**PATHOLOGY USER HANDBOOK**

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**GENERAL PATHOLOGY**

**INTRODUCTION**

The Pathology department is accredited by the United Kingdom Accreditation Service (UKAS) to the International Standard **‘Medical laboratories - Requirements for quality and competence (ISO 15189:2012)’** and maintains a quality service by the implementation of a comprehensive internal Quality Management System coupled with an extensive External Quality Assurance programme.

Our accreditation is limited to those activities described on our UKAS schedule of accreditation available on the UKAS website using the following link: <http://www.ukas.com/search-accredited-organisations> and by entering the Accreditation number **8429**.

Laboratory tests, technology and methodology are evolving, and some tests are not yet accredited and consequently not listed on the schedule of accreditation. Please contact the laboratory for further information about specific tests or if you have any concerns.

This handbook has been produced to provide information which will allow you to make best use of our services.

Information on the services provided and contact telephone numbers are available within this document and on the hospital intranet,

Pathology is part of the Clinical Service Unit- Diagnostics and Screening, hereby referred to as the **Pathology CSU**.

The Pathology CSU provides a comprehensive, consultant lead service by a combination of in-house and external provision. Routine high volume and emergency investigations are provided in-house, other specialist investigations are sent to accredited reference laboratories.

The Pathology CSU comprises Pathology Support Unit, Haematology, Blood Transfusion and Immunology, Chemical Pathology, Microbiology and Cellular Pathology (Histopathology, Diagnostic Cytology and the Mortuary). Cervical Screening Cytology is provided by Norfolk and Norwich University Hospital Cellular Pathology.

Pathology’s Consultants are available to advise on the selection of appropriate tests and help with the interpretation of results.

Pathology results are stored on a centralised Pathology computer system and are made available to Acute Trust service users via eCare. eCare training and passwords are available from the IT helpdesk, extension 87000.

GP requested results are transmitted electronically via the GP Messaging serv

into the GP system in regular downloads and GP practices have access to ICE for requesting and reporting.

Please contact Pathology System Manager or System Support Officer on 01908 995812 if there are any problems.

The main Pathology reception is situated next to Ward 20, on Level 2 of the Hospital building, adjacent to the Haematology Outpatient’s Clinic. The Blood Bank Issue Refrigerator is situated in a room just outside Pathology reception. The room is clearly signposted and approved access is via your hospital ID badge/swipe card. Cellular Pathology and the Mortuary are situated on Level 1 at the rear of Ward 3. Please see the map at the end of this document.

Pathology staff are always willing to be involved in clinical audit, please contact the appropriate Head of Department.

Pathology is pleased to accept samples from non-NHS institutions for private, Category 2 and contract health screening. Prices are available on request, please discuss your requirements with the Pathology Services Manager on 01908 995811or contact the Pathology Business Support Officer on 01908 995794.

We welcome the opportunity that allows our users to visit us on site and gain insight into our laboratory services. Please contact Jill Beech on 01908 995811or or via email: [jill.beech@mkuh.nhs.uk](mailto:jill.beech@mkuh.nhs.uk)

*Despite every effort, it is possible that mistakes will exist in this handbook. If errors are detected please let Pathology know by contacting the Pathology Quality Manager on 01908 995823 or email –* [jessica.dixon@mkuh.nhs.uk](mailto:jessica.dixon@mkuh.nhs.uk). *Please use this handbook and let us know your views so that it can be improved.*

## Measurement Uncertainty

The laboratory considers measurement uncertainty when interpreting measured quantity values. Estimates of measurement uncertainty for test values will be made available on request.

## Feedback and Complaints Procedure

The Pathology Department welcomes feedback from our service users. Should you wish to contact Pathology to lodge a complaint or comment, please contact the appropriate Head of Department, Pathology Services Manager or Quality Manager.

A copy of our complaints procedure [MPCOMP Pathology Complaints Procedure] is available on request and is written to conform to the Trust Complaints Policy available on the intranet

Complaints and comments can be made verbally or in writing (including electronically) and will be documented by pathology staff and investigated as soon as possible.

All written complaints will be acknowledged within 2 working days of receipt by the Pathology Quality Manager or Head of Department.

A written response will be provided following investigation.

## Protection of Personal Information

*The Pathology Department strictly adheres to the requirements of the Data Protection Act and to the requirements of the Trust Data Protection Policy available on the Trust intranet*

**PATHOLOGY CSU TELEPHONE NUMBERS**

**(MKU HOSPITAL SWITCHBOARD: 01908 660033)**

**General Pathology** **Ext. Direct Dial Bleep**

Director of Pathology – Dr Khalid Enver 85681 99 5681

Pathology Services Manager – Jill Beech 85811 99 5811 1007

Pathology Business Support Officer – Alex Badger 85794 99 5794

Pathology Quality Manager – Jessica Dixon 85823 99 5823

Pathology Quality Associate Practitioner - Christine Marsh 85823 99 5823

Pathology Systems Manager – Pirran Salter 85812 99 5812

Pathology Supplies (bottles & cards) 85793 99 5793

**Pathology Support Unit, Haematology, Chemical Pathology & Microbiology**

Results/Enquiries 85768 99 5768

Urgent samples 85842 99 5842

Enquiries re referred samples 85772 99 5772

PSU Manager – Helen Botwood 85769 99 5769

**Blood Science Manager** Grant Barker 85830 995830

Blood Science Business Officer. – Yvonne Brown 85754 99 5754

**Haematology**

Consultant Haematologist - Dr Catherine Hildyard 85817 99 5817 1241

Consultant Haematologist - Dr Subir Mitra 85753 99 5753

Consultant Haematologist - Dr Moez Dungarwalla 85756 99 5756 1163

Consultant Haematologist –Dr Mags Akanni 87573 99 7573 1206

Consultant Haematologist –Dr Sarah Davis 87574 99 7574 1789

Consultant Immunologist - Dr Liz Bateman (01865) 225991 / 225995

Secretarial Support 85814 99 5814

Secretarial Support 85815 99 5815

**Technical Enquiries** 85764 99 5764

**Out of routine hours bleep** 1412

Operational Manager – Alison McEvoy 85780 995780

**Blood Transfusion** 85776 99 5776

Blood Bank Manager - Jasmine Beharry 85832 99 5832

**Technical Enquiries** 85776 99 5776

**Out of routine hours bleep** 1412

Specialist Practitioner of Transfusion – Caroline Lowe 85798 99 5798 1644

Specialist Practitioner of Transfusion – Terrie Perry 85798 99 5798 1644

**Chemical Pathology**

Consultant Chemical Pathologist Dr Farhan Ahmed 85792 99 5792

Operational Manager – Ben Powell 85831 995831

**Technical Enquiries** 85761 99 5791

**Out of routine hours bleep** 1413

**Microbiology Laboratory Manager**

Imran Sheikh 85790 995790

Micro / Cellular Pathology Business Support Office

Brodie Woodgate 85839 995839

**Microbiology**

Consultant Microbiologist - Dr Mansoor Raza 85799 99 5799

Consultant Microbiologist - Dr Poonam Kapila 85786 99 5786

Consultant Microbiologist – Dr Prithriwaj Chakrabarti 85796 99 5796

Secretarial Support 85782 99 5782

Technical Enquiries 58779 99 5779

Chief Biomedical Scientist - Carol Jones 85781 99 5781

Asst Dir of Infection Prevention – Angela Legate 85789 99 5789 1182

**Cellular Pathology**

Consultant Histopathologist - Dr. Ann Abraham 85806 99 5806

Consultant Histopathologist - Dr Angus Molyneux 85807 99 5807

Consultant Histopathologist - Dr Sherly Mathews 85808 99 5808

Consultant Histopathologist - Dr Niveen Abdullah 85809 99 5809

Consultant Histopathologist - Dr Achamma John 85810 99 5810

Consultant Histopathologist - TBA 85836 99 5836

Consultant Histopathologist - Dr Moyna Dyer 85763 99 5763

Secretarial Support 85802 99 5802 / 3 / 4

**Technical Enquiries** / Main Lab 85819 99 5819

**Cellular Pathology Laboratory Manager** - Liz Thwaites 85820 99 5820

**Mortuary**

Mortuary Manager – Joanne Smith 85828 99 5825 / 8

## Normal Working Hours

Routine and urgent services are available during normal opening hours, which are:

Monday to Friday - 09:00 - 17:00 Saturday / Sunday & Bank Holidays - 09:00 - 16.30 (skeleton staff only)

(Cellular Pathology Monday to Friday only)

Mortuary normal hours are 08.00 - 16.00 and emergency out of hours.

Outside of these hours an ‘Emergency Out-of-Hours’ service is provided for Haematology & Blood Transfusion, Chemical Pathology and Microbiology.

## Urgent, Preliminary and Telephoned Results

During normal working hours please notify the laboratory of all urgent requests on ext. 85842 (all Accident & Emergency, Surgical Assessment Unit, Ward 5/Neo-Natal Unit, Medical Assessment Unit, Ambulatory Emergency Care Unit and Department of Critical Care requests are treated as urgent).

Appropriate results will be telephoned or made available on the computer as soon as possible.

During processing, results may be obtained which may affect immediate patient management. These results will either be issued as preliminary/interim reports or telephoned to the requester. The provision of accurate, brief relevant clinical details is

essential to enable workload prioritisation.

Outside normal working hours urgent requests are dealt with by the ‘out-of-hours’ service.

## ‘Out of Hours’ Service

Biomedical Scientists (BMS) for Haematology and Chemical Pathology are always resident in the Hospital and may be contacted via the Switchboard. BMS for Microbiology are resident in the hospital up to 20:45 and may be contacted by bleep via the Switchboard.

Consultant staff are also available for advice and may be contacted via the Switchboard. Arrangements for the provision of advice from a Consultant Microbiologist are as follows: In order to provide a 24/7 advice service 365 days of the year the pool of available Consultants includes colleagues at The Great Western Hospital, Swindon, and cross-site cover is provided in this way. To maintain this arrangement responsibilities are specified and calls will be taken from GPs and Registrar grades and above. Exceptions are in place to specify that calls will be taken from surgical SHOs in the operating theatre and from ITU medical staff.

Switchboard operators are aware of this arrangement and will follow this protocol.

**Haematology and Chemical Pathology:** If you require investigations for IMMEDIATE patient management or examinations that are time sensitive please bleep the relevant ‘on-call’ BMS out of hours who will analyse the sample as soon as possible on receipt.

**Haematology Bleep 1412**

**Chemical pathology Bleep 1413**

Results will be made available on the computer systems or telephoned if critically abnormal. The provision of accurate, brief relevant clinical details is essential to enable workload prioritisation.

**Microbiology:** After 20:45 and up to 08:45 calls for CSFs will be made directly to the BMS via Switchboard to their home/mobile telephone. All other calls will be diverted to the Consultant Microbiologist. If it is deemed necessary for a BMS to come in, the Consultant Microbiologist will request the switchboard to contact and instruct the BMS on call. Microbiology samples other than blood cultures and CSFs should not be sent to the laboratory after 20:45 but refrigerated and sent after 08:45 the following day.

**For all other samples**, if the investigations are NOT required for immediate patient management, you do not need to bleep the on-call BMS. Send the samples to Pathology and they will be processed, and results will be available via the computer systems. Routine enquiries made outside “office hours” run the risk of impeding the lone working BMS in the business of dealing with emergencies.

Blood cultures should be placed in the incubator (specially labelled), in the ‘Blood Bank Issue Room’ outside the laboratory. Significant positive results will be telephoned by laboratory staff.

If you require additional tests on samples already sent to Pathology, an additional request card MUST be sent to verify the request.

## Pathology Supplies

The following items may be requested:

|  |  |
| --- | --- |
| **Blood bottles:** | FBC – LAVENDER |
|  | COAGULATION – BLUE |
|  | TRANSFUSION – PINK |
|  | SST (GEL) – GOLD |
|  | LITHIUM HEPARIN – GREEN |
|  | PLASMA GLUCOSE – GREY |
|  | TRACE ELEMENT – DARK BLUE – please state test required |
|  | PLAIN LITHIUM HEPARIN (FOR TB ELISPOT) – DARK GREEN |
|  | PAED WHITE |
|  | PAED PINK |
|  | PAED BLUE |
|  | PAED ORANGE |
|  | PAED RED |
|  | PAED GOLD |
|  | PAED GREEN |
|  | PAED PURPLE |
|  | SBR’S – BROWN |
|  | PAED GLUCOSE – YELLOW |
|  | |
| **Other Containers:** | BLOOD CULTURE SETS |
|  | BLOOD CULTURE PAEDIATRIC |
|  | 10ML BORIC ACID TUBE FOR PAEDIATRIC URINE SAMPLES |
|  | U-BAG (INFANT) |
|  | WHITE TOP UNIVERSAL CONTAINER |
|  | BLUE TOP STOOL CONTAINERS |
|  | SPUTUM POTS |
|  | FORMALIN POTS (State size required) 20ml,60ml,480ml, 2.5L, 5L, 10L or a dry pot. |
|  | CHLAMYDIA MALE – URINE |
|  | CHLAMYDIA FEMALE – SWAB |
|  | DERMAPAK |
|  | FAECAL IMMUNOCHEMICAL (FIT) TUBE (flat, green lid) |
|  | LBC KITS (SMEAR POT AND BRUSH) |
|  | |
| **Swabs:** | BLACK SWABS (black swab, amies charcoal medium) |
|  | PERNASAL SWABS (black swab, wire shaft) |
|  | MRSA PCR SWAB (RED TOP COPAN DUAL SWAB) |
|  | VIRAL SWABS RED TOP (Viral PCR other than Flu/RSV) |
|  | FLU/RSV (XPERT VIRAL TRANSPORT) SWABS  (approved locations only) |
|  | OTHER ITEMS: |
|  | |
| **Forms:** | PINK BLOOD FORMS |
|  | ANTE-NATAL REQUEST FORMS |
|  | ICE REQUEST FORMS |
|  | ICE REQUEST BAGS |
|  | BLUE & WHITE REQUEST CARDS |
|  | BLUE HISTOPATHOLOGY/CYTOLOGY CARDS |

Please ensure your departments have adequate stock to avoid the necessity for immediate stock replenishment. The Pathology Supplies department is not permanently staffed throughout the day and cannot support same day dispatch.

Please order as below:

·       **ROUTINE** orders are to be made using the [Pathology Supplies Order Form](https://gbr01.safelinks.protection.outlook.com/?url=https%3A%2F%2Fintranet.mkuh.nhs.uk%2Fwp-content%2Fuploads%2F2020%2F08%2FINTERNAL-Supplies-Order-Form-Pathology-1.docx&data=02%7C01%7CCarol.Jones%40mkuh.nhs.uk%7C672631935e60492669cd08d83d3818b5%7Ce96dd0a15d474a949e4a5c1056daa82c%7C0%7C0%7C637326658603954556&sdata=aWwrjMW1Gg9LaSpWIs6HmM9p%2F8bsKVZik7RBF1QUM2s%3D&reserved=0)  or email your order to [pathologysupplies@mkuh.nhs.uk](mailto:pathologysupplies@mkuh.nhs.uk). These orders will be dispatched within two working days (Monday to Friday).

·       **URGENT** orders (not multiple stock requests) can be made by telephoning 01908 995793 or extension 85793 and leaving a message. Please note, this telephone is not staffed and the orders will be dispatched the next working day (Monday to Friday).

·       **24-hour urine** containers are ordered by telephoning 01908 995768 or extension 85768. You will need to specify if a plain or acidified container is required. If this is unknown the test required must be stated. A patients name **must** be given for any requested acidified container. The orders will be dispatched the next working day (Monday to Friday).

The Pathology Porters will deliver within the hospital and the Hospital Transport will deliver to the GP Practices.

## Request Cards

Request forms are not required (unless specified at the time of placing an order) for eCare pathology orders. Requests for investigations generated manually or via ICE must include a form/card. Although the same manual request card is used for Chemical Pathology, Haematology and Microbiology; **please** use one form for Chemical Pathology and/or Haematology tests and a separate form for Microbiology tests (hence 2 separate forms from the same patient). If using the ICE order communications system please use ALL the request forms printed by the system, ensuring that the correct specimen is attached to each form.

For requests made using the eCARE order communications system please print request cards when prompted by the system.

There are separate request cards for Cellular Pathology (blue) and Blood Transfusion (pink).

A combined antenatal request form for infectious diseases, sickle cell and thalassaemia in pregnancy is available.

Request cards for Chromosomes/Genetic Markers are available directly from the Churchill Hospital, Oxford or can be printed from ICE

In many cases the exact tests performed will be greatly influenced by the clinical details supplied. Request cards must be completed with:

1. Patient’s full name,
2. Hospital number or NHS number (if available),
3. Date of birth,
4. Address of patient,
5. Patient’s Consultant and/or GP (This information can be provided using a PID label, please ensure a label is placed on both copies of the joint request card),
6. Destination for report,
7. Date and time of sample,
8. Clinical details including:

a) Clinical features including whether hospital or community acquired

1. Any history of infection with dangerous pathogens such as TB, Neisseria meningitidis, Brucella, Salmonella typhi

c) Details of foreign travel.

d) Onset and duration of illness (esp. for serology).

e) Details of recent (one week), current and intended chemotherapy.

f) Specify site from which samples were taken.

g) Details of other therapies.

h) For pre-op assessment samples state the surgical procedure

1. GP patients – please add a patient telephone number in the event of

results produced out of hours being critically abnormal.

