

Schedule of Accreditation

issued by

United Kingdom Accreditation Service

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

| | | |
|---|---|--|
|  <p>Accredited to ISO 15189:2012</p> | Oxford University Hospitals NHS Foundation Trust | |
| | Issue No: 001 Issue date: 27 July 2016 | |
| | Department of Laboratory Haematology Level 4 John Radcliffe University Hospital Headley Way Headington Oxford OX3 9DU | Contact: Andrew Platt Tel: +44 (0) 1865 (8)57663 E-Mail: andrew.platt@ouh.nhs.uk Website: www.ouh.nhs.uk |
| Testing performed by the Organisation at the locations specified below | | |

Locations covered by the organisation and their relevant activities

Laboratory locations:

| Location details | Activity | Location code |
|--|--|---------------|
| Level 4 John Radcliffe University Hospital Headley Way Headington Oxford OX3 9DU Local contact: Andrew Platt (details as above) | <ul style="list-style-type: none"> • Routine haematology • Routine Coagulation • Blood Grouping and antibody screen • Blood & blood product issue • Immunophenotyping by flow analysis • Haemoglobinopathy analysis • Molecular haematology | JRH |
| Horton Hospital Oxford Road Banbury OX16 9AL Haematology: Mr A Nicklin (01923 229369) Blood Transfusion: Miss J Staves (01865 220334) | <ul style="list-style-type: none"> • Routine haematology • Routine Coagulation • Blood Grouping and antibody screen • Blood & blood product issue | HH |
| Churchill Hospital Laboratory Medicine Old Road Headington Oxford OX3 7LE Mr P Baker (01865 857096) | <ul style="list-style-type: none"> • Routine FBC analysis • Routine Coagulation • Blood & blood product issue | CHLM |
| Churchill Hospital Oxford Haemophilia and Thrombosis Centre Old Road Headington Oxford OX3 7LE Mr P Baker (01865 857096) | <ul style="list-style-type: none"> • Routine Coagulation • Factor Assay analysis • Haemostatic investigations • Thrombophilia investigations | OHTC |



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Site activities performed away from the locations listed above:

| Location details | Activity |
|---|---|
| Churchill Hospital Porters Lodge Old Road Headington Oxford OX3 7LE | Blood storage & issue Fridge managed by Haematology |
| Delivery Suite John Radcliffe Hospital Headley Way Oxford OX3 9DU | Blood storage & issue only Fridge managed by Haematology |
| West Wing Theatres, Level 1 John Radcliffe Hospital Headley Way Oxford OX3 9DU | Blood storage & issue only Fridge managed by Haematology |
| John Radcliffe Theatres John Radcliffe Hospital Headley Way Oxford OX3 9DU | Blood storage & issue only Fridge managed by Haematology |
| Cardio Thoracic Critical Care John Radcliffe Hospital Headley Way Oxford OX3 9DU | Blood storage & issue only Fridge managed by Haematology |
| Churchill Theatres Level 1 Old Road Headington Oxford OX3 7LE | Blood storage & issue only Fridge managed by Haematology |



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|--|---|
| Day Treatment Unit Level 1 Churchill Hospital Old Road Headington Oxford OX3 7LE | Blood storage & issue only Fridge managed by Haematology |
| Haematology Ward Level 1 Churchill Hospital Old Road Headington Oxford OX3 7LE | Blood storage & issue only Fridge managed by Haematology |
| Theatres Nuffield Orthopaedic Centre Windmill Road Headington Oxford OX3 7HE | Blood storage & issue only Fridge managed by Haematology |
| The Foscote Hospital 2 Foscote Rise Banbury OX16 9XP | Blood storage & issue only Fridge monitored by Haematology |
| Treatment Centre Oxford Road, Banbury OX16 7FG | Blood storage & issue only Fridge monitored by Haematology |



