinophils #Basophils #NRBC Metamyelocyte count Myelocyte count Promyelocyte count Blasts count Blast Flag Fragments Flag Immature Granulocyte Flag Platelet Clump Flag Atypical Lymphocyte Flag	Sysmex XN-9100 series – XN10 Optical Analysis (Flow Cytometry) LP-BS-HA001 LP-BS-HA005 LP-BS-HA008 BSL-HUB-HAEM-SOP1 BSL-HUB-HAEM-SOP6	KCH HUB
Whole Blood	Full Blood Count (FBC) Analysis: Total red cell count Haemoglobin Mean cell volume Mean cell haemoglobin Mean cell haemoglobin concentration Haematocrit Red cell distribution width Total white cell count Absolute neutrophil count Absolute lymphocyte count Absolute monocyte count Absolute eosinophil count Absolute basophil count Platelet count Mean platelet volume	Sysmex XN10 & XN20 Fluorescence Flow Cytometry BH-AP-040	PRUH



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HUMAN BODY FLUIDS (cont'd)	<u>Haematology examinations for the purpose of clinical diagnosis</u> (cont'd)	In house documented methods incorporating manufacturers' instructions as required	
Whole Blood	Blood Full Blood Count (PCV, MCV, Hb, RBC, WBC, Hct, MCV, MCH, MCHC, RDW Platelets, Neutrophils, lymphocytes, monocytes, Eosinophils, Basophils Nucleated RBCs)	Beckman Coulter DxH 800 analysers using impedance principle for cell counting and sizing; volume, conductivity and light scatter (VCSn) for differential leucocyte count and measurement of absorbance of stable pigment for haemoglobin concentration. SOPHA00494	RBH HH
Blood	Reticulocytes: Retic absolute Retic Percentage LFR MFR HFR Retic RBC IRF %HYPO Ret-He	Sysmex XN-9100 series – XN10 Optical Analysis (Flow Cytometry) LP-BS-HA001 LP-BS-HA005 BSL-HUB-HAEM-SOP1 BSL-HUB-HAEM-SOP6	KCH HUB
Whole Blood	Reticulocytes	Sysmex XN10 & XN20 Fluorescence Flow Cytometry BH-AP-040	PRUH
Blood	Citrate platelet count	Sysmex XN-9100 series – XN10 Optical Analysis (Flow cytometry) BSL-HUB-HAEM-SOP1 BSL-HUB-HAEM-SOP6	HUB
Whole Blood	Nucleated Red Cell Count	Sysmex XN10 & XN20 Fluorescence Flow Cytometry BH-AP-040	PRUH



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HUMAN BODY FLUIDS (cont'd)	<u>Haematology examinations for the purpose of clinical diagnosis</u> (cont'd)	In house documented methods incorporating manufacturers' instructions as required	
Whole blood	Nucleated RBCs and Reticulocyte Count	Beckman Coulter DxH 800 using New Methylene Blue stain – SOPHA00494	RBH
Blood	Detection and speciation of Malaria parasites: <i>Plasmodium falciparum</i> , <i>P. vivax</i> , <i>P. ovale</i> , <i>P. malariae</i> , <i>P. knowlesi</i> , t trypanosomes, microfilariae, <i>Leishmania</i> spp.	Wright-Giemsa & Giemsa R66 staining followed by Light Microscopy LP-BS-ML008, LP-BS-ML009	KCH
Whole Blood (Blood Film)	Detection & presumptive speciation of Malaria & non-malaria parasites, including: <i>P. falciparum</i> , <i>P. vivax</i> , <i>P. ovale</i> , <i>P. malariae</i> , <i>P. knowlesi</i> % Parasitaemia for <i>P. falciparum</i> & <i>P. knowlesi</i>	Thick and thin film Giemsa (pH7.2) staining BH-AP-002 Nikon Eclipse 80i Olympus BX43 Microscopy BH-AP-10 BH-AP-30	PRUH
	<i>P. falciparum</i> -specific HRP2 antigen Pan-malarial antigen of: <i>P. falciparum</i> , <i>P. vivax</i> , <i>P. ovale</i> , <i>P. malariae</i>	BINAX NOW ICT Immunochromatographic assay LP-BS-ML009	KCH
Whole Blood	Detection of Malaria antigens: <i>Plasmodium falciparum</i> HRP2 & <i>P. falciparum</i> , <i>P. vivax</i> , <i>P. ovale</i> , <i>P. malariae</i> Pan LDH	CareUS Immunochromogenic method BH-AP-002	PRUH
Blood (Thin blood films)	Detection of <i>Plasmodium</i> spp.	Pathway Diagnostics Quantitative Buffy Coat (QBC) kit & Fluorescence microscopy LP-BS-ML009	KCH