There are exceptions to the above:

1. Blood Borne Virus Clinic: samples and cards are labelled with Clinic Number and date of birth only.
2. Infectious diseases in pregnancy screening requests require the following information:

* NHS number, MRN, Forename and Surname.
* Estimated Date of Deliver (EDD)
* First line of address and Postcode
* Date of Birth
* GP name and/or code.
* Name and location of requester
* Maternity unit booked for delivery
* Address for results and reports
* Name and location of sample taker
* Date and time of sample collection
* Priority Status –

Initial antenatal screening sample

Repeat antenatal screening sample

Repeat sample to exclude recent infection

Initial sample after previous decline

* Examinations requested,
  + - accept/decline for each examination
    - known positive/unknown status for each
* Clinical indications for urgent testing if required. (NB for urgent testing please contact the laboratory by telephone).

**Labelling for ‘Danger of Infection’:**

Samples from the following categories of patients must be identified as ‘Danger ofInfection’:

(i) All sputa specimens, whether or not a diagnosis of tuberculosis is clinically suspected.

(ii) Any material suspected to contain M. tuberculosis.

(iii) Blood / Tissue specimens from patients with suspected/confirmed Hepatitis B, Hepatitis C or HIV infections.

(iv) Clinical specimens from known, suspect or at risk patients to transmissible Spongiform Encephalopathy agents.

(v) Patients with suspected Typhoid or Brucellosis

Other high-risk groups exist. If in doubt, please telephone Microbiology, Ext. 58779 for clarification.

**Labelling and packing procedures (details also on request card):**

(i) Label specimen and request card.

(ii) Apply a **DANGER OF INFECTION** label to the specimen and request card.

(iii) Place the sample in the specimen compartment of the bag and seal.

## Sample Collection

Every day Pathology receives numerous patient samples of varying types from the Hospital, GPs and other sources hence it is essential that samples and request cards be correctly identified.

The **minimum** essential requirements for labelling samples are:

Forename, Surname, Date of Birth and NHS number or Hospital number or 1st line of address. Blood Transfusion samples from Acute Trust patients must be labelled with the Hospital number.

**Please DO NOT USE Patient Identification (PID) labels on any blood bottles** unless produced on demand from the Order Communication system. Blood Transfusion samples MUST be handwritten and signed by the person taking the sample.

Additional information required is: The identity of the person collecting the sample and the collection date and time.

It is the Sample Taker’s responsibility to ensure the accuracy of information on both the sample and request card

**SAMPLES AND REQUESTS CARDS WHICH ARE UNLABELLED OR INCORRECTLY LABELLED AND CANNOT THEREFORE BE IDENTIFIED WILL NOT BE ACCEPTED FOR ANALYSIS.**

**CHECK EXPIRY DATES ON ALL STOCK BEFORE USING IT AND RETURN TO PATHOLOGY IF OUT OF DATE STOCK DISCOVERED. SAMPLES RECEIVED IN EXPIRED STOCK WILL NOT BE PROCESSED.**

All procedures carried out on a patient need the informed consent of the patient. For most routine laboratory procedures, consent can be inferred when the patient presents himself or herself at a laboratory / phlebotomy area / ward with a request form and

willingly submits to the usual collecting procedure, for example, venepuncture.

Patients in a hospital bed should normally be given the opportunity to refuse.

Blood samples are received in containers, which may or may not contain an anticoagulant. The purpose of an anticoagulant is to prevent the blood from clotting. Samples must be mixed after collection to avoid clotting.

The more common sample bottles are:

|  |  |  |
| --- | --- | --- |
| **Bottle** | **Anticoagulant** | **Usage** |
| Gold top (gel bottle) | no anticoagulant (clotted blood) | General |
| White top(paed) | no anticoagulant (clotted blood) | General |
| Light Green top | Lithium heparin | ITU, A&E \* |
| Orange top | Lithium heparin | Paed |
| Lavender top | Potassium EDTA | FBC, ESR, Malaria Parasites, GF screen, G6PD |
| Clear lavender top (paed) | Potassium EDTA | FBC, ESR, Malaria Parasites, GF screen, G6PD |
| Red top (neonates) | Potassium EDTA | FBC, Malaria Parasites, GF screen, G6PD (**NOT ESR**) |
| Blue top | Sodium citrate | INR, APTT, Clotting Screen, thrombophilia, lupus, factor assays. |
| Clear Blue top (paed) | Sodium Citrate | INR, APTT, Clotting screen, thrombophilia, lupus, factor assays. |
| Grey top | Sodium fluoride/potassium oxalate | Lactate/Glucose  Please contact Chem for bottle |
| Yellow top (paed) | Sodium fluoride | Lactate/Glucose (Paed)/Plasma Glucose  Please contact Chem for bottle |
| Pink top (plastic round bottom) | EDTA | Grouping/ Crossmatching |
| Pink top(plastic round bottom) | EDTA | Antenatal Grouping & Abs |
| Dark green top | Plain Lithium Heparin | IGRA tests |

\*These bottles along with other specially prepared containers are used for certain tests identified in the handbook; please contact the laboratory to check procedures before commencing.

It is essential to use the correct sample containers with the appropriate amount of blood. This ensures the correct blood-to-anticoagulant ratio and when necessary, minimises the risk of clotting. NEVER MIX BLOOD FROM ONE BOTTLE WITH ANOTHER. The wrong lid on the wrong bottle can also lead to contamination and erroneous results.

Remember to gently mix the sample in the bottle after collection (using full inversions for the number of times stated on the BD collection cards), seal the sample in the plastic bag attached to the card, or in a separated plastic bag for Cellular Pathology, and then despatch the sample as soon as possible.

Other sample containers are detailed later in the discipline-specific parts of this handbook.

## Genetic Studies/Tissue Typing

Sample requirements will be found in the discipline sections of this handbook

i.e. Chromosomes - Chemical Pathology, HLA - Blood Transfusion etc.

## Phlebotomy Service

A morning service to the wards is provided by 18 part-time staff (covering the hours Monday to Friday 08:00 – 12:00) and by one Phlebotomist in the afternoon to bleed patients not available in the morning.

Phlebotomy at weekends / Bank Holidays is provided by a team of 6 phlebotomists working 8 - 11am. It is limited to 8 cards per ward.

Phlebotomy is no longer provided to Campbell Centre or Marlborough House

Requests must be made via eCare placing the orders onto the phlebotomy ward round. These rounds close at 8am 7 days per week.

Any samples taken by phlebotomy staff will not be treated as urgent.

## Transport of samples

GP samples are collected from practices by the community drivers at least once a day, some practices, at their request, have a later second collection.

The Pathology porters collect routine samples from all wards mid-morning and from Outpatients twice a day as part of the ward round, when results and supplies will also be delivered.

CSF, High Risk samples and Blood Gases must **NOT** be sent via the ‘air tube’.

Other samples should be sent via the ‘air tube’ delivery system, instructions are on each station. The reliability of the ‘air tube’ system CANNOT be guaranteed so beware before you use it. If the ‘air tube’ is not functioning, urgent samples should be sent via the general portering system, the pathology porters **do not** collect urgent samples.

## Pathology Reports

Pathology reports are printed centrally in Pathology and distributed only to agreed locations. All GP practices accept electronic download of Pathology results.

## Computer System Enquiries

All authorised results for Haematology and Blood Transfusion, Chemical Pathology, Microbiology and Cellular Pathology are available via eCare/ ICE on the Hospital network. All Medical staff should have a password that enables them to interrogate the computer for Pathology results. This is the preferred method of result notification as it eliminates transcription errors. If you require further instruction on the use of eCare/ICE please contact the IT Training Department on 87000.

**MICROBIOLOGY**

**INTRODUCTION**

Microbiology provides services for:

1. Bacteriological and Virological diagnosis

2. Advice on antimicrobial treatment and prophylaxis

3. Advice on epidemiology and prevention of infection including control of Hospital infection

**TAKING SAMPLES**

Appropriate timing of sample taking is crucial.

1. Please put date and time taken on all request forms (enabling us to assure quality of samples and monitor turnaround time form receipt to report.
2. For **Antibody tests**: please put the onset date of the illness on the request form.
3. For antibody tests on patients in contact with infectious disease, give the date and nature of the contact.

Factor affecting tests:

1. Delayed transport will have a detrimental effect on microbiological investigations. Where delay is unavoidable all samples OTHER THAN Blood Cultures and Chlamydia/Neisseria gonorrhoeae PCR are best stored at 4°C. Urine MC&S samples in boric acid preservative will be stable for up to 96 hours at room temperature.
2. Sample for culture should when possible be collected prior to the commencement of antibiotic therapy.

Requests for additional investigations:

Some investigations may be added following despatch of samples to the laboratory. These requests must be made within 48hrs of despatch and discussed with the laboratory technical staff. Tel 01908 995799. A request form must be sent for any add on test.

Please note that we are unable to share samples with Chemical pathology/haematology unless in exceptional circumstances.

**SPECIFIC NOTES**

### Antibiotic Assays

Vancomycin and Gentamicin assays are carried out by the Chemical Pathology Department. Other antibiotic assays (see list below) are sent to the Microbiology Department and referred to another laboratory for testing. The Consultant Microbiologist provides pre- and post- analysis interpretation and advice on all antibiotic assays regardless of which Pathology department analyses the sample. We encourage users to discuss referred antibiotic assays with the Consultant Microbiologist who will advise on the correct timing for sample collection and ensure that the laboratory is aware and prepared to make arrangements for sample transport to the reference laboratory.

**Vancomycin**: Please refer to Trust Antibiotic Policy, available on the intranet, for therapeutic and monitoring guidelines

**Gentamicin**: Please refer to Trust Antibiotic Policy, available on the intranet, for therapeutic and monitoring guidelines

**Instructions and timing of Serum Gentamicin Levels are given in the** gentamicin calculator which can be found on the Trust Intranet (Pharmacy Policies)

Teicoplanin: Tests are referred for testing. Please discuss with the Consultant Microbiologist before sending samples.

See also Antimicrobials Guidelines on the Trust Intranet (Pharmacy Policies)

|  |
| --- |
| **Antibiotic assays available (referred):** |
| Amikacin |
| Chloramphenicol |
| Colistin |
| Cycloserine  Ethambutol |
| Flucytosine |
| Itraconazole |
| Streptomycin |
| Teicoplanin |
| Tobramycin |
| Voriconazole |

Please discuss with the Consultant Microbiologist before requesting.

Assays for other antimicrobials may be available if required. Please contact the Consultant Microbiologist to discuss.

**For all antimicrobial assays:**

Collect blood sample (minimum 1ml) into a gold topped bottle or plain white paediatric bottle.

**Antimicrobial Assay Result Enquiries:**

If it is necessary to enquire for a referred antibiotic assay result outside normal laboratory hours the reference laboratory can be contacted on 0117 414 6220 or on 0117 414 8469 from 09:00 to 14:00 on Saturday only.

Advice can be given out-of-hours, by contacting the reference laboratory Hospital switchboard (0117 950 5050) and asking them to contact the on-call Medical Microbiologist

### Biopsies for culture and sensitivity

Collect specimens before antimicrobial therapy where possible.

Use aseptic technique.

Collect specimens into sterile plain (white topped) universal containers and place in sealed plastic bags.

If specimen is small, place it in sterile water to prevent desiccation.

Note: Specimens received in formalin are not suitable for culture.

Suspected Legionella species (lung tissue and biopsy)

If specimen is small place it in sterile water to prevent desiccation. Avoid the use of saline, as it is known to be inhibitory to Legionella species.

**3.** **Helicobacter pylori culture from gastric biopsy:**

This test can only be carried out by the Gastroenterology department by prior arrangement with the Microbiology Consultant.

Ideally biopsies should be taken before antimicrobial therapy is begun, however a ‘test and treat’ strategy for the diagnosis of H. pylori is recommend by NICE and therefore most samples referred for culture will be due to treatment failure. A period of at least two weeks should have elapsed since the last dose of antimicrobial therapy before the collection of the specimen.

Gastric biopsy specimens are usually taken from the gastric antrum at endoscopy, and sometimes from the main body of the stomach depending on location of inflammation. Duodenal biopsies will be taken in cases with duodenal ulcers.

The biopsy must be collected directly into sterile saline in a universal container.

Samples must reach the Microbiology department by 4pm on Monday to Thursday only.

### 4.Blood Cultures

Collect specimens before antimicrobial therapy where possible.

Collect specimens as soon as possible after the onset of clinical symptoms. Although blood can be sampled at any time, drawing blood at, or as soon as possible after a fever spike is optimal, except in endocarditis where timing is less important.

* The use of iodine-based disinfectants is not recommended for disinfection of the butyl rubber septum for some commercial systems as this may affect the septum’s integrity.
* The use of blood collection adapters without ‘winged’ blood collection sets is not recommended as it is not possible to accurately judge the sample volume and there may be the potential for backflow of blood culture media to patient veins.
* If blood for other tests such as blood gases or ESR is to be taken at the same venepuncture, the blood culture bottles should be inoculated first to avoid contamination. It is preferable to take blood for culture separately.

A blood culture set is defined as one aerobic and one anaerobic bottle. For infants and neonates, a single paediatric bottle may be requested.

**Sample Volume:**

Blood culture sets (Aerobic and Anaerobic) 8-10ml for each bottle.

Paediatric bottles 1-3ml

Take two consecutive sets from two separate venepuncture sites during any 24hr period for each septic episode. For neonates, take a single aerobic bottle or special low volume bottle.

Take two sets during the first hour in cases of severe sepsis prior to commencing antibiotic treatment, provided this does not significantly delay antibiotic administration.

If 1 set is catheter drawn, draw at least one set for a peripheral vein.

Take at least three sets during a 24hr period where the patient has suspected infective endocarditis.

Specimens should be transported and processed as soon as possible. If pathology reception is closed the bottles should be placed into the incubator in the Pathology Specimen reception room (under bench next to the blood issue fridge).

Samples should not be refrigerated.

|  |  |
| --- | --- |
| **Procedures:** | **Notes:** |
| * Two sets needed to evaluate sepsis. * 8-10mL of blood in each culture bottle for adults   1-3 mL in paediatric culture bottle |  |
| **Assemble supplies:**   * Bottles, * Blood culture collection pack (available from Hospital stores), * Disposable tourniquet, * Vacutainer butterfly needles, Chlorhexidine/alcohol skin cleanser (chloraprep sepp/frepp), * Sanicloth for bottle tops |  |
| * **Hand hygiene:** * Wash hands prior to donning gloves before drawing cultures. * Use alcohol-based hand sanitizer (allow to dry). | Hand hygiene is proven to reduce spread of infection and blood culture contamination. |
| **Prepping Skin: Peripheral Cultures**   * Select site of venepuncture: cleanse with soap and water if unusually dirty. * Wear non-sterile examination gloves. * Apply tourniquet, if necessary. * Cleanse venepuncture site with Chloraprep 1.5mls (Frepp) * Cleanse skin as per Chloraprep guidance * Allow air drying (20 seconds). | * Reduce chance of false positives by reducing potential for sample to contact organisms on patient skin, central lines, or transferred from operator. * Do not blow or fan to speed drying site, can contaminate the cleansed area. Air drying can occur while culture bottles are being prepared. * Allows times for alcohol to act and avoids stinging from alcohol at site. |
| **Prepare Culture Bottles**   * Flip off plastic lid. * Cleanse each rubber top with Chloraprep Sanicloth * Ensure chlorhexidine/alcohol has evaporated before inoculation. | * Ensure sterile access to culture medium. |
| **Drawing/Transfer of Peripheral Site Blood Cultures (Recommended)**   * Apply tourniquet if not already applied. Avoid contamination of prepped area with gloves or tourniquet. After cleansing site: cleanse gloved finger thoroughly with alcohol if it is necessary to touch venepuncture site. * Use a needle and syringe only where vacutainer system is unavailable. * Use single use adaptor for each patient when using vacutainer system. * Perform phlebotomy: release tourniquet and withdraw needle. Apply pressure to site and bandage/non-allergenic tape * Innoculate blood culture bottles: (aerobic bottle first (blue top) in the case of a set). * To avoid Needlestick injury, discard used sharps immediately into a designated sharps bin which has a device for needle removal. Transfer blood from syringe to blood culture bottles using a new sterile needle attached to sample syringe. | * Reduce chance of transfer of contamination organisms from needle or butterfly into culture bottles by removing: may reduce false positives by half from this precaution alone. * Reduce chance of transfer of organisms from gloved finger to clean site. * Inoculate aerobic bottle first to avoid air entry to the anaerobic bottle. |
| * Affix patient label to each bottle with patient’s name, hospital number, date and time of collection and send to lab using standard protocol for specimen submittal. * Blood cultures should be taken to Microbiology department within an hour of collection or sent in the pneumatic tube system. * Blood Cultures collected outside normal hours should be delivered and placed in the incubator next to the Blood Bank collection refrigerator located in the Pathology specimen reception room. | * Assure right patient, right test. * Samples will be rejected if incompletely labelled. * To prevent breakage of specimen and for optimal incubation. |

### Bone marrow cultures

For Mycobacteria,Typhoid and Brucella. These are usually unrewarding, discuss with Consultant Microbiologist before collection.

Use aseptic technique.

Bone marrow specimens should be submitted in sterile white topped universal containers sealed inside plastic bags for transport.