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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 |
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| HUMAN BODY FLUIDS | <u>Blood Transfusion examinations to assist in clinical investigations</u> | Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods: | |
| Blood Plasma | Blood Group by detection and identification of antigens: A Rh D Positive AB Rh D Positive B Rh D Positive O Rh D Positive A Rh Negative AB Rh D Negative B RhD Negative O Rh D Negative Including the presence of A2 antigens | Automated using Immunocor Neo analyser (JRH) HJ 2562 NEO HJ 2559 NEO HJ 2560 NEO Automated using Immunocor Echo analyser (HH) Haem SOP 679 | JRH HH |
| Blood Plasma | Antibody Screen by detection of antibodies to: Rh -C,D,E,c,e, Cw Kell -K, K, Kpa Duffy -Fya, Fyb MNSs- M, N, S, s Kidd -JKa, Jkb Lutheran -Lua P-P1 Anti-Chido/Rogers | Automated using Immunocor Neo analyser HJ 2562 NEO HJ 2559 NEO HJ 2560 NEO | JRH |
| Blood Plasma | Antibody Screen by detection of antibodies to: Rh -C,D,E,c,e, Cw Kell -K, K, Kpa Duffy -Fya, Fyb MNSs- M, N, S, s Kidd -JKa, Jkb | Automated using Immunocor Echo analyser Haem SOP 679 | HH |



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| HUMAN BODY FLUIDS (cont'd) | <u>Blood Transfusion examinations to assist in clinical investigations</u> (cont'd) | Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods: | |
| Blood Plasma | Blood Group by detection and identification of antigens: Rh -C,D,E,c,e, Cw Kell -K, K, Kpa Duffy -Fya, Fyb MNSs- M, N, S, s Kidd -JKa, Jkb P-P1 Anti-Chido/Rogers | Manual tube spin agglutination using Diamed Column gel technology HJ 2513 (JRH) Haem SOP 462 (HH) | JRH HH |
| Plasma | Antibody Screen by detection of antibodies to: Rh -C,D,E,c,e, Cw Kell -K, K, Kpa Duffy -Fya, Fyb MNSs- M, N, S, s Kidd -JKa, Jkb | Manual Diamed Column gel IAT technology HJ 2551 | JRH HH |
| Blood Plasma | Compatibility testing of patients plasma with donor cells | Cross matching using Diamed Column gel IAT technology HJ 2531 (JRH) HH 2639 (HH) | JRH HH |
| Blood | Direct Antiglobulin Test (DAT) | Diamed Column gel IAT technology HJ 2590 | JRH HH |
| Blood Plasma | ABO titrations | Immunocor Neo analyser and microtitre plate analysis Haem SOP 439 NEO | JRH |
| Blood | Heparin Induced Thrombocytopenia Screen (HIT) | Diamed Column gel IAT technology HJ 2637 | JRH |



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| HUMAN BODY FLUIDS (cont'd) | <u>Blood Transfusion examinations to assist in clinical investigations</u> (cont'd) | Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods: | |
| Blood Plasma | Antibody investigation by detection, identification and quantification of: Rh –C,D,E,c,e, Cw Kell –K, K, Kpa Duffy –Fya, Fyb MNSs- M, N, S, s Kidd –JKa, Jkb | Diamed Column gel IAT technology HJ 2603 HJ 2597 HJ 2609 HJ 2604 HJ 2737 HJ 2605 HJ 2594 HJ 2576 HJ 2516 for including elution's, auto and cross absorptions | JRH HH |
| Blood | Foetal Maternal Haemorrhage | Clin Tech Shepard kits Kleihauer Acid Elution Technique HJ 2569 | JRH |
| Blood | <u>Haematological examinations for the purpose of clinical diagnosis</u> Erythrocyte Sedimentation Rate (ESR) | Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods: Automated Starrsed analysers, modified Westergren method Haem SOP 905 ESR Auto Compact | JRH HH |
| Plasma | Plasma Viscosity | Haem SOP 854 ESR Flex Modified Westergren method (manual) H 543 Benson Viscometers, capillary sheer stress measurements Haem SOP 110 | JRH JRH HH JRH |