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HUMAN BODY FLUIDS (cont'd)	<u>Haematology examinations for the purpose of clinical diagnosis</u> (cont'd)	In house documented methods incorporating manufacturers' instructions as required	
Blood	Morphological analysis & differential cell count	Light Microscopy LP-BS-ML007, LP-BS-ML018	KCH
Blood	Erythrocyte Sedimentation Rate (ESR)	Sysmex Starrsed Interrliner XN FRL Westergren method LP-BS-HA002	KCH
		Sysmex Starrsed Interliner XN9100 BSL-HUB-HAEM-SOP6	HUB
Whole Blood	Erythrocyte Sedimentation Rate (ESR)	StaRRsed Autocompact Red cell sedimentation BH-AP-011	PRUH
Blood	Infectious Mononucleosis screen: Detection of the heterophile antibody	Accusay Mono Solid-phase immunoassay and: LP-BS-ML003	KCH
Plasma	Prothrombin/ INR* Activated partial thromboplastin time/APTT* Clauss Fibrinogen* Thrombin Time /TT*	Sysmex CN6500 (tests marked with * for urgent workflow) LP-BS-HT0013 Sysmex CS-5100 Clotting method LP-BS-HT001 LP-BS-HT003 LP-BS-HT004	KCH
Whole Blood	Clauss Fibrinogen	Sysmex CS-2500 Spectrophotometry BH-CP-002	PRUH
Plasma	Prothrombin time/PT Activated partial thromboplastin time/APTT APTT ratio Fibrinogen D-Dimer INR INR and APTT 50/50 correction Reptilase time	Sysmex CN6000 BSL-ALL-HT-SOP1	HUB



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Plasma (citratated)	<u>General Coagulation tests:</u> Prothrombin time (PT) and International Normalised Ratio (INR) Activated partial thromboplastin time (APTT) Fibrinogen assay Thrombin time (TT) D-dimer assay Anti-Xa assay Antithrombin assay PT mix APTT mix	Instrumentation Laboratory Werfen ACL TOP 550 coagulometers. Measurement principles include turbidometric clot endpoint detection, chromogenic assays and immunoturbidometric assays. SOPHA00429 Routine use of ACL TOP 550 SOPHA00037 Prothrombin time (PT) and International Normalised Ration (INR) SOPHA00104 Activated partial thromboplastin time (APTT) SOPHA00296 Fibrinogen assay SOPHA00106 Thrombin time (TT) SOPHA00040 D-dimer assay samples (ELISA method) RBH SOPHA00092 Anti-Xa assay SOPHA00137 Antithrombin assay SOPHA00037 PT mix SOPHA00104 APTT mix	RBH HH
Plasma (citratated)	<u>Specialist Coagulation tests:</u> Protein C Free protein S DRV/T (lupus anticoagulant) APCR vWf antigen vWf functional assay (Ristocetin cofactor) Plasminogen assay Factor assays: II, V, VII, VIII, IX, X, XI, XII Anti-Factor Xa Assay for Direct Xa Inhibitors	Instrumentation Laboratory Werfen ACL TOP 550 coagulometers. Measurement principles include turbidometric clot endpoint detection, chromogenic assays and immunoturbidometric assays. SOPHA00429 Routine use of ACLTOP 550 coagulometer SOPHA00111 Protein C SOPHA00112 Free protein S SOPHA00122 DRV/T (lupus anticoagulant) SOPHA00042 APCR SOPHA00113 vWf antigen SOPHA00113 vWf functional assay (Ristocetin cofactor) SOPHA00229 Plasminogen assay SOPHA00129 (Factor Assays APTT) SOPHA00130 (Factor Assays PT) SOPHA00536 (Apixaban and Rivaroxaban assays) SOPHA00537 (Direct Thrombin Inhibitor assay (Dabigatran))	RBH