Bone marrow specimens for TB culture should be inoculated into Lithium Heparin vacutainer blood bottles without gel (dark green cap). A volume of at least 2ml is required. These will be referred to the Mycobacterium reference laboratory for TB culture.

### Bronchoscopic samples

If possible collect samples before antimicrobial therapy is started.

Biopsy samples: bring upright in 0.5 mL saline in a sterile universal container or sputum container to laboratory.

BAL samples: send in sterile universal container.

Brush samples: send brush in sterile universal container.

To ensure urgent processing please contact the laboratory before sending these samples.

### Catheters

Do not send urinary catheters, these are unsuitable for microbiological analysis. Send Mid-stream or Catheter Urine instead (see section 20). Urinary catheters will not be tested.

### Cerebrospinal fluid (CSF)

For microscopy, culture and examination for Xanthochromia

Collect 4 specimens into sterile plain (white topped) universal containers, numbering the containers consecutively:

* + First (label **1**) 1 mL
  + Second (label **2**) 1 mL
  + Third (label **3**) 1 mL
  + Fourth (label **4**) 1 mL

Send samples **1** and **3** for Microbiology investigation

Sample **2** should be sent for Biochemical analysis (protein and glucose)

Sample **4** should be sent to Biochemistry for Xanthochromia investigation (protect from light)

For *Mycobacterium* species (TB investigation)*,* collect at least 10mL into tube 3 where possible.

Minimum sample volumes required for additional CSF tests:

TB PCR: 0.5mL

CSF PCR panel (Viral, Bacterial and Fungal): 0.2mL

16s PCR: 0.2mL

Cryptococcal antigen testing: 0.1mL

Cryptococcal PCR: 0.2mL

For paediatric samples the volumes listed above are preferred, however if the volume collected is insufficient for all tests we will liaise with the Consultant Microbiologist and requesting Clinician to decide which tests are prioritised.

**Microbiology CSF samples must reach the laboratory within an hour.**

Remember to take a simultaneous serum sample to measure glucose concentration.

For cytology (malignancies) send a minimum of 0.5 mL directly to Cellular Pathology unless the patient is known to have leukaemia or lymphoma, in which case the Haematology department should be consulted.

### Chlamydia trachomatis/Neisseria gonorrhoeae CT/NG testing:

Neisseria gonorrhoea PCR can be carried out simultaneously with Chlamydia PCR on the same sample. Request the combined Chlamydia/Gonorrhoea PCR test. If requesting Neisseria gonorrhoeae PCR please also send a cervical or urethral swab for MC&S.

**In women:**

i) Endocervical swab remains the best sample

ii) self- taken, low vaginal swabs are acceptable

The testing of first catch urine specimens from women may result in lower sensitivity and is not recommended by national or European guidelines. In women, urine is not the optimal sample for *N. gonorrhoeae/C. trachomatis* combined NAATs.

**In men:**

i) “First void” urine sample is the preferred sample

**Sample Stability**

Within the Roche Cobas PCR sample tube both swabs and urines are stable at 2-30oC for 12 months. Urines not in the Roche Cobas PCR sample tube will be rejected as the quality of the specimen cannot be assured.

**Sample Collection**

**Endocervical swabs:**

Handling precautions for Cobas PCR Female Swab collection: **Do NOT** pre-wet collection swabs with the collection media before obtaining the endocervical specimen. Use care to avoid splashing of contents.

* Label the Cobas tube with the patient’s details.
* Using one of the swabs provided in the Cobas PCR Female Swab Sample packet, remove excess mucus from the cervical os and surrounding mucosa. Discard this swab after use OR take bacteriology swab(s) first which will effectively clean the neck of the cervix of pus and mucus.
* To collect the Chlamydia specimen, insert the other provided swab into the endocervical canal. Gently rotate the swab 5 times in one direction in the endocervical canal. Do not over rotate. Carefully withdraw the swab, avoiding any contact with the vaginal mucosa.
* Remove the cap from the Cobas PCR media tube and lower the swab specimen into the tube until the visible dark line on the swab shaft is aligned with the tube rim. The tip of the swab should be just above the media surface near the hexagonal Roche logo.
* Carefully leverage the swab against the tube rim to break the swab shaft at the dark line; discard the top portion of the swab.
* Tightly re-cap the Cobas PCR media tube and swirl the tube 3 to 5 times. The specimen is now ready for transport.
* Ensure that the tube is labelled with the Patient’s details, swab site and date and time of specimen.

**NOTE:** Swab samples with frank bloodstaining will not be tested, as the presence of blood can affect the result.

**Self-taken, low vaginal swabs:**

* Label the Cobas tube with the patient’s details
* Do NOT dip the swab into the tube before the test.
* Insert the swab into the vagina, as if inserting a tampon. Twirl the swab for approximately 5 seconds, ensuring it touches the inside of the vagina.
* Remove the swab and snap off into the Cobas PCR medium tube. Screw cap firmly.
* Ensure that the tube is labeled with Patient’s details, swab site, date and time of collection. Transport to the laboratory.

Instructions for collection appear on the sample collection kit pack.

**NOTE:** Swab samples with frank bloodstaining will not be tested, as the presence of blood can affect the result.

**First void urine:**

The patient should not have passed urine in the previous 1 hour

* Label a Cobas PCR media tube with the Patients details.
* Collect the **first 10-25 mL** of the stream into a sterile universal container (the label is graduated, to give a guide to collection volume). **This is NOT a mid-stream urine sample.**
* Transfer the urine into the Cobas PCR media tube (yellow cap) using the disposable pipette to add urine to between the two black fill lines on the tube.

Take care to avoid splashing of contents

* Tightly recap the tube. Check that the tube is labelled with patient information plus date and time of specimen.
* Mix by inverting the tube 5 times. The specimen is now ready for transport
* If the urine specimen cannot be transferred immediately it can be stored at

2º C -30ºC for up to 24 hours. It must be transferred prior to sending to the

laboratory to maintain the integrity of the sample. Samples received in the

laboratory which are not in the Roche Cobas PCR tube will not be tested.

Instructions for urine collection appear on the sample collection pack and are available in poster format from Microbiology.

**NOTE:** Samples which are NOT filled to between the two black lines will not be tested, since the result may be unreliable. Please stress to the patient the importance of filling the tube to the correct level (between the two fill lines).

### Drains

Send drainage fluid in a plain (white topped) sterile universal container rather than drain tips. Transport to the laboratory as soon as possible.

### Faeces

Collect into a sterile blue capped universal container. Minimum required for routine culture is 1-2g (approx. 1-2mL); but please note that for *C. difficile* testing samples less than 4mL will not be tested.

Please ensure that all details are on both the specimen and accompanying request form

The following methods can be used to collect a specimen:

The patient or carer should wear disposable gloves

Contamination with urine should be avoided

* Toilet paper can be crumpled into the toilet bowl or suspended across the toilet bowl in a cross to make a sling.
* A clean plastic container can be positioned in the toilet bowl
* Cling film can be stretched across the top of the toilet bowl

A portion of faeces can then be collected with a wooden tongue depressor or the

spoon provided in the specimen pot and transferred to the specimen container

The specimen pot should then be sealed into the specimen bag.

All materials should be placed in a plastic bag which is sealed before disposal in

the refuse bin.

Up to three samples collected on different days may be necessary in order to exclude bacterial infections. Repeated sampling over longer periods may be necessary if *Giardiasis* is suspected, as intermittent excretion of cysts is not uncommon.

Routine specimens are cultured for *Salmonella spp., Shigella spp., Campylobacter spp.* and *E. coli O157*. A routine screening test is also applied for *Cryptosporidium* *spp*. if the sample is diarrhoeal. Other investigations will only be performed if indicated by the clinical history, erg. Rotavirus antigen, *Vibrio* culture.

For in patients, routine culture will only be performed if the patient was admitted 3 days ago or less. For patients who have been inpatients for 4 days or more the first sample from each admission will be tested and subsequent samples will not be tested.

**Microscopy for parasites**: The examination for Ova, Cysts and parasites will only be performed on samples with the appropriate clinical or travel history.

Ideally three stool specimens collected over no more than a 10-day period. It is recommended that specimens are collected every other day. Unless the patient has severe diarrhoea or dysentery, no more than one specimen should be examined within a single 24 hour period, as shedding of cysts and ova tends to be intermittent.

If *E. histolytica* or *G. duodenalis* are suspected and the first three specimens are negative, ideally three additional specimens should be submitted at weekly intervals.A 'sellotape preparation' using a pinworm collection device should be sent for threadworm investigations. Instructions for use are issued with each device. It is recommended that samples should be taken for at least four to six consecutive days. If the results of all these are negative the patient can be considered free from infection.

**Pseudomembranous colitis/*Clostridium difficile* toxin A and B**

Please ensure that you request *C. difficile* toxin where clinically indicated.

Samples will be tested a maximum of two times per episode of diarrhoea.

Samples taken less than 28 days after a positive *C. difficile* toxin test will not be processed.

A minimum of 4ml faeces must be sent for *C. difficile* testing. Smaller samples will not be tested.

Please discuss with the Consultant Microbiologist before sending samples for C. difficile on patients who are on NG or PEG feeds or have received laxative/aperients/enema or bowel prep in the last 24 hours.

### Lines and shunts

Line tips e.g. CVP or Hickman lines, swabs of cannula insertion sites:

Collect specimens before starting antimicrobial therapy where possible.

Use aseptic technique.

**Cannulae**

Disinfect the skin around the cannula entry site, remove cannula using aseptic technique.

Cut off 4cm of the tip into a sterile white topped universal container using sterile scissors.

Place in sealed plastic bags for transport.

Cannulae should only be sent if there is evidence of infection. Swabs and blood cultures should also be submitted:

**Swabs**

Sample the inflamed area / exudate around the catheter insertion site using an Amies swab with charcoal medium.

**Blood Cultures**

At least two blood cultures should be obtained when catheter infection is suspected. One set should be collected by peripheral venepuncture and one through the vascular catheter.

Paediatric blood culture bottle - blood volume 0.5-5mL, ideal volume 1-3mL

### MRSA

Sample collection: Use swabs with Amies charcoal transport medium. For sampling of skin surfaces, nose, and groin - sample collectors should moisten swabs in sterile water prior to swabbing skin surfaces. Take the moistened swab and roll it at the entrance of both sides of the nose, using the same swab for both nostrils. Return the swab to the transport tube. Send to the laboratory with a completed request form.

**Patients:**

* **Samples always collected:** Nose/ groin swabs (plus axilla and perineum swabs in neonates)
* **Samples collected where applicable:** Wounds, other skin lesions, insertion of IV catheters, tracheostomies, catheter urines and sputum.
* **Samples collected from patients with persistent nasal carriage:** Throat.

**Staff:**

* **Samples always collected:** Nose
* **Samples collected where appropriate:** Lesions, sites of abnormal skin, others as decided by the Infection Prevention and Control Team.

**Follow up of new positive cases/screening of previous positives/transfers from other hospitals:**

* **Samples always collected**: Nose/axilla/groin
* **Samples collected where appropriate**: Wounds, other skin lesions, insertion sites of IV catheters, tracheostomies, catheter urines and sputum.

**MRSA Rapid Screening (approved locations only)**

Department of Critical Care (DOCC), Neonatal Unit and other locations approved by the Infection Prevention and Control Team and the Consultant Microbiologists may submit nasal swabs for rapid MRSA screening by PCR.

This method of screening is appropriate only on admission to the requesting location, and only for patients not known to be previously MRSA positive. It is not appropriate as a test of cure.

The sample required is a red-topped double Copan nasal swab (red top). Each Copan swab has two individual swabs affixed to the lid. Do not detach these but treat as one unit when sampling. Moisten the swab in sterile water.

Take the moistened swab and roll it at the entrance of both sides of the nose, using the same swab for both nostrils. Return the swab to the transport tube.

The laboratory must be informed that the swab has been taken, in order for it to be processed urgently. After 5pm on weekdays and any time at weekends the on call Microbiology BMS should be contacted via the switchboard. Results will be available on eCare/ICE within 2 hours.

MRSA PCR samples received after 9pm will be held for processing the following morning.

### Screening for CPE (Carbapenemase producing Enterobacteriaceae)

Indications for screening are hospital treatment abroad, hospitalisation in an area known to have cases of CPE, previous CPE or close contact with a case.

**For further guidance refer to the Infection Prevention and Control pages on the Trust Intranet.**

* Rectal Swab- (standard bacteriological swab) should be taken with visible material identified. If it is not possible to obtain a rectal swab, a stool sample can be sent.

All specimens should be labelled on the form clearly as ‘**possible CPE colonisation or infection’**

### Screening for ESBL (Extended Spectrum Beta-Lactamase producers)

ESBL producing Enterobacteriaceae may be isolated from various specimens during routine processing.

Screening for ESBL producers is carried out for patients being repatriated to Neo-natal unit or as requested by the Consultant Microbiologist/Infection Control and Prevention team. **For further guidance refer to the Infection Prevention and Control pages on the Trust Intranet.**

### Mycology

Special specimen collection packs (Dermapak) are available from the laboratory (ask for Dermapak).

**Skin**

Patients’ skin and nails can be swabbed with 70% alcohol prior to collection of the specimen, this is especially important if creams, lotions or powders have been applied. The edges of skin lesions yield the greatest quantities of viable fungus. Lesions should be scraped with a blunt scalpel blade, collecting scrapings into a Dermapak. Label with the patient details and the sample site. Seal in a plastic bag for transport to the laboratory. If insufficient material can be obtained by scraping and being placed in a container, then a swab or sticky tape can be pressed on the lesion and transferred to a clean glass slide for transport to the laboratory (‘stripping’).

**Nail**

It should be specified whether the sample is from the fingernails or toenails. Material should be taken from any discoloured, dystrophic or brittle parts of the nail. The affected nail should be cut as far back as possible through the entire thickness and should include any crumbly material. Nail drills, scalpels and nail elevators may be helpful but must be sterilized between patients. When there is superficial involvement (as in white superficial onychomycosis) nail scrapings may be taken with a curette. If associated skin lesions are present samples from these are likely to be infected with the same organism and are more likely to give a positive culture. Sample from associated sites should be sent in separate packets. Label each packet clearly with the sample site. Place in sealed plastic bags for transport to the laboratory.

**Hair**

Samples from the scalp should include skin scales and hair stumps. Cut hairs are not suitable for direct examination as the infected area is usually close to the scalp surface. Scraping for direct examination is the preferable sample collection method, however plastic hairbrushes, scalp massage pads, swabs or plastic toothbrushes may be used to sample scalps for culture where there is little obvious scaling. If sufficiently long, hairs should be plucked with forceps and wrapped in black paper or commercial transport packs together with flakes of skin. Place the collection pack into a sealed plastic bag for transport to the laboratory.

Results of direct films will normally be reported within 48 hours, but final culture results will be issued when positive or after four weeks.

**Blood**

Blood samples may be referred for Serum-Beta-Glucan or other fungal serology. See serology section.

### Sputum

Sputa samples must be sent in sterile 60ml screw top sputum containers. Place in the bag attached to the request form with ‘DANGER OF INFECTION’ labels applied to specimen and form. The value of this specimen depends critically on the care taken to collect it. Do not send saliva. Salivary samples will not be tested.

Indicate if the sample is a suction (endotracheal) sample. Make sure the lid or suction trap is firmly sealed.

Culture for TB will only be carried out if this is specifically indicated on the request form.

Early morning freshly expectorated sputum is recommended for *Mycobacterium* species.

For sputum specimens the material required is from the lower respiratory tract, expectorated by deep coughing. When the cough is dry, physiotherapy, postural drainage or inhalation of an aerosol before expectoration may be helpful. Saliva and pernasal secretions are not suitable.

Ideally, a minimum volume of at least 1mL is required.

Sputum samples can be referred for *Pneumocystis jirovecii* (PCP) PCR if indicated. Please send a separate sample for this test.

### Swabs

Collect specimens before antimicrobial therapy where possible.

For swabs which cannot be sent to the laboratory immediately, refrigeration is preferable to room temperature storage.

Where possible send a specimen of pus in a sterile white topped universal container rather than a swab.

Moisten swabs before taking from dry areas of skin (if these need swabbing) with sterile water.

**ENT swabs**

Use Amies with Charcoal (black) swabs.

**Genital tract swabs**

Use Amies with Charcoal (black) swabs.

Ideally, inoculation of specimens for *N. gonorrhoeae* is made directly to culture media at the bedside (at the sexual health clinic) and incubated without delay.

Transport time should be as short as possible.

**High vaginal swabs (HVS)**

HVS is not recommended in routine cases as it is unlikely to be of significant diagnostic value.

The situations where an HVS is indicated are:

· Postnatal infection

· Pre & post termination of pregnancy

· Pre & post-operative gynaecological surgery

· Acute presentation to emergency gynaecology or persistent/recurrent symptoms

· Symptoms not characteristic of candida or BV

After the introduction of the speculum, the swab should be rolled firmly over the surface of the vaginal vault. The swab should then be placed in Amies transport medium with charcoal.

HVS is suitable for diagnosis of *Trichomonas vaginalis,* Yeasts and some bacterial infections. All HVS are screened for clue cells, which may indicate bacterial vaginosis. Further cultures may be put up if indicated by the clinical details. HVS is unsuitable for diagnosis of *Neisseria gonorrhoeae* infection.

**Cervical swabs**

After introduction of the speculum to the vagina, the swab should be rotated inside the endocervix. The swab should then be placed in Amies transport medium with charcoal.