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| HUMAN BODY FLUIDS (cont'd) | <u>Haematological examinations for the purpose of clinical diagnosis</u> (cont'd) | Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods: | |
| Blood | Full Blood Count (FBC) analysis of the following parameters: WBC RBC HGB HCT MCV MCH MCHC RDW-CV PLT NEUT LYMPH MONO EOS BASO Nucleated red blood cell (NRBC) count Reticulocyte count | Sysmex XN automated analysers, differential laser light scatter Haem SOP 879 | JRH CHLM HH |
| Plasma Serum | Glandular Fever Screen | Biokit Monogen commercial kit, latex micro particle immunoassay H 559 | JRH HH |
| Blood | G6PD | Commercial (Trinity Biotech) kit, functional enzyme colourimetric assay H 541 | HH |
| Plasma | Haptoglobin analysis | Commercial (Siemens NOR-Partigen Haptoglobin) kit, immunodiffusion H 644 | HH |



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| HUMAN BODY FLUIDS (cont'd) | <u>Haematological examinations for the purpose of clinical diagnosis (cont'd)</u> | Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods: | |
| Blood | Malarial antigens of: <i>Plasmodium falciparum</i> and pan-detection for: <i>Plasmodium malariae</i> , <i>Plasmodium ovale</i> & <i>Plasmodium vivax</i> | CareStart commercial kit H 3010 | JRH HH |
| | <u>Haemostasis and thrombosis examinations for the purpose of clinical diagnosis</u> | Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods: | |
| Plasma | Routine Coagulation testing for: Prothrombin Time (PT) Activated Partial Thromboplastin Time (APTT) Fibrinogen (Clauss) Thrombin Time (TT) | Sysmex CS5100 automated analysers, optical end-point detection and in-house procedures: Haem SOP 808 Haem SOP 809 Haem SOP 855 Haem SOP 830 H 1016 | JRH HH CHLM OHTC |
| Plasma | Anti-IIa (Dabigatran) | Haem SOP 1045 | OHTC |
| Plasma | Reptilase | Haem SOP 943 | OHTC |
| Plasma | Echis Time | Haem SOP 847 | OHTC |
| Plasma | High Molecular Weight Kininogen (HMWK) | Haem SOP 953 | OHTC |



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| HUMAN BODY FLUIDS (cont'd) | <u>Haemostasis and thrombosis examinations for the purpose of clinical diagnosis</u> (cont'd) | Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods: Sysmex CS5100 automated analysers, optical end-point detection and in-house procedures: | |
| Plasma | Factor Assay analysis: FII FV FVII FVIII FIX FX FXI FXII | Haem SOP 817 Haem SOP 818 Haem SOP 819 Haem SOP 834 Haem SOP 835 Haem SOP 820 Haem SOP 836 Haem SOP 837 | OHTC |
| Plasma | Factor Acquired Inhibitor Screen FVIII FIX | HC 1222 HC 1225 | OHTC |
| Plasma | Factor Inhibitor (Bethesda) analysis | Modified Nijmegen HC 1224 | OHTC |
| Plasma | Acquired Thrombophilia (Lupus) Screen (DRVVT screen / confirm, Actin FSL) | Haem SOP 1046 | OHTC |
| Plasma | Factor assay analysis FVIII FXIII | Sysmex CS5100 automated analysers, chromogenic assay and in-house procedures: Haem SOP 590 Haem SOP 839 | OHTC |
| Plasma | Inherited Thrombophilia Screen: Protein C AT | Haem SOP 1012 Haem SOP 1019 | OHTC |