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HUMAN BODY FLUIDS (cont'd)	<u>Haematology examinations for the purpose of clinical diagnosis</u> (cont'd)	In house documented methods incorporating manufacturers' instructions as required	
Whole Blood	Clotting Screen: Prothrombin Time (PT) Activated Partial Thromboplastin Time (APTT)	Sysmex CS-2500 Spectrophotometry Correction studies (where required) BH-CP-002	PRUH
Plasma	Factor Assays: FII FV FVII FVIII FIX FX FXI FXII	Sysmex CN6500 LP-BS-HT0013 Sysmex CS-5100 Clotting method LP-BS-HT001 LP-BS-HT003 LP-BS-HT004	KCH
Plasma	LA Screen (DRVVT) Activated Protein C Resistance	Sysmex CS-5100 Clotting method LP-BS-HT005 LP-BS-HT006	KCH
Plasma	DDimer VWF Antigen Free Protein S Antigen	Sysmex CN6500 (tests marked with * for urgent workflow) LP-BS-HT0013 Sysmex CS-5100 Latex bead Immunoturbidimetric Assay LP-BS-HT003 LP-BS-HT006 LP-BS-HT008	KCH
Whole Blood	D-dimer	Sysmex CS-2500 Spectrophotometry BH-CP-002	PRUH
Plasma	Anti-Xa: Rivaroxaban Apixaban Edoxaban Dabigatran Antithrombin Activity Protein C Activity	Sysmex CS-5100 Chromogenic method LP-BS-HT007 LP-BS-HT005	KCH



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Blood	Haemoglobin H bodies identification	Supra-vital staining with brilliant Cresyl blue and microscopic examination LP-HAE-RC-HB009	KCH
Blood	Haemoglobin S and Haemoglobin F quantification	Bio-Rad Variant II HPLC analyser LP-HAE-RC-HB003	KCH
Blood	Investigation for Hereditary Spherocytosis	EMA dye binding to red blood cells and flow cytometry analysis on Becton Dickinson FACS Canto II LP-HAE-RC-NEH-007	KCH
Whole Blood (Blood Film)	Red Cell morphology White Cell morphology Platelet morphology White Cell Differential	RAL Stainer Methylene Blue and May-Grünwald stains Methanol Free, Chromatic Detection BH-AP-039 Nikon Eclipse 80i Olympus BX43 Microscopy BH-AP-10 BH-AP-30	PRUH
Blood film prepared from whole blood specimen	Blood Film Examination for Assessment for cell typing and morphology including white blood cell differential for enumeration of cell types.	DxH Slidemaker/Stainer (RBH) Shandon Varistain 12 (HH) SOPHA00503 Preparation & staining of blood films SOPHA00474 Microscope use & maintenance.	RBH HH
Urine	Iron stain for Urine hemosiderin in the diagnosing of intravascular haemolysis.	Manual light microscopy of urine smears using manual Iron (Perl's stain) SOPHA00330	RBH
Plasma	Plasma Free Haemoglobin	HemoCue plasma/low Hb photometer using conversion of haemoglobin to azidemethaemoglobin followed by absorbance measurement SOPHA00271	RBH HH



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Materials/Products tested	Type of test/Properties measured/Range of measurement	Standard specifications/ Equipment/Techniques used	Location code
HUMAN BODY FLUIDS (cont'd)	<u>Blood transfusion examinations for the purpose of clinical diagnosis</u>	In house documented methods incorporating manufacturers' instructions as required	
Blood	ABO / RhD Blood Group antigens: A, B & D	Automated using: Biorad Banjo ID reader Biorad IH1000 Biorad IH500 LP-HAE-BT2-001, BT-3-A	KCH
Blood	<u>Automated Blood Grouping</u> A, B, O and AB groups	BioRad IH500 using Biorad micro column cards SOPHA000538	RBH HH
	Rh groups, antigens:-C, c, D, E, e, Kell blood group system, antigens: K	SOPHA00172	
Blood	<u>Manual Blood Group/Phenotype</u> A, B, O and AB groups	Manual method using Biorad micro column cards (as back-up to automated method) SOPHA00176	RBH HH
	Rh groups, antigens:-C, c, D, E, e, Kell blood group system, antigens: K	SOPHA00172	
Blood	Red Cell Antibody detection	Manual method using Biorad micro column cards and 3 cell panel SOPHA00176	RBH HH
Blood	Red Cell Antibody Identification: Indirect antiglobulin test (IAT) panel Enzyme panel	Manual method using Biorad micro column cards SOPHA00172	RBH HH
Blood	/Crossmatching Indirect antiglobulin test (IAT) crossmatch Emergency issue	IAT crossmatch carried out by manual method using Biorad microcolumn cards. SOPHA00164 Emergency issue of blood cross site BSL-RBH-BT-SOP3 Crossmatching & red cell issue	RBH HH