Cervical swabs will be tested for *Neisseria gonorrhoeae*. Further cultures will be put up if indicated by the clinical details or if no HVS has been received with the cervical swab.

**Urethral swabs**

Contamination with micro-organisms from the vulva or the foreskin should be avoided.

Thin swabs are available for collection of specimens.

The patient should not have passed urine for at least one hour. For males, if a discharge is not apparent, attempts should be made to "milk" exudate from the penis. The swab is gently passed through the urethral meatus and rotated. Place the swab in Amies transport medium with charcoal.

Urethral swabs are suitable for diagnosis of *Neisseria gonorrhoeae*, and Yeasts.

**Rectal swabs**

Rectal swabs are taken via a proctoscope. Use swabs with Amies with charcoal transport medium.

**Throat swabs**

Throat swabs should be taken from the tonsillar area and/or posterior pharynx avoiding the tongue and uvula.

**Wound/Ulcer/Skin swabs**

Use swabs with Amies charcoal transport medium.

Note that all ulcers are invariably colonised by a polymicrobial flora and swabs should only be taken if a clinical diagnosis of infection has been made.

For **Chlamydia PCR** see section 8.

1. **Intrauterine contraceptive devices (IUCDs)**

The entire device should be sent in a sterile, white topped universal container.

1. **Fluids and pus**

These are taken from the fallopian tubes, tubo-ovarian and Bartholin’s abscesses, etc. during surgery. Minimum volume 1mL. Collect into a sterile white topped universal. Specimens should be transported and processed as soon as possible. If processing is delayed, refrigeration is preferable to storage at ambient temperature.

1. **Pernasal and postnasal swabs for whooping cough** – use pernasal swab with flexible ultrafine wire shaft Amies charcoal transport medium. Collect before antimicrobial therapy where possible.

A pernasal swab is inserted through a nostril and advanced along the floor of the nose until it reaches the nasopharynx. It has been suggested that the swab is held against the posterior nasopharynx for up to 30s or until the patient coughs. In practice, it is more likely that a patient will only be able to tolerate this for a few seconds.

1. **Swabs for Viral investigation:**

**CMV PCR Swab**

Saliva swab in viral transport medium

**Herpes simplex**

Send swab from lesions in viral transport medium red capped tube with red liquid transport medium.

### Nasopharyngeal aspirate or swab for Respiratory Virus Detection

**Nasopharyngeal aspirate**

Aspirate into suction trap. Remove the suction tubing and replace with the screw cap. Place into a plastic bag and transport to the laboratory as soon as possible. Please note that a nasopharyngeal swab is the preferred sample for RSV testing.

**Nasopharyngeal swab**

Using a GeneXpert nasopharyngeal virus collection kit, swab the nasopharynx.

Aseptically remove the cap from the tube of transport medium.

Insert the swab into the tube.

Break the swab shaft by bending it against the tube wall.

Replace the cap on the tube and close tightly.

Label with appropriate patient information.

Place into a plastic bag and transport to the laboratory.

A rapid PCR test for RSV, Flu A and Flu B is performed in house.

An extended virus panel is available in selected cases. Please discuss with Consultant Microbiologist before sending samples.

### SARs-CoV-2 PCR

Screening and diagnostic samples are referred for testing at the John Radcliffe Hospital, Oxford.

Sample type is a combined nose and throat swab in virus transport medium, in a grip bag with absorbent pad, inside a Biohazard bag.

Use a the swab supplied to sample the throat, and then using the same swab sample the nose. Full instructions for collecting samples are available on the Trust Intranet

SARs-CoV-2 rapid testing on site is available for selected cases and may be requested by the Duty Manager or DOCC Consultant.

### Urines

Collect specimens before antimicrobial therapy where possible.

Numbers and frequency of specimen collection are dependent on clinical condition of patient

Fill to the line marked on containers with boric acid preservative according to manufacturers’ instructions.

For urine samples in plain white topped universal containers a minimum volume of 1mL is required.

Containers and Kits:

All routine urine samples for Microscopy Culture and Susceptibility (MC&S) testing from patients over 5 years old should be collected into 10mL containers with boric acid, CE marked leak proof containers, filled to the line on the container and transported as soon as possible to the laboratory, in sealed plastic bags with the request form.

These containers are issued as a kit including plastic collection cup to aid filling the tube and avoid contamination of the outside of the container. For paediatric patients, 10mL boric acid containers are available. These must be filled to the line marked on the container.

Underfilled boric acid urine containers will not be tested.

White top universal will ONLY be accepted on patients <5 years, who are unable to produce 10mL urine. Please ensure they reach the laboratory promptly, >4hours at room temperature may affect the quality of the results. If transport likely to be greater than 4 hours, the sample can be refrigerated up to 48 hours.

Boric acid is a preservative (white powder) that maintains the integrity of the sample up to 96 hours at room temperature. By preserving the white blood cells, reducing the numbers of mixed and false positives, which in turn improves quality of patient care by reducing unnecessary antibiotic treatment. In the long term reduced antibiotic usage, will help to minimise the development of antibiotic resistance in the bacteria population of the hospital and local community.

Specimens NOT received in Boric acid and /or > 96 hours will be rejected unless they are a specimen type (detailed below) which can be collected in white topped containers. These specimens should reach the laboratory within **one** **hour** unless refrigerated, to ensure they are processed within a 4 hour window from collection to processing. If delayed or collected between 20:45 and 08.45 specimens may be refrigerated for up to 48 hours before processing.

Note: boric acid may affect the results given by some brands of urinary dip sticks.

It is important that urines are collected in a manner which minimises contamination

(aseptically) as this avoids 'difficult to interpret' results. Within the Trust, sterile Midstream Urine (MSU) collection containers are available from HSDU, in the community urines can be collected direct into the boric acid container or white topped container, or similar sterile vessel, as appropriate to testing.

**Urine Microscopy, Culture & Susceptibility**

Urine samples will be cultured only if this is indicated by the microscopy result or clinical information provided, for example pregnant or paediatric samples. If culture has been omitted a report is released with a comment ‘if culture still indicated based on clinical condition, please contact the laboratory as soon as possible. Provide the sample is in boric acid and < 96hours, the specimen can be cultured.’

**Mid-stream urine (MSU)**

MSU is the recommended routine collection method. A detailed Patient Information Leaflet is available on the Trust intranet; ‘How to collect a urine sample’ or can be supplied by the laboratory.

Periurethral cleaning is recommended (water is considered sufficient).

Mid-stream urines should be collected as follows:

(a) Carefully wash the external genitalia and dry with a clean towel.

(b) Start passing urine, allowing the first part to flow into the pan.

(c) Collect the next part of the specimen into the container (women should separate the labia with the fingers of the hand which is not holding the container)

(d) Screw the lid on firmly and label.

The first part of voided urine is discarded and, then without interrupting the flow, 20 mL is collected into a red topped universal container with boric acid. The remaining urine is discarded into the toilet. If boric acid preservative is used, white powder, the container is filled up to the mark in a similar manner and the contents mixed well.

**Clean-catch urine**

A reasonable alternative to MSU.

Periurethral cleaning is recommended. The whole specimen is collected and then a 20mL aliquot sent for examination in a red topped universal container with boric acid.

**Suprapubic aspirate (SPA)**

Minimum volume 1mL

Urine is obtained aseptically, directly from the bladder by aspiration with a needle and syringe. The use of this invasive procedure is usually reserved for clarification of equivocal results from voided urine (e.g. in infants and small children). Ultrasound guidance should be used to show presence of urine in the bladder before carrying out SPA. Use a white topped sterile universal and transport to the laboratory within one hour, mark urgent.

**Catheter urine (CSU)**

Minimum sample volume 20mL (fill to the line).

The sample may be obtained either from a transient (‘in and out’) catheterisation or from an indwelling catheter. In the latter case, the specimen is obtained aseptically from a sample port in the catheter tubing or by aseptic aspiration of the tubing. The specimen should not be obtained from the collection bag. Transfer the sample into a red topped universal container with boric acid.

**Bag urine**

Minimum volume 1mL

Used commonly for infants and young children. The sterile bags are taped over the freshly cleaned and dried genitalia, and the collected urine is transferred to a sterile universal container. There are frequent problems of contamination with this method of collection.

**Pad urine**

Minimum volume 1mL

An alternative to collecting bag urine from infants and young children. After washing the nappy area thoroughly, a pad is placed inside the nappy. As soon as the pad is wet with urine (but no faecal soiling), push the tip of a syringe into the pad and draw urine into the syringe. Transfer specimen to a sterile universal container. If difficulty is experienced in withdrawing urine, the wet fibres may be inserted into the syringe barrel and the urine squeezed directly into the container with the syringe plunger.

**Ileal conduit – urostomy urine**

Minimum sample volume 20mL (fill to the line).

Urine is obtained via a catheter passed aseptically into the stomal opening after removal of the external appliance. Results from this type of specimen may be difficult to interpret.

Collect the sample into a red topped container with boric acid.

**Cystoscopy urine**

Minimum volume 1mL

Urine is obtained directly from the bladder using a cystoscope. Use a white topped sterile universal and transport to the laboratory and mark urgent

**Ureteric urine**

Minimum volume 1mL

Paired urine samples are obtained from each ureter during cystoscopy via ureteric catheters inserted from the bladder.

Urine samples may also be sent following nephrostomy, other surgical procedures, or bladder washout. Use a white topped sterile universal and transport to the laboratory within 1 hour and mark urgent

**Other specimens**

Minimum volume 1mL

Other specimens obtained during or as a result of surgery include those from ileal conduit, cystoscopy, nephrostomy and urostomy, prosthetic massage/secretions. Specimens may also be taken after bladder washout. Collect into a white topped container and transport to the laboratory within 2 hours.

**Urine for *Salmonella typhi* and *Salmonella paratyphi* cultures**

Minimum volume 1mL

Any urine samples from suspected cases or contacts of cases. Label the sample and form with danger of infection stickers, and give relevant clinical details including travel history. Collect the sample into a white topped universal container.

**Early morning urine for TB culture**

Three entire, first voided, early morning urine specimens are required for culture for   
*M. tuberculosis.* Special containers are available from Pathology Supplies upon request. Label the sample and form with a danger of infection sticker.

**Legionella antigen:**

Minimum volume 1mL in sterile white topped universal container or 20mL in red topped container (fill to line).

Detection of Legionella antigen in urine can confirm the diagnosis of Legionellosis. Urine should be sent in a sterile white topped universal container. Relevant clinical information including date of onset should be provided. Collect in either a sterile white topped universal container or a boric acid container, filled to line.

The samples can be stored at room temperature (15-30°C) if assayed within 24 hours of collection.

Alternatively, specimens may be stored at 2-8°C for up to 14 days or at -10°C to -20°C for longer periods before testing.

**Pneumococcal antigen**

Minimum volume 1mL in sterile white topped universal container or 20mL in red topped container (fill to line).

Relevant clinical information and date of onset must be provided. Collect in a sterile white topped universal container or boric acid filled to line

Store at room temperature (, 15-30°C) if assayed within 24 hours of collection. Alternatively, store urine at 2-8°C or frozen for up to 14 days before testing.

**Urine for *Schistosoma haematobium* detection**

Minimum volume 10mL in sterile white topped universal container.

Total urine collected between 10am and 2pm (period of maximum activity) is the preferred specimen, alternatively a 24 hour collection of terminal urine may be obtained.

It is preferable to obtain total urine collected over the time period between 10am and 2pm as it has been shown that a maximum concentration of eggs is excreted at this time. Sterile containers without boric acid must be used. In patients with haematuria eggs may be found trapped in the blood and mucus in the terminal portion of the urine specimen.

For terminal urine collection, collect the last 10-20mL of urine each time of urination within one 24 hour period.

### Virus detection

VIRAL (Bacterial and Fungal also) PCR can be carried out on CSF samples collected into a sterile white topped universal container.

For herpes simplex PCR send swabs from lesions in Virus transport medium (red capped tube with red liquid transport medium).

Nasopharyngeal aspirate can be submitted for respiratory virus PCR.

Nose and throat swabs may be submitted for respiratory virus PCR.

CMV PCR can be carried out on urine or virus swabs.

**Salivary swabs for CMV PCR (neonates and infants only).**

Use a virus transport swab. Place the swab inside the baby’s mouth between the lower gum and cheek for at least a minute. Once the tip is soaked place the swab in a tube with universal or virus transport medium (red capped tube with red liquid transport medium). If the baby is breast fed, then the swab should be taken at least 1 hour after the feed”

1. **In House Microbiology Tests:**

| **Test:** | **Sample:** | **Container:** | **Turnaround:** | **Notes:** |
| --- | --- | --- | --- | --- |
| AFB (TB) Culture | Sputum/other | Sputum pot or 150ml urine pot | 3-8 weeks |  |
| AFB Microscopy | Sputum/other | Sputum pot | 1 working day | Positive results will be telephoned to Ward or clinician as soon as available |
| \*Blood Culture | Blood Culture | Blood culture | 5-7 days | Microscopy result will be telephoned to Ward or clinician as soon as available |
| Bordetella pertussis culture | Pernasal swab | Amies wire shaft charcoal swab | 10 days |  |
| C difficile toxin | Faeces | Blue top universal | <24 hours | Positive results will be telephoned to Ward as soon as available |
| Chlamydia/Gonorrhoeae PCR | Genital swab/urine | Roche CT/NG collection kit | 5 days |  |
| Covid-19 (Novel Coronavirus PCR) – selected cases only | Combined nose and throat swab | Virus transport medium | 2-4 hours subject to analyser availability | Telephone laboratory |
| CSF MC&S | CSF | Sterile White top Universal | Microscopy within 2 hours of sample  collection  Culture 16-72 hours | Telephone laboratory when sending.  Microscopy result will be telephoned to ward or clinician as soon as available |
| CSF PCR Panel, Viral Bacterial and Fungal | CSF | Sterile plain universal | 2 hrs | Telephone laboratory |
| Dermatophyte culture (Mycology) | Skin/Nail | Dermapak | 10-30 days |  |
| Flu A/B and RSV PCR | NP Swab | GeneXpert collection tube | <24 hours | Telephone laboratory |
| Legionella antigen | Urine | White topped universal | <24 hours |  |
| MRSA culture | Nose/other swab | Amies charcoal swab | 2-5 days |  |
| MRSA PCR | Nose swab | Red top Copan swab | 2 hours if notified urgently between 9am-9pm | Notify the laboratory when sending |
| Norovirus PCR | Faeces | Blue top universal | <24 hours |  |
| Pneumococcal antigen | Urine | White topped universal | <24 hours |  |
| RSV and Flu A/B PCR | NPA or NP Swab | Suction trap or GeneXpert collection tube | <24 hours | Telephone laboratory |
| Extended Respiratory Virus Panel | NPA or NP swab, CNT swab | Swab in Virus transport | <24 hours | Telephone laboratory |
| Faeces culture | Faeces | Blue top universal with spoon | 2-5 days |  |
| Ova, Cysts and Parasites | Faeces | Blue top universal with spoon | 2-5 days |  |
| Cryptosporidium/Giardia EIA | Faeces | Blue top universal with spoon | 2-5 days |  |
| Sputum culture | Sputum/BAL | Sputum pot | 2-5 days | 7 days if Burkholderia cepacia culture is indicated |
| Swab culture | Swab | Amies charcoal swab | 5 days | Longer if actinomyces or Fusobacterium culture is indicated |
| Urine Culture | Urine | Boric acid (red top) universal or 10mL tube | 2-5 days |  |
| Urine Microscopy | Urine | Boric acid (red top) universal or 10mL tube | <24 hours |  |

*\*Blood cultures are incubated for 5 days before being reported as negative. When a blood culture becomes positive the laboratory will telephone the requesting location (in core laboratory hours) or the appropriate Clinician (outside core laboratory hours) as soon as possible, with details of organisms seen.*

*An interim report will be issued showing the microscopy result. The culture result, detailing any isolate with appropriate sensitivities will be sent as soon as available, normally 1-2 days later. The Consultant Microbiologists can advise on appropriate therapy pending the final report.*

**SAMPLE TYPE CODES**

BAL Bronchioalveolar lavage

BM Bone Marrow

CB Clotted blood. Gold cap.

CSF Cerebrospinal fluid

EDTA EDTA blood. Purple cap.

F Faeces

GBX Gastric Biopsy

JASP Joint Aspirate

LH Plain lithium heparin (no gel). Dark green cap

SA Spleen aspirate

SKB Skin biopsy

SP Sputum

SW Swab

TIS Tissue

U Urine

VS Viral Swab

### Viral and other infectious disease serology

All antibody tests require a 5 mL clotted blood tube.

Give as much relevant clinical history as possible, including the date of onset, details of any contact with known or suspected cases.

**Ante-Natal Patient with rash illness/in contact with rash illness**

Sera from pregnant women with rash/in contact with rash should be tested for Parvovirus IgG and IgM, and also for Rubella IgM and IgG is there is no clear history of past infection or full immunisation.

If there is contact with measles, immunity can also be tested, ideally within 6 days of the contact.