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| HUMAN BODY FLUIDS (cont'd) | <u>Haemostasis and thrombosis examinations for the purpose of clinical diagnosis (cont'd)</u> | Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods: Sysmex CS5100 automated analysers, chromogenic assay and in-house procedures: | |
| Plasma | Fibrinolysis Screen: Plasminogen Alpha-2 Anti plasmin | Haem SOP 949 Haem SOP 838 | OHTC |
| Plasma | Anti-Xa | Haem SOP 919 | JRH |
| Plasma | D-Dimer | Sysmex CS5100 automated analysers, latex optical end-point detection and in-house procedures: Haem SOP 864 | JRH CHLM OHTC HH |
| Plasma | vWD analysis (Antigen) | HC 1286 | OHTC |
| Plasma | vWD analysis (Activity) | HC SOP 591 | OHTC |
| Plasma | Inherited Thrombophilia Screen: Free Protein S | Haem SOP 1022 | OHTC |
| Plasma | Ristocetin Co-Factor (RiCOF) | Sysmex CS5100 automated analysers, platelet aggregation HC 1285 | OHTC |
| Plasma | vWD Type 2N analysis | Asserachrom vwf:FVIII B manual ELISA HC 1231 | OHTC |
| Plasma | Collagen Binding Assay (CBA) | Reads Corgenix manual ELISA HC 1279 | OHTC |



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| HUMAN BODY FLUIDS (cont'd) | <u>Haemostasis and thrombosis examinations for the purpose of clinical diagnosis</u> (cont'd) | Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods: | |
| Plasma | ADAMTS13 Antibody Screen Activity | Technozym manual ELISA Haem SOP 459 Haem SOP 460 | OHTC |
| Blood Bone marrow | <u>Morphology and immunophenotyping examinations for the purpose of clinical diagnosis</u> Blood / bone marrow film analysis | Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods: May Grunwald-Giemsa staining Manual H 542, HJ 3020 or automated using Sysmex SP-10 (JRH) Haem SOP 895 or Siemens Hematek (HH) H 504 HJ 3021 Routine Staining of Blood Films and Bone Marrows-Shandon staining machine Varistain 24-4 (JRH) | JRH HH |
| Blood Bone marrow | Blood / bone marrow film analysis | Manual light microscopy HJ 3004 (blood) HJ 3006 (blood) HJ 3012 (bone marrow) HJ 3013 (bone marrow) HC2309 (bone marrow) | JRH HH |
| Blood Bone marrow | Blood / bone marrow film analysis | Automated blood film analysis using Sysmex DM analyser, light microscopy Haem 33913 HJ3400 HJ3401 Haem SOP 900 | JRH |



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| HUMAN BODY FLUIDS (cont'd) | <u>Morphology and immunophenotyping examinations for the purpose of clinical diagnosis</u> (cont'd) | Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods: | |
| Blood Bone Marrow Urine | Iron deposits | Light microscopy following Perl's Prussian Blue iron staining using commercial (TCS Biosciences) Kit HJ 3103 | JRH |
| Blood | Detection & speciation (thin films) of malaria parasites: <i>Plasmodium falciparum</i> <i>Plasmodium malariae</i> <i>Plasmodium ovale</i> <i>Plasmodium vivax</i> | Preparation of thick and thin films, manual May Grunwald-Giemsa staining and light microscopy HJ 3009 H555 | JRH HH |
| Blood Bone Marrow | Acute Leukaemia antigen detection and quantification. Panel to include: CD-45 CD-7 CD-33 CD-10 CD-19 CD-34 CD-15 CD-117 HLA-DR CD-13 CD-14 CD-64 CD-41 CD-38 | BD FacsCanto II analyser, flow cytometry HJ 3354 Haem SOP 330 HJ 3340 HJ 3306b HJ 3305 | JRH |



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| HUMAN BODY FLUIDS (cont'd) | <u>Morphology and immunophenotyping examinations for the purpose of clinical diagnosis</u> (cont'd) | Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods: | |
| Blood Bone Marrow | Acute Leukaemia antigen detection and quantification. Panel to include (cont'd): CD-11c Glycophorin-A CD-56 NG2 CD-9 CD-20 IgM CD-25 CD-1a CD-4 CD-8 CD-2 CD-57 | BD FacsCanto II analyser, flow cytometry SOPs as above | JRH |
| Blood Bone Marrow | Chronic Leukaemia antigen detection and quantification. Panel to include: sCD-3 CD-16 + 56 Cyt MPO Cyt CD-79a Cyt TdT Cyt Lysozyme Cyt CD-3 Cyt IgM CD-25 IgG1 PE IgG1 FITC CD-19 | BD FacsCanto II analyser, flow cytometry HJ 3354 Haem SOP 330 HJ 3340 HJ 3306b HJ 3305 | JRH |