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Synnovis Analytics LLP
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Materials/Products tested	Type of test/Properties measured/Range of measurement	Standard specifications/ Equipment/Techniques used	Location code
HUMAN BODY FLUIDS (cont'd)	<u>Blood transfusion examinations for the purpose of clinical diagnosis</u>	In house documented methods incorporating manufacturers' instructions as required	
Blood	Antibody Screen for: D, C, Cw, E, c, e, K, Fya, Fyb, Jka, Jkb, S, s, Lea, Leb, M, N & Lua	Automated using: Biorad Banjo ID reader Biorad IH1000 Biorad IH500 LP-HAE-BT2-001, BT3-A	KCH
Blood (Red cells, Plasma)	ABO / RhD typing by detection & identification of antigens: A Rh D Positive AB Rh D Positive B Rh D Positive O Rh D Positive A Rh D Negative AB Rh D Negative B Rh D Negative O Rh D Negative	Automated using: CAT BioRad IH500 & IH1000 BT-PP-041 Manual using: CAT BioRad ABO and reverse grouping cards and A1 & B reagent cells BT-PP-032	PRUH
Blood (Plasma)	Antibody screen for detection of antibodies: Rh – C, D, E, c, e, Cw K, k, Kpa Fya, Fyb M, N, S, s Jka, Jkb Lea, Leb Lua P, P1	Automated using: CAT BioRad IH500 & IH1000 SOP?? BSL-KDP-BT-SOP5 Manual using: CAT BioRad IAT technology, anti-IgG IAT gel cards and BioRad 3 cell antibody screening reagent cells BT-PP-002	PRUH
Blood (Plasma)	Antibody Identification of antibodies: Rh – C, D, E, c, e, Cw K, Kpa Fya, Fyb M, N, S, s Jka, Jkb Lea, Leb Lua P1	Automated using: CAT BioRad IH500 & IH1000 BT-PP-041 Manual using: CAT BioRad IAT technology, anti-IgG IAT gel cards BT-PP-002	PRUH



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HUMAN BODY FLUIDS (cont'd)	<u>Blood transfusion examinations for the purpose of clinical diagnosis</u> (cont'd)	In house documented methods incorporating manufacturers' instructions as required	
Blood	DAT (polyspecific, monospecific)	Automated using: Biorad Banjo ID reader Biorad IH1000 Biorad IH500 LP-HAE-BT2-001, BT2-05, BT3-D	KCH
Blood (Red cells)	Direct Antiglobulin Testing (DAT)	Automated using: CAT BioRad IH500 & IH1000 BT-PP-041 Manual using: CAT BioRad polyspecific & monospecific IgG and CD3d DAT cards BT-PP-033	PRUH
Blood	Direct antiglobulin test (DAT)	Manual using Poly specific and mono specific (anti-IgG and anti-C3d) Biorad micro column cards SOPHA00527	RBH HH
Blood	RHK phenotype antigens: D,C,c, E, e, K	Automated using: Biorad Banjo ID reader Biorad IH1000 Biorad IH500 LP-HAE-BT2-001, BT2-003	KCH
Blood (Red cells)	Rh/K phenotyping of antigens: C, c, E, e, K	Automated using: CAT BioRad IH500 & IH1000 BT-PP-041 Manual using: CAT BioRad Rh/K antigen typing cards BT-PP-006	PRUH
Blood	Non-RHK phenotype antigens: Jka, Jkb, Lea, Leb, S, s, M/N, Fya, Fyb	Automated (except Fya/Fyb) using: Biorad Banjo ID reader Biorad IH500, ID- Incubator 37 SI, ID- Centrifuge 12 S, DiaCent-12 LP-HAE-BT2-003, BT3-008	KCH
Blood	Non-RHK phenotype antigens: Fya, Fyb (Biorad automated IH500)	Biorad column agglutination technique Biorad Banjo ID reader LP-HAE-BT2-019-Automated Sample processing using IH-500 LP-HAE-BT2-003 Automated and Manual Cell Typing	KCH