Varicella Zoster:

ICE request: VZV Contact Antenatal

eCARE request: VZV Antenatal Serology, blood

Parvovirus:

ICE request: Parvovirus Contact Antenatal

eCARE request: Parvovirus Antenatal Ab, blood

Rubella (patient with rash):

ICE request: Rubella ?infection

eCARE request: Rubella infection, blood

Rubella (contact):

ICE request: Rubella immunity

eCARE request: Rubella immunity, blood

Measles (immunity):

ICE request Measles immunity

eCARE request: Measles immunity, blood

Where a stored sample exists, the tests will be carried out on stored booking blood to demonstrate immunity prior to the contact or onset of rash. The current sample will also be tested if indicated.

Information required from the requester:

* Gestation of pregnancy (LMP date)
* Clinical features (type and distribution of rash, joint pain, lymphadenopathy)
* Date of onset
* History of rubella immunisation/antibody testing; date, place and results of testing
* History of contact with rash illness, type and date of contact

**Routine infectious diseases in pregnancy (antenatal) screening**

Samples should be collected into 5mL gold topped tubes.

The sample should normally be collected at the booking appointment.

Manual requests use the combined request form For Infectious Diseases, Sickle Cell and Thalassaemia in Pregnancy

ICE or eCARE: request Antenatal Infectious Disease Screening

Samples will be tested for HIV, Hepatitis B surface antigen and Syphilis antibodies unless declined by the patient.

Accept/Decline response for each test is captured on the hand-written request form or on eCare/ICE, along with known positive or unknown status for each infection.

Declined tests are noted on the laboratory report.

During the requesting process the sample priority status will also be captured:

Initial antenatal screening sample

Repeat antenatal screening sample

Repeat sample to exclude recent infection

Initial sample after previous decline

Positive reports are copied to the Pre-Natal Screening Co-Ordinator as well as being notified to the Pre-Natal Screening team via email.

Any declined test, inconclusive result or inadequate sample is also notified via email to the Pre-Natal Screening team, to facilitate the timely collection of a repeat sample.

Samples received from patients >24 weeks or presenting in labour with no prior screening results will be tested urgently and results made available within 48 hours of receipt in the laboratory; however, the laboratory MUST be informed by telephone when such a sample is to be sent.

**HIV Antibodies**

ICE Request: HIV

eCARE Request: HIV 1 & HIV 2 Ag + Ab

It is the responsibility of the requesting clinician to obtain consent from the patient. If a sample is received in the laboratory, it will be assumed that consent has been obtained and the sample will be tested.The requesting doctor must make arrangements to convey the result to the patient.

Samples and request forms must be clearly labelled DANGER OF INFECTION. See 'Procedure for Sending Specimens with a Special Risk of Infection'.

**Requesting of Hepatitis Tests**:

5 mL of clotted blood is required where Hepatitis B or C is suspected or known, samples and request forms must be clearly labelled **‘Danger of Infection’**.

Information required from the requesters:

* Full clinical details
* Date of onset
* Immunisation history
* Details of any inoculation injury, including the hepatitis status of the source, if known

Please specify which tests you require:

***Hepatitis A:***

* To test for immunity: ICE: Hepatitis A immunity

eCARE: Hepatitis A total antibody, blood

* To test for current or recent infection: ICE: Hepatitis A IgM

eCARE: Hepatitis A IgM screen, blood

***Hepatitis B:***

To test for current or past Hepatitis B infection request the following:

ICE: Hepatitis B ?infection

eCARE: Hepatitis B infection screen, blood

Samples testing positive for either Hepatitis B core antibody or Hepatitis B surface antigen will go on to have further tests and may be referred to the reference laboratory for a full panel of infectivity markers.

If required Hepatitis B core antibody and Hepatitis B surface antigen can be requested individually:

* For evidence of past or present infection: Hepatitis B core antibody

ICE: Hepatitis B core IgG

eCARE: Hepatitis B core IgG, blood

To test for infectivity: Hepatitis B surface antigen

ICE: Hepatitis B surface Antigen

eCARE: Hepatitis B surface Ag, blood

* To test for immunity Anti Hepatitis B

ICE: Hepatitis B immunity

eCARE: Hepatitis B immunity, blood

***Hepatitis C:***

* For evidence of past or present infection: Hepatitis C IgG

ICE: Hepatitis C IgG

eCARE: Hepatitis C virus IgG, blood

* To test for infectivity Hepatitis C RNA

ICE: Hepatitis C RNA

eCARE: Hepatitis C virus RNA, blood

* Genotyping

ICE: Hepatitis C genotyping

eCARE: Hepatitis C virus genotyping, blood

Needlestick or body fluid exposure:

The full policy on management of inoculation injury is available in the Trust Policy “Managing Exposures to Blood Borne Viruses” available on the Trust Intranet.

A clotted blood sample should be collected in a gold top tube.

Ensure that clinical details clearly state that this is a Needlestick sample and whether the sample is from the donor (source) or the recipient (victim). On eCare/ICE the tests can be found by searching for Needlestick.

ICE: Needlestick Source Hep B, Hep C, HIV

or Needlestick Recipient (Victim) blood

eCARE: Needlestick Source blood

or Needlestick Recipient blood

Needlestick recipient samples are not tested but are stored for two years.

Needlestick donor (source) samples are routinely tested for HIV, Hepatitis B surface antigen and Hepatitis C IgG. If the patient declines any of these tests, please state clearly in the clinical details.

Telephone the laboratory to ensure urgent processing of Needlestick donor samples.

### TB T spot (Elispot) IGRA test

Sample must arrive in Microbiology by 2pm Monday to Thursday only.

For adults collect a minimum of 6ml blood in plain lithium heparin sample (dark green top, available from the laboratory upon request).

For paediatric patients from 2-9 years old: Minimum 2mls of blood in 2ml plain lithium heparin (light green translucent top). These samples are referred for testing.

ICE: TB Elispot Test

eCARE: TB Elispot, blood

### In House Serological Tests

| **Serological tests** | **Sample** | **Container** | **Turnaround** | **Notes** |
| --- | --- | --- | --- | --- |
| Antenatal infectious disease screen | CB | Gold top | ≤8 days excluding any referred test.  <48hrs for urgent sample | Telephone the laboratory when sending urgent samples |
| CMV IgG & IgM | CB | Gold top | ≤5 days |  |
| EBV serology | CB | Gold top | ≤5 days |  |
| Hepatitis A serology | CB | Gold top | ≤5 days |  |
| Hepatitis B serology | CB | Gold top | ≤5 days |  |
| Hepatitis C serology | CB | Gold top | ≤5 days |  |
| HIV 1 & 2 Ab/Ag | CB | Gold top | ≤5 days |  |
| Measles immunity | CB | Gold top | ≤5 days |  |
| Needlestick donor | CB | Gold top | HIV <24 hrs  Hep B&C up to 48hrs | Telephone laboratory when sending |
| Rubella IgG & IgM | CB | Gold top | ≤5 days |  |
| Syphilis | CB | Gold top | ≤5 days |  |
| Toxoplasma IgG & IgM | CB | Gold top | ≤5 days |  |
| VZV IgG | CB | Gold top | ≤5 days |  |
| VZV IgG antenatal contact | CB | Gold top | <24 hours | Telephone laboratory when sending |
| All serology tests can require referral for confirmation of in house results, in which case the turnaround time will be extended.  Expected turnaround times are valid during normal working hours 9am to 5pm Monday to Friday unless the samples is designated as urgent and agreed with the laboratory. | | | | |

*Most serological tests can be carried out on day of receipt by arrangement with the laboratory.*

### Referred Microbiology/Serology tests:

| **Test** | **Sample type** | **Ref. lab** | **Turnaround time (from receipt in ref. lab)** | **Notes** |
| --- | --- | --- | --- | --- |
| 16S PCR | CSF | MIC | 48HRS- 7 days |  |
| ACYCLOVIR ASSAY | CB | BRS | Same day result by phone |  |
| ADENOVIRUS SEROLOGY | CB | BRI | 5 days |  |
| ALPHAVIRUS SEROLOGY | CB | POR | 5 days |  |
| AMIKACIN ASSAY | CB | BRS | Same day result by phone |  |
| AMOEBIC F.A.T. | CB | HTD | 2 days |  |
| ANAPLASMA PHAGOCYTOPHILUM | CB | POR | 5 days | 10 Working Days |
| ANTHRAX INVESTIGATION | Biospy, eschar, washings, culture | POR | 3 working days | Discuss with Consultant Microbiologist |
| ANTIBIOTIC REFERENCE TESTS | Bacterial isolates | ARU | 15 days | Depends on species |
| ANTI-DNASE B | CB | BRI | 6 days |  |
| ARBOVIRUS SEROLOGY | CB | POR | 5 days |  |
| ASPERGILLUS ANTIGEN (GALACTOMANNAN)/PCR | CB/BAL | BML | 1 day |  |
| ASPERGILLUS PCR REF. | CB/EDTA/BAL/CSF | BML | 3 days |  |
| ASPERGILLUS PRECIPITIN | CB | CHU | 7 days |  |
| ATYPICAL PNEUMONIA SCREEN | CB | NOR | 2 days |  |
| AVIAN PRECIPITINS | CB | CHU | 7 days |  |
| BORD. PERTUSSIS SEROLOGY | CB | RSI | 10 days |  |
| BRUCELLA SEROLOGY | CB | NOR | 2 days |  |
| CAMPYLOBACTER SEROLOGY | CB | PRE | 7 days |  |
| CANDIDA PRECIPITINS | CB | BRI | 4 days |  |
| CHLAMYDIA SEROL (GENITAL) | CB | BRI | 5 days |  |
| CHLAMYDIA SEROLOGY (RESP) | CB | BRI | 5 days |  |
| CHLAMYDIA TYPING LGV | RECTAL SWAB (chlamydia tube) | STB | 6 days |  |
| CHLORAMPHENICOL ASSAY | CLOTTED BLOOD | BRS | Same day result by phone | Not tested on Saturday without prior arrangement |
| CJD | CSF | TSE | Contact lab | Discuss with Consultant Microbiologist |
| CLOST.PERFRINGENS TOXIN |  | GBRU | 5 days |  |
| CMV IGM CONFIRMATION | CLOTTED BLOOD | RFH | 7 days |  |
| CMV PCR | U, EDTA CB, VS | VJR | Variable | performed twice/week |
| COLISTIN ASSAY | CB | BRS | Same day result by phone |  |
| COXIELLA QFEVER SEROLOGY | CB | BRI | 5 days |  |
| COXSACKIE SEROLOGY | CB | EPS | 8 days |  |
| CRYPTOCOCCAL ANTIGEN | CSF  CB | BML | 1 day |  |
| CYCLOSERINE ASSAY | CB | BRS | 3 days |  |
| CYSTICERCOSIS REFERRAL | CB, CSF if indicated | HTD | 10 days |  |
| DENGUE FEVER | CB | POR | 5 days |  |
| DIPHTHERIA SEROLOGY | CB | DIP | 21 days |  |
| E. COLI O157 SEROLOGY | CB | LGP | 8 days |  |
| EBV PCR | EDTA | VJR | 3 days |  |
| ECHOVIRUS SEROLOGY | CB | EPS | 8 days |  |
| ENTEROVIRUS DETECTION | CSF, F, VS | EPS | 7 days |  |
| ENTEROVIRUS IGM | CB | EPS | 7 days |  |
| ETHAMBUTOL ASSAY | CB | ANU | 7 days |  |
| FILARIAL SEROLOGY | CB | HTD | 10 working days |  |
| FLAVIVIRUS SEROLOGY | CB | POR | 5 days |  |
| FLUCYTOSINE ASSAY | CB | BRS | Same day result by phone |  |
| GIARDIASIS SEROLOGY | CB | HTD | 10 days |  |
| H. PYLORI CULTURE | GBX | LGP | 15 days |  |
| HAEMOPHILUS(HIB) AB | CB | CHU | 7 days |  |
| HCV RNA QUAL/QUANT | EDTA | VJR | 8 days |  |
| HEP A IGM REFERRAL | CB | BIR | 5 days |  |
| HEP B CONFIRMATION | CB | VRD | 9 working days |  |
| HEP B DNA HEALTH WORKER | CB | BIR | 8 days |  |
| HEPATITIS B GENOTYPE | EDTA | VRD | 28 days |  |
| HEPATITIS B VIRAL LOAD | EDTA | VJR | 14 days |  |
| HEPATITIS C GENOTYPING |  | VJR | 8 days |  |
| HEPATITIS D (DELTA) REFER | CB | VRD | 15 days |  |
| HEPATITIS D RNA | EDTA | VRD | Contact lab |  |
| HEPATITIS E SEROLOGY | CB | VRD | 8 days |  |
| HEPC REF LAB IGG CONFIRM | CB | VJR | 8 days |  |
| HERPES IGG SEROLOGY | CB | VJR | 7 days |  |
| HERPES SIMPLEX PCR | VS | LEE | 14 days |  |
| HERPES TYPE SPEC SEROLOGY | CB | MAN | 7 days |  |
| HIV GENOTYPIC RESISTANCE | EDTA | BIR | 5-20 working days |  |
| HIV REF TESTS | CB | VRD | 9 working days |  |
| HIV VIRAL LOAD | EDTA | BIR | 14 working days |  |
| HTLV | CB | VRD | 8 days |  |
| HUMAN HERPES VIRUS 6 | CSF, CB, EDTA | NEW | Contact lab |  |
| HUMAN HERPES VIRUS 8 | EDTA | VRD | 15 days |  |
| HYDATID SEROLOGY | CB | HTD | 10 working days |  |
| INTRACONAZOLE ASSAY | CB | BRS | Same day result by phone | Only if advance warning given |
| JC/BK VIRUS DETECTION | CSF | VRD | 10 days |  |
| LEGIONELLA SEROLOGY | CB | RSI | 8 days |  |
| LEISHMANIA PCR & CULTURE | BM, SA, SKB | HTD | 20 days | Unless positive |
| LEPTOSPIRA SEROLOGY | CB | POR | 4 days |  |
| LYME DISEASE SEROLOGY | CB | POR | 5 days |  |
| LYME PCR | TIS, CSF, JASP | POR | 7 working days |  |
| MEASLES SEROLOGY REF | CB | ERN | 5 days |  |
| MENINGO/SPN PCR R LAB | CSF | MAN | 2 days |  |
| MERS CO-V | Discuss with Consultant Microbiologist | BAR | Contact lab | Contact lab prior to collection |
| MUMPS IMMUNITY | CB | VJR | 5 days |  |
| MUMPS SEROLOGY | CB | PRE | 2 days |  |
| MYCOPLASMA REFERENCE | CB | NOR | 2 days |  |
| NOROVIRUS PCR (COMMUNITY) | F | CAM | 2 days |  |
| PARASITE SEROLOGY | CB | HTD | 7-15 days |  |
| PARVO VIRUS ANTE NATAL | CB | VJR | 3 days |  |
| PARVOVIRUS PCR |  | VRD | 10 days |  |
| PARVOVIRUS SEROLOGY | CB | VJR | 7 days |  |
| PHLEBOVIRUS SEROLOGY | CB | POR | 2-5 days |  |
| PNEUMOCOCCAL ANTIBODIES M | CB | CHU | 7 days |  |
| PNEUMOCOCCAL PCR |  | MAN | 2 days |  |
| PNEUMOCYSTIS CARINII PCR | SP/BAL | MIC | 1 day |  |
| PROVIRAL HIV | EDTA | VRD | 8 days |  |
| RABIES IMMUNITY | CB | VET | Contact laboratory |  |
| RICKETTSIA SEROLOGY | CB | POR | 5 days |  |
| RUBELLA REFERENCE | CB | PRE | 2 days |  |
| SARS-CoV-2 (Covid) PCR (Novel Coronavirus) | CNT | MJR | 1-3 days |  |
| SARS CoV-2 (Covid) ANTIBODY | CB | VJR | 5 days |  |
| SCHISTOSOMIASIS SER. | CB | HTD | 7 working days |  |
| SCRUB TYPHUS | CB | POR | 5 days |  |
| STAPH TOXIN DETECTION | Bacterial isolate | ARU | 7 days |  |
| STREPTOCOCCUS GROUP B PCR |  | GOS | 1 days |  |
| STREPTOMYCIN ASSAY | CB | BRS | Same day result by phone |  |
| STRONGYLOIDES SEROLOGY | CB | HTD | 7 working days |  |
| SYPHILIS REFERENCE TEST | CB | STB | 7 working days |  |
| TB ELISPOT TEST (IGRA) | PLH | ODL | 2 days |  |
| TB PCR | SP | MRU | 1 working day |  |
| TEICOPLAININ ASSAY | CB | BRS | Same day result by phone | Only if advance warning given |
| TETANUS ANTIBODIES | CB | CHU | 7 days |  |
| THERMOPHILIC PRECIPITINS | CB | CHU | 7 days |  |
| TOBRAMYCIN ASSAY | CB | BRS | Same day result by phone |  |
| TOXOCARA ANTIBODIES | CB | HTD | 7 working days |  |
| TOXOPLASMA REFERRAL | CB | SWA | 10 working days |  |
| TRICHINELLA REFERRAL | CB | HTD | 5 working days |  |
| TRYPANOSOMA BRUCEI | CB, CSF if indicated | HTD | 10 working days |  |
| TRYPANOSOME SEROLOGY | CB | HTD | 10 days |  |
| VIRAL HAEMORRHAGIC FEVER |  | POR | 10-15 days | Discuss with Consultant Microbiologist before sending sample. |
| VIRAL PCR | Viral Swab | LEE | 14 days |  |
| VORICONAZOLE ASSAY | CB | BRS | Same day result by phone | Only if advance warning given |
| VZV IGM | CB | EPS | 5 days |  |
| WORM ID | WORM | HTD | 5 working days |  |
| YELLOW FEVER SEROLOGY | CB | POR | 4 days |  |
| ZIKA VIRUS | CB | POR | 7 days |  |

**SAMPLE TYPE CODES**

BAL Bronchioalveolar lavage

BM Bone Marrow

CB Clotted blood. Gold cap.