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| <p>HUMAN BODY FLUIDS (cont'd)</p> <p>Blood Bone Marrow</p> | <p><u>Morphology and immunophenotyping examinations for the purpose of clinical diagnosis</u> (cont'd)</p> <p>Chronic Leukaemia antigen detection and quantification. Panel to include (cont'd):</p> <p>CD-20 CD-22 CD-23 FMC-7 CD-79b CD-79a sKappa sLambda CD-5 CD-4 CD-8 CD-45 CD-38 CD-200 sIgM CD-10 CD-103 CD-25 CD-11c CD-2 CD-3 CD-7 CD-57 CD-16 + 56 TCR-a/b TCR-g/d CD-56 CD-138 Cyt Kappa Cyt Lambda</p> | <p>Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods:</p> <p>BD FacsCanto II analyser, flow cytometry SOPs as above</p> | <p>JRH</p> |



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| HUMAN BODY FLUIDS (cont'd) | <u>Morphology and immunophenotyping examinations for the purpose of clinical diagnosis</u> (cont'd) | Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods: | |
| Blood | Paroxysmal nocturnal haemoglobinuria (PNH) clone antigen detection and quantification. Panel to include: CD-24 Flaer CD-14 CD-45 CD-15 CD-33 CD-59 Glyophorin A | BD FacsCanto II analyser, flow cytometry HJ 3323 HJ 3324 | JRH |
| Blood | <u>Haemoglobinopathy examinations for the purpose of clinical diagnosis</u> Adult Haemoglobinopathy analysis (including Antenatal) by detection of haemoglobin variants: Hb A Hb A2 Hb F Hb S Hb C Hb D Punjab Hb E | Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods: BioRad Variant II analyser, HPLC HC 2006 HC 2039 HC 2030 | JRH |



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| HUMAN BODY FLUIDS (cont'd) | <u>Haemoglobinopathy examinations for the purpose of clinical diagnosis</u> (cont'd) | Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods: | |
| Blood Dried Blood Spots | Neonatal Haemoglobinopathy analysis by detection of haemoglobin variants: Hb A Hb F Hb S Hb C Hb D Punjab Hb E | Perkin Elmer DBS Puncher and BioRad Variant NBS, HPLC HC 2026 HC 2028 | JRH |
| Blood | 2 nd Line Haemoglobinopathy analysis by detection of haemoglobin variants: Hb A Hb A2 Hb F Hb S Hb C Hb D Punjab Hb E | Perkin Elmer (Resolve FR9120), iso-electric focussing method HC 2005 | JRH |
| Blood | Sickle cell solubility | Microgen Bioproducts commercial kit, modified Itano solubility technique H 1509 | JRH HH |



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| HUMAN TISSUES AND FLUIDS | <u>Molecular genetic analysis for genetic mutations and variants</u> | Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods: | |
| Blood Bone Marrow CVS Amniotic Fluid FFPE tissue | | Manual and automated DNA extraction and quantification using the QIAGEN Symphony and in-house procedures: Haem SOP 940 HC2271 HC2316 | JRH |
| Blood Bone Marrow | | Manual and semi-automated RNA extraction and cDNA preparation using the QIAGEN Qiacube and in house procedures HC 2302 | JRH |
| Blood CVS Amniotic Fluid | Clotting deficiency: F8 gene Intron1 and 22 inversion mutations | In house method for Inversion PCR using Biometra thermocyclers HC 2252 HC 2251 HC 2100 | JRH |
| Blood CVS Amniotic Fluid | Clotting deficiencies: F5, F7, F8, F9, F10, F11, F13A genes Coagulation disorder: VWF gene Abnormal Fibrinogen: FGA, FGB, FGG genes Abnormal Antithrombin: SERPINC1 gene | In house method for Sanger sequencing using the ABI 3730 Genetic Analyser, Hotspot and full sequencing HC 2100 | JRH |
| Blood | FVL / PG20210A mutation detection | In house, multiplex PCR using Biometra thermocyclers, Qiagen Multiplex Kit and restriction digest HC 2380 | JRH |