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HUMAN BODY FLUIDS (cont'd)	<u>Blood transfusion examinations for the purpose of clinical diagnosis</u> (cont'd)	In house documented methods incorporating manufacturers' instructions as required	
Blood	Antibody Titres of: Anti-A Anti-B	Manual using: Biorad Banjo ID reader BioRad cards/gels, ID-Centrifuge 12, ID-Incubator 37I LP-HAE-BT5-033	KCH
Blood	Foetal maternal haemorrhage	Kleihauer acid elution test using Microscopy LP-HAE-BT3-J	KCH
Blood (Red cells)	Foetal Maternal Haemorrhage	Kleihauer Acid Elution & microscopy BT-PP-005	PRUH
Blood	Antibody identification of: D, C, Cw, E, c, e, K, Fya, Fyb, Jka, Jkb, S, s, Lea, Leb, M, N & Lua	Automated or manual using: Biorad Banjo ID reader Biorad IH500, ID-Incubator 37 SI, ID-Centrifuge 12 S LP-HAE-BT2-002, BT3-B	KCH
Blood	Baby ABO / RhD Blood Group antigens: A, B, AB & D	Automated using: Biorad Banjo ID reader Biorad IH1000 Biorad IH500 LP-HAE-BT3-A, BT2-001	KCH
Blood	ABO / RhD Blood Group antigens: A, B & D	Manual using: BioRad cards/gels, Biorad Banjo ID reader DiaCent 12 Centrifuge LP-HAE-BT4-008	KCH
Blood	DAT (polyspecific, monospecific)	Manual using: Biorad Banjo ID reader Biorad column agglutination technique LP-HAE-BT2-05	KCH
Blood	Antibody Screen for: D, C, Cw, E, c, e, K, Fya, Fyb, Jka, Jkb, S, s, Lea, Leb, M, N & Lua (manual)	Manual using: Biorad Banjo ID reader Biorad column agglutination technique LP-HAE-BT3-A	KCH
Blood	Non-RHK phenotype antigens: Jka, Jkb, Lea, Leb, S, s, M/N, Fya, Fyb (manual)	Manual using: Biorad Banjo ID reader Biorad column agglutination technique LP-HAE-BT2-003	KCH



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HUMAN BODY FLUIDS (cont'd)	<u>Blood transfusion examinations for the purpose of clinical diagnosis</u> (cont'd)	In house documented methods incorporating manufacturers' instructions as required	
Blood	RHK phenotype antigens: D,C,c, E, e, K (manual)	Manual using: Biorad Banjo ID reader Biorad column agglutination technique LP-HAE-BT2-003	KCH
Blood	Antibody Titres of: Anti-A, Anti-B (Biorad automated IH500)	Biorad column agglutination technique Biorad Banjo ID reader LP-HAE-BT2-019-Automated Sample processing using IH-500 LP-HAE-BT5-033 Titration Of ABO Antibodies	KCH
Blood	Baby ABO / RhD Blood Group antigens: A, B, AB & D (manual)	Manual using: Biorad Banjo ID reader Biorad column agglutination technique LP-HAE-BT3-A	KCH
Blood	Compatibility testing of patient plasma with donor cells (Manual)	Manual using: Biorad Banjo ID reader Biorad column agglutination technique LP-HAE-BT4-004	KCH
Blood	Compatibility testing of patient plasma with donor cells (Automated)	Biorad automated IH1000 & IH500 Biorad column agglutination technique Biorad Banjo ID reader LP-HAE-BT2-019-Automated Sample processing using IH-500 LP-HAE-BT4-004 Serological (Full) Crossmatching/Compatibility Testing	KCH
Blood (Red cells, Plasma)	Compatibility testing (Crossmatch)	Manual using: CAT BioRad Anti-IgG IAT gel cards BT-PP-003	PRUH
Blood (Red cells, Plasma)	Compatibility testing (Crossmatch)	Automated using: CAT BioRad IH500 & IH1000 SOP??BSL-KDP-BT-SOP5	PRUH
END			