CSF Cerebrospinal fluid

EDTA EDTA blood. Purple cap.

F Faeces

GBX Gastric Biopsy

JASP Joint Aspirate

LH Plain lithium heparin (no gel). Dark green cap

SA Spleen aspirate

SKB Skin biopsy

SP Sputum

SW Swab

TIS Tissue

U Urine

VS Viral Swab

### Microbiology Referral Laboratories

| **Code:** | **Laboratory:** |
| --- | --- |
| ANU | PHE Anaerobe Reference Unit, Public Health Wales Microbiology Cardiff, University Hospital of Wales, Heath Park, Cardiff, CF14 4XW |
| ARU | Antimicrobial resistance and healthcare associated infections (AMRHAI), Public Health England, 61 Colindale Avenue, London, NW9 5EQ |
| BAR | Public Health England National Mycobacterium Reference Services – South (NMRL), National Infection Service, 61 Colindale Avenue, London NW9 5HT |
| BIR | Public health laboratory Birmingham, Heart of England NHS Foundation Trust, Bordesley Green East, Birmingham, B9 5SS |
| BML | National Mycology Reference Laboratory, Myrtle Road, Kingsdown, Bristol, BS2 8EL |
| BRI | Public health laboratory Bristol, Myrtle Road, Kingsdown, Bristol, BS2 8EL |
| BRS | Regional Antimicrobial Reference Laboratory, Microbiology, Lime Walk Building, Southmead Hospital, Westbury on Trym, Bristol, BS10 5NB |
| CAM | Clinical Microbiology and Public Health Laboratory (CMPHL), CMPHL Level 6, Box 236, Addenbrooke's Hospital, Cambridge, CB2 0QW |
| CHU | Immunology Dept., Churchill Hospital, Old Road, Headington, Oxford. OX3 7LJ |
| CRU | Cryptosporidium Reference Unit, Public Health Wales Microbiology ABM, Singleton Hospital, Sgeti, Swansea, SA2 8QA |
| DIP | Respiratory and vaccine preventable bacteria reference unit (RVPBRU), Public Health England, 61 Colindale Avenue, London, NW9 5EQ |
| EPS | Virology Department, St Helier Hospital and Queen Mary's Hospital for Children, Wrythe Lane, Carshalton, Surrey, SM5 1AA |
| GBRU | Gastrointestinal bacteria reference unit (GBRU), Public Health England, 61 Colindale Avenue, London, NW9 5EQ |
| GOS | Microbiology Laboratory, Level 4, Camelia Botnar Laboratories, Great Ormond Street Hospital for Children NHS Trust, Great Ormond Street, London, WC1N 3JH |
| HTD | National parasitology reference laboratory (NPRL), Department of Clinical Parasitology, Hospital for Tropical Diseases, 3rd floor Mortimer Market Centre, Mortimer Market, London, WC1E 6JB |
| LEE | Department of Microbiology, Old Medical School, Leeds General Infirmary, Thorseby Place, Leeds, LS1 3EX |
| LGP | Gastrointestinal bacteria reference unit (GBRU), Public Health England, 61 Colindale Avenue, London, NW9 5EQ |
| LHI | Antimicrobial resistance and healthcare associated infections (AMRHAI), Public Health England, 61 Colindale Avenue, London, NW9 5EQ |
| LIV | Brucella reference unit, Liverpool Clinical Laboratories, Virology Department, Royal Liverpool and Broadgreen University Hospital NHS Trust, Prescott Street, Liverpool, L9 8XP |
| MAN | Meningococcal reference unit (Men RU) Manchester, Clinical Sciences Building 2, Manchester Royal Infirmary, Oxford Road, Manchester, M13 9WL |
| MIC | Micropathology Ltd, University of Warwick Science Park, Venture Centre, Sir William Lyons Road, Coventry, CV4 7EZ, |
| MRU | National Mycobacterium Reference Laboratory (NMRL), Abernethy Building, Institute of Cell and Molecular Science (ICMS), 2 Newark Street, London, E1 2AT |
| MYC | Public Health England National Mycobacterium Reference Services – South (NMRL), National Infection Service, 61 Colindale Avenue, London NW9 5HT |
| NEW | PHE Newcastle Molecular Laboratory, Newcastle, Royal Victoria Infirmary, Newcastle upon Tyne, NE1 4LP |
| NOR | Health Protection Agency, Norfolk and Norwich University Hospital, Colney Lane, Norwich, NR4 7UY |
| ODL | Oxford Diagnostic Laboratories, Oxford Immunotec Ltd, 94C Innovation Drive, Milton Park, Abingdon, Oxfordshire, OX14 4RZ |
| POR | Rare and imported pathogens laboratory (RIPL), Public Health England, Manor Farm Road, Porton Down, Wiltshire, SP4 0JG |
| PRE | Food, water and environmental microbiology laboratory Preston, Royal Preston Hospital, Sharoe Green Lane, Fulwood, Preston, PR2 9HT |
| RFH | Virology Department, Royal Free Hospital, Pond Street, London. NW3 2QC |
| RSI | Respiratory and vaccine preventable bacteria reference unit (RVPBRU), Public Health England, 61 Colindale Avenue, London, NW9 5EQ |
| STB | Sexually Transmitted Bacteria Reference Laboratory (STBRU), Public Health England, 61 Colindale avenue, London, NW9 5EQ |
| SWA | Toxoplasma reference laboratory (TRL), Department of Microbiology, Singleton Hospital, Sgeti, Swansea, SA2 8QA |
| TSE | Virus Reference Department (VRD), Public Health England, 61 Colindale Avenue, London, NW9 5HT |
| UCH | Chlamydia Laboratory, Clinical Microbiology and Virology, University College London Hospitals NHS Foundation Trust, 60 Whitfield Street, London, W1T 4EU |
| VET | Sample Reception, AHVLA, Weybridge, New Haw, Addlestone, Surrey, KT15 3NB |
| VJR | Virology, John Radcliffe Hospital, Headley Way, Headington, Oxford, OX3 9DU |
| VRD | Virus Reference Department (VRD), Public Health England, 61 Colindale Avenue, London, NW9 5HT |

**HAEMATOLOGY AND BLOOD TRANSFUSION**

**INTRODUCTION**

Haematology offers both an analytical and clinical service. Clinical follow up is instigated when necessary. Most routine tests are performed daily and reported the same day. Significantly abnormal results are telephoned.

The Consultant Haematologists operate out-patient clinics listed below and admit patients to Ward 22 at Milton Keynes Hospital. A unit for day attendees is also currently based in the Macmillan Unit & Ward 22.

Consultant Immunology cover is provided both remotely and 1 session per week in the laboratory.

## Haematology Out Patients Clinics:

|  |  |  |  |
| --- | --- | --- | --- |
| Day | Clinic | Clinic Lead | Location |
| Monday | General Haematology | Dr Mitra (am) | Pathology HOPD |
|  | General Haematology | Dr Dungarwalla (am) | Cancer Centre |
|  | General Haematology | Dr Akanni (am) | Pathology HOPD |
|  | General Haematology | Dr Mitra (pm) | Cancer Centre |
| Tuesday | Anticoagulant Therapy | Anticoagulant Nurse (am) | Pathology HOPD |
|  | General Haematology | Dr Akanni (am) | Pathology HOPD |
|  | General Haematology | Dr Dungarwalla (am) | Cancer Centre |
|  | General Haematology | Dr Davis  (pm) | Pathology HOPD |
| Wednesday | General Haematology | Dr Dungarwalla (am) | Pathology HOPD |
|  | General Haematology | Dr Mitra (am) | Cancer Centre |
|  | General Haematology | Dr Hildyard (am) | Cancer Centre |
|  | General Haematology | Dr Mitra (pm) | Cancer Centre |
| Thursday | General Haematology | Dr. Mitra (am) | Pathology HOPD |
|  | General Haematology | Dr Davis (pm) | Pathology HOPD |
| Friday | General Haematology | Dr Dungarwalla (am) | Cancer Centre |
|  | General Haematology | Dr Hildyard (am) | Cancer Centre |
|  |  |  |  |
| \*Every Second Friday | Haemoglobinopathy | Dr Akanni (am) | Pathology HOPD |

## Anticoagulant Therapy

In-Patients

Prior discussion of the planned perioperative anticoagulant regime with the Haematology consultant and relevant physician and surgeon is helpful. A protocol is available in the Haematology Department for patients with atrial fibrillation, artificial heart valve or recurrent DVT requiring surgery. Please contact ext. 85814 or 85815 for a copy.

Discuss increased risk of bleeding/thrombosis with the patient.

Many surgical procedures (but not closed biopsies) can be performed with an INR of around 1.5 - 2.0.

If surgery is planned and continued anticoagulant cover is necessary, aim to gradually reduce the INR to cover the operation. Some patients will require the Warfarin to be stopped and intravenous Heparin instituted to cover the perioperative period

If urgent surgery is required, and the INR is greater than 2.0 the Warfarin can be temporarily reversed with Prothrombin Complex. In all cases contact the duty Consultant Haematologist via switchboard.

## Anticoagulant Clinic

To arrange outpatient anticoagulation control please phone the Haematology secretaries on extension 85814 or 85815. An Anticoagulation Clinic referral form

(i.e. Anticoagulation chart) will need to be completed and the following information given:

1) The patient’s name

2) Patient’s address (is hospital transport required?)

3) Hospital number

4) Indication for anticoagulation

5) Desired INR

6) Duration of anticoagulation

7) Recent Warfarin doses and INR results.

8) GP

9) Other medication

Generally, we can only see patients in the anti-coagulant clinic if the GP has been asked and is unwilling to anti-coagulate the patient. We can only see patients who are stable and require not more than weekly INRs.

## Investigation of Patients with Thrombophilia

If a thrombophilia screen is required, please arrange referral by writing to the Haematology Department with the personal and family history and whether VTE (venous thrombo-embolism) was spontaneous or precipitated (please give details). Note: Samples sent for thrombophilia screening from sources other than the Haematology Clinic will not be processed by the laboratory unless previously agreed.

Thrombophilia Screening Guidelines:

Ia) Unprovoked, recurrent or hormone related venous thrombo-embolism (VTE) **plus** a positive family history **or** in a patient less than 50 years old with children or siblings – **Full screen**. (If no positive family history and not less than 50 years with children or siblings, just do lupus anticoagulant (L.I) and anticardiolipin antibody testing (ACA).

Ib) Provoked VTE in a woman planning to be or currently pregnant – **Full screen**.

Ic) Family history of unprovoked, recurrent or hormone related VTE in a relative with a known thrombophilic abnormality identified – **Partial screen** (no lupus inhibitor screen or anticardiolipin studies).

IIa) Stroke in a patient < 50 years old – lupus inhibitor screen and anticardiolipin antibody screen

IIb) Three or more pregnancy losses **or** late (> 20 weeks) unexplained fetal loss – **Full screen**

|  |  |
| --- | --- |
| **Full Screen** | **Partial Screen** |
|  |  |
| Full Blood Count | Full Blood Count |
| Activated Partial Thromboplastin Time | Activated Partial Thromboplastin Time |
| Prothrombin Time + Thrombin Time | Prothrombin Time + Thrombin Time |
| Protein C, Free Protein S | Protein C, Free Protein S |
| Anti-thrombin III | Anti-thrombin III |
| Factor V Leiden + PT G20201A Mutation | Factor V Leiden + PT G20201A Mutation |
| Lupus Inhibitor Screen |  |
| Anticardiolipin antibody |  |

*(References:*

1. *Walker ID, Greaves M, Preston FE. Guideline: investigation and management of heritable thrombophilia. Br J Haematol, 2001 114****:*** *512-28.*
2. *Personal Communication. David Keeling, Churchill Hospital, Oxford, May 2004)*

## Guidelines for D-Dimer Testing in Suspected Deep Vein Thrombosis (DVT)

D-dimer testing currently should only be conducted in patients at low or moderate risk of DVT. All patients who have had a previous DVT or PE are immediately classified as high risk. Pregnant women or women in post-natal period should be managed outside this protocol and currently D-dimer testing is not appropriate. In suspected pulmonary embolus (PE) patients are currently managed outside this protocol.

|  |  |
| --- | --- |
|  | Points |
| Active cancer (patient receiving treatment for cancer within the previous 6 months or currently receiving palliative treatment) | 1 |
| Paralysis, paresis, or recent plaster immobilisation of the lower extremities | 1 |
| Recently bedridden for 3 days or more, or major surgery within previous 12 weeks | 1 |
| Localised tenderness along the distribution of the deep venous system | 1 |
| Entire leg swollen | 1 |
| Calf swelling at least 3 cm larger than the asymptomatic leg (measured 10 cm below tibial tuberosity) | 1 |
| Pitting oedema confined to the symptomatic leg | 1 |
| Collateral superficial veins (nonvaricose) | 1 |
| Alternative diagnosis at least as likely as deep vein thrombosis | -2 |

In cases in which it is unclear whether there is an alternative diagnosis, the assumption of no alternative diagnosis will ensure highest level of safety.

|  |  |
| --- | --- |
| **Score** | **Probability** |
| <1 | Low |
| 1 – 2 | Moderate |
| >2 | High |

Low/Moderate probability

+

D Dimer normal

DVT excluded

DISCHARGE

Low probability

+

D Dimer raised

High probability

+

D Dimer raised/

Normal/Not Done

**Doppler Ultrasound**

negative

Moderate probability

+

D Dimer raised

negative

positive

Repeat ultrasound 1 week

negative

DVT

diagnosed

*(D Dimer*

*Normal)*

*(D Dimer raised*

**Doppler Ultrasound**

The clinical score must be written on the venogram and D-dimer request cards and clearly written in the notes. In general D-dimer testing should not be carried out in high risk patients/pregnant women.

## Pulmonary Embolus and D-Dimers

Guidelines for D Dimer testing are available as a Clinical Guideline on the intranet.

## TESTS: Bottles Required Frequently and expected TAT of Routine Tests

**The viability of the sample indicates the time limit the sample will be valid for additional tests if requested. If additional tests are required, please check with the laboratory on availability and sample volume.**

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **Test** | **Bottle** | **Analysis** | **Sample Viability** | **Expected Turnaround time from receipt of sample** |
| *General* |  |  |  |  |
| Blood count | Lavender | Daily | 24 hours | 1 hour – urgent  2 hours - routine |
| Malaria | Lavender | Daily | 12 hours | 1 hour – urgent  2 hours - routine |
| ESR | Lavender | Daily | 12 hours | 2hours |
| Glandular fever test | Lavender | Daily | 24 hours | 1 hour – urgent  24 hours - routine |
| G6PD | Lavender | Weekly | 1 week | 7 days  If urgent discuss with lab |
|  |  |  |  |  |
|  |  |  |  |  |
| *Coagulation* |  |  |  |  |
| APTT | Blue | Daily | 4 hours | 1 hour – urgent  2 hours - routine |
| PT, INR | Blue | Daily | 12 hours | 1 hour – urgent  2 hours - routine |
| Full clotting screen | Blue | Daily | 4 hours | 1 hour – urgent  2 hours - routine |
| Platelet Function Testing | By appointment | By arrangement | 4 hours | Variable\*\* |
| Thrombotic screen | By appointment | By arrangement | 4 hours | 14 days |
| Factor assays | By appointment | By arrangement | 4 hours | 14 days |
| D-Dimers | Blue | Daily | 4 hours | 1 hour – urgent  2 hours - routine |
|  |  |  |  |  |
| *Haemoglobinopathies* |  |  |  |  |
| Sickle test | Lavender | Daily | 72 hours | 1 hour urgent |
| HbA2+F | Lavender | Three times per week | 72 hours | 7 days\* |
| Haemoglobin Variants | Lavender | Three times per week | 72 hours | 7-14 days |
|  |  |  |  |  |
| *Immunology* |  |  |  |  |
| Autoantibodies | Gold | Weekly | 1 week | 4 weeks (referral) |
| Endomysial antibodies | Gold | Weekly | 1 week | 4 weeks (referral) |
|  |  |  |  |  |
| Complement | Gold/Green | Daily | 48 hours | 3 days |
| Cardiolipin | Gold | \*\* | 1 week | 4 weeks (referral) |
| DNA/ENA | Gold | \*\* | 1 week | 4 weeks (referral) |
| Rheumatoid factor | Gold/Green | Daily | 24 hrs | 3 days |
|  |  |  |  |  |
| *Haematinics* |  |  |  |  |
| B12 & folate | Gold/green | Daily | 5 days | 24 hours |
| Red cell folate | Lavender | Weekly | 24 hours | 7 days |
| Ferritin | Gold/green | Daily | 5 days | 24 hours |
| Intrinsic Factor Antibodies | Gold/green | Daily | 48 hours | 24 hours |
|  |  |  |  |  |
| *Blood Transfusion* |  |  |  |  |
| Blood group & antibody screen | Pink 6 mL | Daily | 7 days | 24 hours |
| Crossmatch | Pink 6 mL | Daily | See request card | 24 hours  Urgent – discuss with laboratory |
| Kleihauer | 2 x Pink 6 mL | Daily | 48 hours | 48 hours |
| Ante-natal serology | Pink 6 mL | Daily | 48 hours | 48 hours |
| HIT screens | Referred tests – please discuss with laboratory | | | |
| HLA testing for platelet refractoriness |
| Neutrophil Antibodies |
| FMH flow cytometry |