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| HUMAN TISSUES AND FLUIDS (cont'd) | <u>Molecular genetic analysis for genetic mutations and variants</u> (cont'd) | Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods: | |
| Blood CVS Amniotic Fluid | Platelet disorders: GP1BA, GB1BB, GP9, MYH9, ITGA2B, ITG3B genes | In house method for Sanger sequencing using the ABI 3730 Genetic Analyser, Hotspot and full sequencing HC 2100 | JRH |
| Blood CVS Amniotic fluid | F7, F8, F9, F10, F11, SERPINC1 and VWF genes, deletions and duplications | Commercial MRC Holland kit for MLPA using ABI 3130 Genetic analyser system Haem SOP 612 | JRH |
| Blood CVS Amniotic Fluid | Alpha thalassemia: Identification of all known point mutations and deletions within the alpha globin gene cluster. Mutations listed on the Globin Gene Server: http://globin.cse.psu.edu/ | In house method, Gap-PCR using Biometra thermocyclers, Sanger sequencing using the ABI 3130 DNA analysis system, and commercial MRC Holland MLPA kit used for large deletions. Pyrosequencing using Qiagen Pyromark Q24. HC 2041 HC 2042 HC 2043 HC 2044 HC 2045 HC 2046 HC 2047 HC 2048 HC 2049 HC 2050 HC 2053 HC 2055 HC 2052 HC 2054 HC 2057 | JRH |



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| HUMAN TISSUES AND FLUIDS (cont'd) | <u>Molecular genetic analysis for genetic mutations and variants</u> (cont'd) | Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods: | |
| Blood CVS Amniotic Fluid | Beta thalassemia: Identification of all known point mutations and deletions within the beta globin gene cluster. Mutations listed on the Globin Gene Server:- http://globin.cse.psu.edu/ | In house method, Gap-PCR ARMS-PCR, RE-PCR using Biometra thermocyclers, Pyrosequencing using Qiagen Pyromark Q24. Sanger sequencing using ABI 3130 DNA analysis system and commercial MRC Holland MLPA kit for large deletions. HC 2101 HC 2102 HC 2103 HC 2104 HC 2106 HC 2122 HC 2075 HC 2130 HC 2142 HC 2138 HC 2143 | JRH |
| Blood CVS Amniotic Fluid | Delta-Beta thalassemia / HPFH: Identification of all known deletions within the beta globin gene cluster. Mutations listed on the Globin Gene Server:- http://globin.cse.psu.edu/ | In house method, Gap-PCR using Biometra thermocyclers, Sanger sequencing using the ABI 3130 DNA analysis system and commercial MRC Holland MLPA kit for large deletions. HC 2110 HC 2111 HC 2112 HC 2054 | JRH |



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| HUMAN TISSUES AND FLUIDS (cont'd) | <u>Molecular genetic analysis for genetic mutations and variants</u> (cont'd) | Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods: | |
| Blood CVS Amniotic Fluid | Haemoglobin S investigations | In house method, Gap-PCR, ARMS-PCR, RE-PCR using Biometra thermocyclers, Sanger sequencing using the ABI 3130 DNA analysis system, Pyrosequencing using the QIAGEN pyromark Q24. HC 2062 HC 2074 HC 2067 | JRH |
| Blood CVS Amniotic Fluid | Haemoglobin Variants: Identification of all structural variants within the beta globin gene and alpha globin gene clusters. Mutations listed on the Globin Gene Server: http://globin.cse.psu.edu/ | In house method, Gap-PCR, ARMS-PCR, RE-PCR using Biometra Thermocyclers, Pyrosequencing using Qiagen Pyromark Q24, Sanger sequencing using ABI 3130 DNA analysis system HC 2075 HC 2106 HC 2074 HC 2061 HC 2063 HC 2069 HC 2066 HC 2064 HC 2073 HC 2071 HC 2072 HC 2068 HC 2065 HC 2070 | JRH |