\* Pre-natal samples 3 routine working days

## \*\* if urgent, results available on the same day by telephone. If not urgent results available within 7 days.

## Haematology Normal Ranges

Reference ranges or ‘normal values’ in Haematology may vary with age, sex and situation. Some variation from the stated range will occur for paediatric patients.

|  |  |  |  |
| --- | --- | --- | --- |
|  | **Male** | **Female (if different from male)** | **Unit** |
| Haemoglobin | 130 – 170 | 110 – 150 | g/L |
| Haematocrit | 0.4 – 0.5 | 0..36 – 0.46 | l/l |
| Red cell count | 4.6 – 6.2 | 3.8 – 4.9 | x1012/L |
| Mean cell volume (MCV) | 80 – 101 |  | Fl |
| Mean cell haemoglobin (MCH) | 27 – 32 |  | Pg |
| MCHC | 290 – 360 |  | g/L |
| White cell count | 3.7 – 11.1 |  | x109/L |
| Differential white cell count |  |  |  |
| Neutrophils | 1.7 – 7.5 |  | x109/L |
| Lymphocytes | 0.9 – 3.2 |  | x109/L |
| Monocytes | 0.2 – 1.0 |  | x109/L |
| Eosinophils | 0 – 0.5 |  | x109/L |
| Basophils | 0 – 0.1 |  | x109/L |
| Platelet count | 150 – 450 |  | x109/L |
| Erythrocyte Sedimentation Rate (ESR) <50yrs | 1 – 10 | 1 - 12 | Mm/hr |
| 50 - 60yrs | 1 – 12 | 1 - 19 | Mm/hr |
| 60-70yrs | 1 – 14 | 1 – 20 | Mm/hr |
| >70yrs | 1 – 30 | 1 – 35 | Mm/hr |
| Reticulocyte count | 0.2 – 2.0 |  | % |
|  |  |  |  |
| Red cell folate | –140-836 |  | ng/mL |
| Serum B12 | 150 – 900 |  | pg/mL |
| Serum folate | 3.1 – 19.9 |  | Ug/L |
|  |  |  |  |
| Hb F | <1.0 |  | % |
| Hb A2 | 2.2 – 3.4 |  | % |
| Ferritin |  |  |  |
| Male | 23.9 – 336.2 |  | ng/mL |
| Female | 11.0 – 306.8 |  | ng/mL |
|  |  |  |  |
| Prothrombin | 10.1 - 13.7 |  | seconds |
| Activated partial thromboplastin time (APTT) | 28.1 - 40.3 |  | seconds |
| Thrombin Time (TT) | 12.5 – 16.5 |  | seconds |
| Fibrinogen Assay | 1.8 – 4.5 |  | g/L |
|  |  |  |  |
| Immunology |  |  |  |
| C1 esterase inhibitor | 0.22 – 0.38 |  | g/L |
| C3 | 0.9 – 1.8 |  | g/L |
| C4 | 0.1 – 0.4 |  | g/L |

## Key Factors Affecting Results

|  |  |  |  |
| --- | --- | --- | --- |
| **TEST** | **KEY FACTORS AFFECTING RESULTS** | | |
| *General* | | | |
| Blood count | Haemolysis  Bacterial Contamination | Lipaemic  Pre-analysis storage temperature | Clotted  Cold Agglutinins  Sample age |
| Malaria  ***Note that there is no commercially available Rapid Diagnostic Test (RDT) that can specifically detect Plasmodium Knowlesi.*** | Haemolysis | Pre-analysis storage temperature | Sample age |
| ESR | Haemolysis | Lipaemic  Pre-analysis storage temperature | Clotted  Cold Agglutinins  Sample age |
| Glandular fever test | Haemolysis | Lipaemic | Sample age  ***See Note 1*** |
| G6PD | Post transfusion sample | Haemolysis | Clotted  Sample age |
| *Coagulation* | | | |
| PT, APTT | Underfilled/overfilled sample bottles Haematocrit >50  Patients on anticoagulation | Lipaemic  Pre-analysis storage temperature | Clotted  Sample age |
| INR | Underfilled/overfilled sample bottles Haematocrit >50  Patients on anticoagulation | Lipaemic Pre analysis storage temperature | Clotted  Sample age |
| Full clotting screen | Underfilled/overfilled sample bottles Haematocrit >50  Patients on anticoagulation | Lipaemic Pre analysis storage temperature | Clotted  Sample age |
| Platelet Aggregation | Aspirin and other drugs | Lipaemic Pre analysis storage temperature | Clotted  Sample age |
| Thrombotic screen | Underfilled/overfilled sample bottles Haematocrit >50  Patients on anticoagulation  Patients on Oral contraception,  Pregnant patients | Lipaemic Pre analysis storage temperature | Clotted  Sample age |
| Factor assays | Underfilled/overfilled sample bottles Haematocrit >50  Patients on anticoagulation | Lipaemic Pre analysis storage temperature | Clotted  Sample age |
| D-Dimers | Underfilled/overfilled sample bottles Haematocrit >50 | Lipaemic Pre analysis storage temperature | Clotted  Sample age  ***See Note 2*** |
| *Haemoglobinopathies* | | | |
| Sickle test | Post transfusion sample | Hb level | ***See Note 3*** |
| HbA2+F | Post transfusion sample | Hb level | ***See Note 3*** |
| Haemoglobin Variants | Post transfusion sample | Other Hb variants | ***See Note 3*** |

|  |  |  |  |
| --- | --- | --- | --- |
| **TEST** | **KEY FACTORS AFFECTING RESULTS** | | |
| *Immunology* | | | |
| Autoantibodies | Only gel/gold top suitable | Bacterial  contamination |  |
| Anti DNA | Only gel/gold top suitable |  |  |
| Complement | Gold or Green top suitable | Grossly lipaemic samples should be avoided |  |
| Cardiolipin | Only gel/gold top suitable |  |  |
| ENA | Only gel/gold top suitable |  |  |
| Rheumatoid factor | Only gel/gold top suitable | Bacterial contamination |  |
| *Haematinics* | | | |
| B12 & folate | Post transfusion sample | Haemolysis | ***See Note 2*** |
| Red cell folate | Post transfusion sample | Haemolysis | ***See Note 2*** |
| Ferritin | Post transfusion sample | Haemolysis  Acute phase protein | ***See Note 2*** |
| Intrinsic Factor Antibody | High levels of vitamin B12 |  | ***See Note 2*** |
|  | | | |
| *Blood Transfusion* | | | |
| Blood group & antibody screen | Post transfusion sample | Haemolysis |  |
| Crossmatch | Post transfusion sample | Haemolysis |  |
| Kleihauer | Post transfusion sample HPFH | Haemolysis |  |
| Ante-natal serology | Post transfusion sample | Haemolysis |  |

**Note 1 Glandular Fever screening**

Glandular Fever tests

Occasionally detectable levels of Heterophile antibodies are late in developing in patients symptomatic for Infectious Mononucleosis. If symptoms persist it is recommended to repeat the test after several days. Some patients may remain persistently negative, especially children and adolescent. It has been reported that only 80 to 90% of adults and less than 50% of young children develop Heterophile antibodies.

False positive reactions when have been reported in serum samples collected from patients with recent cytomegalovirus, hepatitis A virus, parvovirus and leptospira infection.

*Detectable levels of Heterophile antibodies may persist for months, and more rarely for years, in some individuals.*

**Note 2 Warning to Users with regards to HAMA antibodies on serum B12, serum Folate and Ferritin assays.**

HAMA antibodies seen in Haematinic Assays and D Dimer results

For assays employing antibodies, the possibility exists for interference by heterophilic antibodies in the patient sample. Patients who have been regularly exposed to animals or have received immunotherapy utilising immunoglobulins may produce antibodies e.g. HAMA, that interfere with immunoassays. Such antibodies may cause erroneous results.

HAMA (Human anti-mouse antibodies) could be seen in the following assays:

Serum B12

Serum Folate

Ferritin

Red Cell Folate

Intrinsic Factor antibodies

D Dimer

It is well documented as a phenomenon of possible interference and results should be interpreted in light of the total clinical presentation of the patient, including symptoms, clinical history, data from additional tests and other appropriate information.

**Note 3 UK National Screening Programme for Infectious Diseases, Sickle Cell &Thalassemia in Pregnancy**

A Family Origin Questionnaire (FOQ) **must** be completed and sent with all requests for Antenatal booking bloods. This requester **must** provide the gestation date and the EDD and complete the ethnic origin of both mother and father. In low prevalence areas the FOQ is used as a tool to identify women who are at highest risk of having a baby with a haemoglobin variant or disorder.

In addition, the joint paper request/FOQ form will require details of consent. If declined, the reason for decline must be given. The FOQ should be completed and returned with the booking bloods at <10 weeks of gestation.

For clinical advice regarding Antenatal screening please contact:

Infectious Diseases Antenatal Screening Programme Advice – Consultant Microbiologist - **Dr Poonam Kapila 85786 99 5786**

Sickle Cell and Thalassaemia Antenatal Screening Programme Advice - Consultant Haematologist – **Dr Mags Akanni 87573 99 7573**

**BLOOD GROUPING AND TRANSFUSION**

A copy of the Trust Policy for Blood Transfusion, the Trust Policy for the Treatment of Patients Refusing Blood/Blood Products or their Derivatives and the Trust Policy for the Management of Massive Blood Loss can be found on every ward. **Every member of staff involved in the transfusion process should read these.**

This policy can also be accessed under Clinical Policies on the Trust intranet.

**PATH-GL-03 Blood Transfusion Policy**

Please give 24 hours’ notice for non-urgent cases. Note that patients with positive antibody screens or who are known to have antibodies may not have suitable blood on site and it may take 48 hours for the provision of suitable blood.

A 6 mL EDTA Transfusion bottle is required for blood grouping, cross matching and antibody screening for adult patients. 4 mL EDTA samples (if correctly labelled) will be accepted for adult patients with compromised venous access or paediatric patients.

1.3 mL EDTA samples will be accepted for paediatric requests

The minimum sample volume for adult requests is 2 mL and 1 mL for paediatric requests

Note issue of crossmatched group specific red cells is dependent upon there being at least two blood group results being available. Where this criterion cannot be met group O RhD Positive or Negative red cells will be provided.

The sample should be clearly labelled with patient’s surname, forename, hospital number, date of birth and date sample taken and signed by the taker**.** Samples labelled with patient Addressograph labels will be discarded, as this is contrary to hospital policy. Full details of the sample labelling and request form documentation requirements can be found in the hospital policy: **PATH/GL/14 Acceptance of Samples in the Pathology Department**

The only blood collection point is the main Blood Bank refrigerator. Blood kept in theatre fridges is only to cover intra-operative blood loss.

The instructions regarding the labelling of samples for the purpose of blood transfusion, along with the guide on when new samples are required, can be found on the back of the request forms and further on in this section.

## Transfusion Procedures and Practice

1. An urgent blood group & antibody screen takes 45 minutes from the sample arriving in the laboratory. Should blood be required before this the laboratory will issue group O red cells until there are two blood group results available from two separate samples
2. Non urgent group & antibody screen requests shall be processed within 4 hours of receipt
3. For patients who have been previously grouped and screened (twice) and require blood:
4. Cross-match will take 30 minutes.
5. If criteria are met an electronic cross-match will be performed, blood will be available in 5 minutes (contact the laboratory regarding the eligibility of patient for this procedure).
6. Where blood group results are available for both a historical sample and a current sample (see f below) but where antibody screen results are not yet available group specific (matching the patient ABO type) uncrossmatched blood can be provided. This is as safe for patient use as group O Negative blood.
7. Failing the above, uncross-matched O blood can be issued immediately. This is a precious resource and should only be used in life threatening situations. Note: male patients and female patients who no longer have child bearing potential may be issued with O Positive blood
8. For patients requiring transfusion in emergency situations before compatibility testing can be performed 4 units of O Negative and 4 units of O Positive blood are available for immediate use. They are kept within the pathology Blood Issue Fridge. The blood transfusion laboratory must be informed if these units are collected and to whom they have been transfused as soon as possible. Use of this blood may increase risk of transfusion reaction. If possible, the laboratory should be contacted prior to the use of these units as it may be possible to provide uncrossmatched blood that is more suitable for the patient. A further 2 units of emergency O Negative are located in Phase 1 & 2 Theatre blood fridges.
9. Blood is reserved from the date requested for 24 hours.
10. Repeat sample requirements for blood product requests

**Patients pregnant or transfused within previous 3 months or who have a history of red cell antibodies – sample must be taken <72 hours before the start of a new transfusion episode**

**Patient not pregnant or transfused within previous 3 months and with no history of red cell antibodies – sample must be taken <7 days before the start of a new transfusion episode**

Samples are only stored for 7days

1. Appropriate prescribing of Blood Products

Guidance for appropriate prescribing of blood products including surgical blood order schedule can be found in the Trust Blood Transfusion Policy copies of which can be found on each ward but is also universally available on the hospital intranet under clinical policies:

**PATH-GL-03 Blood Transfusion Policy and in associated Platelet and Plasma Product Guidelines.**

Further guidance for appropriate prescribing of blood including for medical patients can be found in the Trust policy**: PATHOLOGY/GL/09 Prescribing Red Cell Transfusion in Adults:**

Guidance for the management of paediatric patients can be found in the Trust **PATH/GL/20 Paediatric Transfusion Policy**

and **PAED/GL/48 Neonatal Red Cell Transfusion Guideline**:

Specific guidance on the use of Prothrombin Concentrate Complex (PCC) can be found in the Trust **Guideline for the Prescribing & Administration of Prothrombin Concentrate Complex (Octaplex) PATHOLOGY/GL/22**

Guidance for the management of massive blood loss or major haemorrhage is available in the Trust policy **PATH/GL/05**

Guidance on the management of patients that refuse transfusion of blood or blood products can be found in the Trust policy **PATH/GL/04**

1. Patients presenting with suspected transfusion reaction

Guidance for the management of these patients is provided in the Hospital Blood Transfusion Policy (see above). In all cases the transfusion should be stopped; the laboratory and duty Haematology Consultant informed immediately; a transfusion reaction form completed and sent to the laboratory with appropriate samples; the suspected reaction must also be reported on the hospital incident reporting system (Datix). It is the responsibility of the attending clinician to ensure these tasks are completed.

1. Responsibilities

Currently clinicians are responsible for the prescribing of all blood and blood products and the management of patients with suspected transfusion reactions. Collection of blood components can only be performed by staff trained and certified as competent. Access to the Blood Issue Fridge is restricted to staff that meet this requirement. For further information about the collection and administration of blood products contact the Transfusion Specialist Practitioner Ext. 85798 or Bleep 1644 Monday-Friday.

**IMMUNOLOGY LABORATORY SERVICE**

##### Principles of the Tests and Clinical Implications

ANA Screen The term connective tissue disease refers to a group of disorders involving the protein-rich tissue that supports organs and other parts of the body. These disorders often involve the joints, muscles, and skin but they can also involve other organs and organ systems, including the eyes, hearth, lungs, kidneys, gastrointestinal tract, and blood vessels. They are o inflammatory diseases of unknown origin with humoral and cellular disturbances, systemic organ failure and chronic disease. These diseases are characterised by overlapping symptoms, thus making them difficult to diagnose.

When systemic autoimmune disease is suspected a common practice is to perform a general screening test for antinuclear antibodies. The ANA test if positive, leads to screens for the antibodies to RNP-70, RNP/Sm., Sm., SS-A, SS-B, Scl 70, Centromere and Jo-1.

Anti-dsDNA Systemic lupus erythematosus (SLE) is characterised by the presence of autoantibodies against native double stranded DNA (dsDNA). Additionally, these patients also exhibit autoantibodies against the single stranded form of DNA. Together with the determination of antinuclear antibodies (ANA) the determination of dsDNA antibodies are the most important serological criteria for diagnosing SLE. Autoantibodies against dsDNA have a diagnostic specificity of about 96% and a sensitivity of about 91% for diagnosing SLE. Anti-dsDNA antibodies can be useful in some patients to monitor therapy and to predict disease progression.

ENA Screen This test determines the IgG class autoantibodies directed against the extractable nuclear antigens SS-A, SS-B, Sm, RNP/Sm, Scl-70 Jo-1 & Centromere.

Anti SS-A(Ro)/SS-B(La) Autoantibodies against SS-A(Ro) and SS-B(La) are found in most patients suffering from primary Sjögrens syndrome. Inflammatory processes of the salivary and lachrymal glands are characteristic of this disorder. Patients with sicca syndrome also exhibit these antibodies. More importantly, anti SS-A(Ro) antibodies are present in >80% of mothers of infants with congenital heart block.

Anti RNP/Sm RNP is associated with patients suffering with systemic lupus erythematosus and undifferentiated connective tissue disease. Autoantibodies against Sm proteins are of pathognomonic importance for diagnosing SLE. A negative finding of anti-Sm does not exclude SLE.