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| HUMAN TISSUES AND FLUIDS (cont'd) Blood DNA | <p><u>Molecular genetic analysis for genetic mutations and variants</u> (cont'd)</p> <p>Iron metabolism disorder diagnostic gene panel. Mutations in:</p> <p>TFR2; (NM_003227.3), SCL40A1, (NM_014585.5), HFE; (NM_139006.2), HFE2; (NM_213653.3), HAMP; (NM_021175.2), FTH1; (NM_002032.2), Atransferrinaemia: TF; (NM_001063.3), Hyperferritinaemia: SCL40A1, (NM_014585.5), FTL; (NM_000146.3), Hereditary Hyperferritinaemia Cataract Syndrome (HHCS): FTL; (NM_000146.3) , Iron-refractory iron deficiency anaemia (IRIDA): SLC11A2; (NM_001174125.1), TMPRSS6; (NM_153609.2, Iron deficiency: HEPH; (NM_138737.3), iron and neurodegeneration:- , CP (Aceruloplasminaemia), (NM_000096.3), FTL (Hereditary ferritinopathy), X-linked Sideroblastic anaemia:- ALAS2 ; (NM_000032.4), Genes involved in the BMP/SMAD pathway:- BMP4; (NM_001202.3), BMP6; (NM_001718.4) SMAD4; (NM_005359.5).</p> | <p>Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods:</p> <p>MiSeq Next Generation Sequencing and custom designed commercial panel (Illumina TSCA v1.5 assays) HC 2507 HC 2509</p> | JRH |



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| HUMAN TISSUES AND FLUIDS (cont'd) Blood DNA | <p><u>Molecular genetic analysis for genetic mutations and variants</u> (cont'd)</p> <p>Somatic myeloid disorder diagnostic gene panel. Mutations in:</p> <p>ASXL1 (NM_015338.5), ATRX (NM_000489.3), CBL (NM_005188), CBLB (NM_170662.3), CBL (NM_012116.3), CSF3R (NM_156039.3), DNMT3A (NM_022552), ETV6 (NM_001987.4), EZH2 (NM_004456.4), HRAS (NM_005343.2), IDH1 (NM_005896.2), IDH2 (NM_002168.2), FLT3 (NM_004119), JAK2 (NM_004972.3), KIT (NM_000222.2), KRAS (NM_033360.2), MPL (NM_005373), NPM1 (NM_002520.6), NRAS (NM_002524.4), PDGFRA (NM_006206.4), PHF6 (NM_032458.2), PTEN (NM_000314), RUNX1 (NM_001754.4), SETBP1 (NM_015338.5), SF3B1 (NM_012433.2), SRSF2 (NM_001195427.1), TET2 (NM_001127208.2), TP53 (NM_000546.5), U2AF1 (NM_001025203.1), WT1 (NM_024426.4) and ZRSR2 (NM_005089.3).</p> | <p>Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods:</p> <p>MiSeq Next Generation Sequencing and custom designed commercial panel (Illumina TSCA v1.5 assays) HC 2507 HC 2509</p> | JRH |



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| HUMAN TISSUES AND FLUIDS (cont'd) | <u>Molecular genetic analysis for genetic mutations and variants</u> (cont'd) | Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods: | |
| Blood | HFE gene screen (C282Y / H63D) | In house method RE-PCR using Biometra thermocyclers HC 2263 | JRH |
| Blood DNA | HFE, HFE2, HAMP, SLC40A1 and TFR2 genes / DNA | ABI 3130 Genetic Analyser and commercial MRC Holland MLPA kit used for large deletions HC 2297 | JRH |
| Blood Bone Marrow | BCR-ABL analysis (ALL, CML) Diagnosis and monitoring of MRD (p210, p190 only) | Published Multiplex RT-PCR method using Qiagen, Rotorgene, Commercial QPCR (Qiagen) kit for MRD, Pyrosequencing using QIAGEN Pyromark Q24 HC 2301 HC 2300 HC 2303 HC 2306 | JRH |
| Blood Bone Marrow | Acute Leukaemia Diagnosis: FLT-3 D835 | RE-PCR using Biometra Thermocycler HC 2118 | JRH |
| | NPM1 | In house fragment analysis method using ABI 3130 Genetic Analyser HC 2119 | JRH |
| | c-KIT D816V | Pyrosequencing using QIAGEN Pyromark Q24 HC 2123 | JRH |
| Blood Bone Marrow | Acute Leukaemia MRD monitoring (PML-RARA, Inv16, AML-ETO) | Commercial QPCR (Qiagen) kit for MRD using QIAGEN Rotorgene, HC 2304 | JRH |



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| HUMAN TISSUES AND FLUIDS (cont'd) | <u>Molecular genetic analysis for genetic mutations and variants</u> (cont'd) | Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods: | |
| Blood Bone Marrow Tissue | Acute Leukaemia Diagnosis (cont'd): Lymphoma B Clonality (IgH rearrangements) | Commercial (Invivoscribe) fragment analysis using ABI 3130 Genetic Analyser HC 2273 | JRH |
| | Lymphoma T Clonality (TCR re-arrangements) | HC 2268 HC 2267 HC 2270 | JRH |
| Blood Bone Marrow | TP53 gene mutation analysis (CLL only) | In house Sanger sequencing using the ABI 3730 Genetic Analyser HC 2305 | JRH |
| Bone Marrow | Jak-2 Exon 12 mutation analysis | Haem SOP 250 | JRH |
| Blood Bone Marrow | Jak-2 mutation screen (V617F) | In house ARMS-PCR using Biometra thermocycler, Pyrosequencing using the QIAGEN Rotorgene HC 2261 | JRH |
| Blood Bone Marrow | Chimerism analysis | Promega Powerplex Fusion commercial kit, Q-PCR using ABI Real Time 7500 HC 2120 HC 2127 | JRH |



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| HUMAN TISSUES AND FLUIDS (cont'd) | <u>Molecular genetic analysis for genetic mutations and variants</u> (cont'd) | Procedures documented in manufacturer's equipment manuals in conjunction with documented in-house procedures by the following methods: | |
| FFPE DNA | EGFR common mutation screen | COBAS 4800 system, RT-PCR and commercial (COBAS) kits Haem SOP 578 | JRH |
| FFPE DNA | KRAS mutation screen (codon 12, 13, 61) | Haem SOP 444 | JRH |
| FFPE DNA | BRAF mutation screen (V600E) | Taqman assay Haem SOP 579 | JRH |
| FFPE DNA | Somatic Cancer 50 gene diagnostic panel. Mutations in: TP53 NM_000546.5, PTEN NM_000314.4, PIK3CA NM_006218.2, PDGFRA NM_006206.4, KRAS NM_004985.3, NRAS NM_002524.4, KIT NM_000222.2, EGFR NM_005228.3, BRAF NM_004333.4, ERBB4, FGFR1, ERBB2, MET, FLT3, FGFR3, GNAS, SMARCB1, CTNNB1, CDKN2A, ABL1, NOTCH1, ATM, PTPN11, SMO, SMAD4, VHL, NPM1, MPL, CSF1R, HRAS, JAK3, AKT1, IDH1, CDH1, FGFR2, SRC, KDR, ALK, JAK2, RB1, MLH1, HNF1A, APC, RET, STK11, FBXW7, EZH2, GNA11, GNAQ and IDH2 | Thermofisher Ampliqseq Next Generation Sequencing commercial panel and Ion Torrent next generation sequencing. Haem SOP 624 Haem SOP 491 Haem SOP 625 | JRH |
| END | | | |