Anti-Scl-70 Scl-70 antibodies are directed against the enzyme DNA Topoisomerase I and are highly specific for Scleroderma. Progressive Systemic Sclerosis is an autoimmune disorder of the connective tissue leading to slowly progressive fibrosis and later to sclerosis. Patients suffering from scleroderma develop often develop tight skin as the first manifestation with involvement of the heart, kidneys and lungs.

Patients with Scl-70 Abs are more likely to have facial skin involvement and disease in heart, kidneys and lungs in comparison with anti-Scl-70 Ab-negative scleroderma patients. Anti centromere antibodies are found in the CREST subset of scleroderma.

Rheumatoid Factor: Rheumatoid Factor is a non-specific serological indicator included in the diagnosis of RA. A weak positive, clinically insignificant result may be found in normal elderly subjects. RF negative RA patients do exist

Anti-Cardiolipin: Autoantibodies directed against the anionic phospholipid Cardiolipin are primarily found in patients suffering from primary and secondary antiphospholipid syndrome. These patients have clinical manifestations such as thrombosis of the veins and arteries, thrombocytopaenia and recurrent foetal loss.

Anti- CCP : Antibodies to cyclic citrullated peptide are found in early Rheumatoid Arthritis and are predictive of a more severe arthropathy. Anti-CCP antibodies are more specific for RA than RF.

Anti-β2 GPI Antibodies to this co-factor for cardiolipin may be useful in early diagnosis of the anti-phospholipid syndrome.

Anti-PR3 c-ANCA: Antineutrophil antibodies were first reported in 1982 in patients with necrotizing glomerulonephritis. This led to the discovery of autoantibodies detected in systemic vasculitic disorders. Proteinase 3 is the classic autoantigen in Wegener’s granulomatosis. Approximately 66% of patients in the early stages of the disease exhibit anti -PR3 and it can be detected in more than 95% of all untreated patients (may disappear after immunosuppressive treatment).

Anti-MPO p-ANCA Churg -Strauss syndrome exhibits autoantibodies against lysosomal Myeloperoxidase (MPO-ANCA). Churg-Strauss is an allergic granulomatosis and angiitis. The autoantibodies are also found in patients with microscopic polyangiitis.

Anti Thyroid Peroxidase: Anti-TPO autoantibodies are detected in cases of Hashimoto’s thyroiditis, myxoedema and Graves’ disease.

Anti Tissue Transglutaminase: IgA antibodies to tTG is a main screening investigation for Coeliac Disease.

Endomysial Antibody: This is also used to confirm a diagnosis of Coeliac Disease performed by IFA: together with anti-Tissue Transglutaminase IgA, it forms a screen for adults.

[Coeliac screening](http://www.ouh.nhs.uk/immunology/diagnostic-tests/tests-catalogue/coeliac-antibodies.aspx) is carried out by performing a combination of anti-endomysial and anti-TTG antibody screens.

**COELIAC ANTIBODIES**

**Also known as:** Endomysial Antibodies; Endomysium Antibodies; IgA Endomysial Antibodies

IgA endomysial antibodies are currently the most specific assay for Coeliac Disease screening with a sensitivity of 70 - 90% and specificity 90 - 100% and will detect virtually all cases of untreated coeliac disease except those with co-existing IgA deficiency. For this reason, a total serum IgA is performed alongside, if this is low (<0.06 g/l) and the endomysial antibody is negative, the serum is tested for IgG endomysial antibodies. There is some controversy as to whether there is a good correlation of endomysial antibody with disease activity, There is evidence that poor compliance with a gluten free diet is associated with a return to antibody positivity. IgA endomysial antibodies are also present in patients with Dermatitis Herpetiformis (DH).

If suspect Coeliac disease, please ensure the patient is on a gluten containing diet before serological testing.

Associated tests

* TTG antibody
* IgG endomysial antibodies

See ng20 September 2015: NICE guidelines

For further information contact

Dr Elizabeth Bateman FRCpath  
Department of Immunology  
Churchill Hospital  
Old Road  
Headington  
Oxford  
OX3 7LE  
(01865) 225991 or 225995

## Clinical Indications and Screening Tests

|  |  |  |
| --- | --- | --- |
| **Clinical Indications** | **Initial Screen** | **Further Tests** |
|  |  |  |
| SLE, RA, other connective Tissue diseases | Auto Immune Profile (to include ANA) | Depends on clinical symptoms |
|  |  |  |
| Pernicious Anaemia | Anti smooth muscle |  |
| Primary biliary cirrhosis | Anti gastric parietal |  |
| Chronic active hepatitis | Anti mitochondrial |  |
| Auto Immune Hepatitis | Anti Liver Kidney Microsomal |  |
|  |  |  |
| Arthritis**/**RA/ | Rheumatoid Factor , ANA | ENA, anti CCP if RF negative |
| Rheumatoid joint pain |  |  |
|  |  |  |
| SLE, lupus, UCTD | ANA, C3,C4 | ENA, DNA |
|  |  | (ENA, Cardiolipin in pregnancy) |
|  |  |  |
| Monitoring SLE | dsDNA |  |
|  |  |  |
| Connective tissue | ENA |  |
| Disease |  |  |
|  |  |  |
| Goodpasture’s Syndrome | Glomerular Basement Membrane Ab |  |
|  |  |  |
| Drug induced lupus | ANA | DNA |
|  |  |  |
| Recurrent thrombosis | ANA, Cardiolipin/β2GPI | DNA, ENA, C3/ C4 |
| antiphospholipid syndrome |  |  |
| recurrent miscarriage/abortion |  |  |
|  |  |  |
| Raynauds | ANA, Centromere | ENA, DNA, |
|  |  |  |
| Wegener’s granulomatosis | cANCA IIF confirmed by anti PR3 & anti MPO | Anti PR3 & Anti MPO |
|  |  |  |
| Vasculitis, pauci immune | pANCA | DNA |
| Glomerulonephritis, |  |  |
| Churg-Strauss |  |  |
|  |  |  |
| Sjögren’s syndrome/sicca | ANA, ENA | DNA |
| syndrome Dry eyes, mouth |  |  |

|  |  |  |
| --- | --- | --- |
| **Clinical Indications** | **Initial Screen** | **Further Tests** |
|  |  |  |
|  |  |  |
| Scleroderma.CREST | ANA, Centromere | Scl-70,ENA,DNA |
| Systemic sclerosis |  |  |
| Myositis | ANA | Jo-1, ENA,DNA |
|  |  |  |
| Liver disease/ Hepatitis/ CAH PBC/abnormal LFT/pruritus | Mitochondrial/Smooth Muscle & LKM Ab | M2 specific mitochondrial for PBC |
|  |  |  |
| Pernicious anaemia/B12 deficiency | Gastric Parietal Cell | Intrinsic Factor Abs |
|  |  |  |
| Thyroid disease | Thyroid peroxidase antibody | TSH Receptor Ab in Graves’ |
|  |  |  |
| Addison’s Disease | Adrenal antibodies |  |
|  |  |  |
| Infertility/Amenorrhoea | Ovarian antibodies |  |
|  |  |  |
| Diabetes/IDDM(Stiff man | Islet cell Antibody, GAD antibody |  |
| Syndrome) |  |  |
|  |  |  |
| Coeliac disease/ abdominal pain | IgA anti tTG-Ab | Endomysial Ab to confirm |
| Diarrhoea/FTT/Malabsorption |  | Check for IgA deficiency |
|  |  |  |
| Renal failure | ANCA, GBM, ANA C3/C4 | DNA, ENA, MPO, PR3 |
| Glomerulonephritis |  |  |
|  |  |  |
| Stroke, TIA (<60 yr) | Cardiolipin IgG | β2GPI Ab |
|  |  |  |
| Pemphigoid | Basement membrane Ab | Direct skin biopsy IF |
| Pemphigus | Intercellular cement Ab | Direct skin biopsy IF |
|  |  |  |
| Myasthenia Gravis | Acetylcholine receptor ab |  |
| Thymoma in Myasthenia |  | Skeletal muscle antibody |
| Other myasthenic syndromes (Lambert-Eaton etc) | VGCC Abs |  |

ANCA testing is limited to specific conditions and locations. Please refer to the relevant protocol on eCare/ICE or contact the laboratory for advice.

**CELLULAR PATHOLOGY**

**INTRODUCTION**

The Cellular Pathology Department provides a diagnostic histology and cytology service to Milton Keynes Hospital, local GPs, the Breast and Bowel Cancer Screening Programmes and other local private health care providers. This is supported by a comprehensive immunohistochemistry service and referral for Her2 testing and molecular diagnostics.

The department acts as delivery point for specimens from the Cervical Cytology Screening Programme. The service is provided by South Bucks NHS Trust at High Wycombe Hospital.

The department provides a comprehensive autopsy service for Milton Keynes Hospital and HM Coroner

**REQUESTING A CELLULAR PATHOLOGY INVESTIGATION**

Quality and patient safety in Cellular Pathology starts from the moment the sample is taken. Please follow the instructions regarding request form and sample requirements in order to provide the most accurate result for the patient.

You can request Cellular Pathology investigations either on hand-written blue request cards or electronically with eCare/ ICE. Requests for histology and cytology on the same patient should be made as separate requests on eCare/ ICE or each have a blue request form.

All electronic requests must be accompanied by a signed request form.

All requests must contain 3 matching legible patient identifiers between the form and each pot. The following are mandatory:

* + Unique identification number (MRN or NHS number, both if available)
  + Full name
  + Date of birth

In addition to the mandatory information, we also require relevant clinical and sample information in order to provide the most accurate result.

* + Specimen information to include type and anatomical site of specimen. Multiple specimens from a single patient must be clearly labelled and differentiated, and corresponding information provided on the request form and the pot
  + Sender’s details (Full name) and bleep / phone number
  + Location of where the procedure took place
  + All relevant clinical information. This should be as extensive as practical and should include nature, appearance and site of the lesion, duration and nature of symptoms, results of relevant investigations including imaging, relevant past medical history and clinical differential diagnosis.
  + Infection risk
  + Priority (e.g. urgent, 2WW)
  + Date and time specimen was taken (Essential to determine the length of time of fixation)

**ACCEPTANCE CRITERIA**

Accurate identification details on cellular pathology specimens are vital for patient safety.

It is the responsibility of the clinician requesting the investigation to ensure that the specimens are correctly labelled and that the request details are completed to the above standard. Specimen and request details must be compatible.

Where essential information is missing from a specimen or request form the laboratory will attempt to contact the requesting clinician and will issue a Specimen Rejection Form. The requesting clinician (or other responsible person who has been given the authority by the clinician to identify the specimen) will be required to attend the laboratory to complete or amend the details before the specimen can be processed.

This person must sign the form to confirm that they have agreed to take responsibility for the identification of the specimen and/or any amendments made.

Where the missing information includes the requesting clinician a printed hard copy of the report may be delayed or unavailable. In this case the report may be issued with “Unknown Sender” on the laboratory computer system.

**SPECIMEN REQUIREMENTS**

* Except where indicated below all specimens should be sent to the laboratory in an adequately sized pot containing formalin fixative (10% Neutral Buffered Formal Saline).
* Poor fixation can compromise the quality of the specimen and subsequent histological examination so specimen pots should be large enough to easily accommodate the specimen. The volume of formalin should be at least 10 x the volume of the specimen, so it is important not to squeeze the specimen into a container that is too small.
* Specimens should be placed into fixative as soon as possible after removal from the patient. With small biopsies in particular, it is very important not to let the specimen dry out
* Small endoscopic biopsies should be placed into a ‘mini’cassette before placing into formalin. This allows safe handling of the small biopsies in the laboratory. (‘Mini’ cassettes are available from the laboratory on request)
* Prostate needle core biopsies should be placed between moist sponges and placed into a blue processing cassette. Care should be taken not to allow the cores of tissue to overlap. The cassette should be placed into a pot of ‘pink’ formalin which contains a drop of eosin to colour the tissue cores making them easier to handle in the laboratory. This coloured formalin should not be used for other larger specimen types. (Cassettes, sponges and ‘pink’ formalin pots are available from the laboratory on request)
* Pots should be labelled as specimens are placed into them. It is poor practice to label pots in advance of a procedure.
* All pot labels should include full patient identifiers, specimen details and relevant hazard indicators.
* Where possible specimens should be sent from theatres and clinics to the laboratory regularly throughout the day and should not be batched to be sent at the end of the day. This ensures that urgent diagnostic biopsies are processed as soon as possible and allows a more efficient and timely production of results.
* Upon receipt in the laboratory large resection specimens require immediate opening and/or slicing to ensure adequate fixative penetration. Large specimens that are delayed reaching the laboratory may suffer irreversible tissue damage which will compromise the quality of the final report.

Containers of formalin should be securely closed before transport to the laboratory. Formaldehyde vapour is a well-recognised respiratory irritant and possible carcinogen. Inhalation of formalin vapour and skin contact should be avoided as repeated exposure may cause skin sensitisation and allergic contact dermatitis.

Spillage kits should be available in all areas that store formalin pots.

Specimens in formalin must not be sent via the hospital air tube system.

**High Risk/ Danger of Infection specimens:**

Specimens potentially infected (known or suspected) with a Hazard Group 3 organisms must be clearly marked as such, and the nature of the risk described. Laboratory staff need to be able to work safely and process the specimen appropriately.

High risk specimens for routine histology must be fixed in 10% formalin for at least 24 hours in the laboratory prior to processing. As a result, there will be a subsequent 24 hour delay in reporting for small specimens and 48 hours for large resection specimens.

Please do NOT send specimens for disposal to the laboratory.

**SPECIAL SAMPLE COLLECTION**

### Frozen Section for Rapid Intra-operative Diagnosis

Whenever possible, frozen sections should be booked in advance on Ext. 58521

Without advance arrangement the laboratory cannot guarantee that a pathologist will be available to report the biopsy.

The specimen must be sent in a labelled clean dry container. DO NOT place the specimen into formalin.

Please identify the theatre in which the procedure is being performed together with a contact number for the surgical team on the request card.

To avoid unnecessary delays in reporting specimens for frozen section must be brought directly to the Histopathology Laboratory (Behind Ward 3, next to the mortuary) and **NOT** taken to main Pathology Reception

The turnaround time from receipt of the specimen in the laboratory to the issue of a telephoned report is 20-30 minutes.

Please contact the laboratory to cancel if the frozen section is no longer required.

### Direct Immunofluorescence studies on skin biopsies

All of the skin samples must be delivered immediately to the laboratory in a Universal containing Saline available from the laboratory, accompanied by an eCare/ ICE or blue request card, clearly stating that the specimen is for direct immunofluorescence studies.

**These specimens are sent to John Radcliffe Hospital for testing. Results are usually available on eCare and ICE within 5-7 days**

### Cytogenetics

All samples should be sent to the laboratory in Tissue transport medium – available from Churchill Hospital, Oxford and must be accompanied by a Churchill request form. Tissue Transport medium is stored on Labour Ward – the laboratory does not keep a stock

### Foetus’s & Pregnancy Remains

This area is regulated by The Human Tissue Authority. It is vital that the request cards that accompany specimens state the gestational age. The fate of these specimens depends on having this information. The laboratory will not process specimens that do not have this clearly stated in the clinical information and will be returned.

* **Less than 14 weeks gestation**

These samples should be sent to the laboratory in 10% formal saline. After reporting, the laboratory sends these specimens for cremation.

* **14 – 18 weeks gestation**

These samples should also be sent to the laboratory in 10% formal saline. After reporting, the laboratory will pass the specimen to the mortuary who will send these cases separately for cremation.

* **Greater than 18 weeks**

Foetus’ greater than 18 weeks are sent direct to the mortuary. All such specimens should be sent to the mortuary in a clean, dry container without fixative, together with a *blue request card*. The placenta should also be sent in a dry container and should always accompany the foetus. If parents request no investigations, it must be clearly stated on the request form. t

* If a PM / Histopathology is required then a *PM consent form* is required, **regardless of the age of the foetus.** These are then passed on to the Paediatric Pathology Unit at the John Radcliffe Hospital in Oxford. It must be ascertained from the parents of every case that is sent to the mortuary for post mortem, whether the mother would like the return of the placental tissues, for burial or cremation with the baby or disposal via Oxford’s hospital protocol. If the mother opts for burial or cremation, residual wet tissue will be returned from Oxford.

The options are that Oxford will return the placental tissue, with the baby to MK, so that disposal can be arranged (cremation or burial) or the placenta will be incinerated in line with Oxford’s hospital protocol.

**DIAGNOSTIC CYTOLOGY**

All specimens must be sent as soon as possible to the laboratory before degeneration of the cells occurs. If this is not possible, ensure that the specimen is kept refrigerated.

**PLEASE SEND SEPARATE SPECIMENS AND MAKE SEPARATE REQUESTS IF REPORTS REQUIRE BOTH MICROBIOLOGY AND CYTOLOGY INVESTIGATION**

### Sputum Cytology

This is recognised as a specimen of limited or no clinical value. Where patients are unfit for bronchoscopy, three separate sputum samples collected on different days could be sent for cytological examination. Nebulised saline may be used to induce sputum production in appropriate clinical circumstances. Guidance should be given to the patient on producing a deep cough sample. A salivary sample is inadequate for cytology. The whole of the expectorated sample could be sent in a pot (60ml pot) already labelled by pathology **‘sputum only’.** Please put the sample in the refrigerator if there is unavoidable delay (overnight).

### Body Cavity and Cyst Fluid Cytology

Fluid for examination should be sent in a 20ml Universal container.

Deliver to the laboratory as soon as possible on the day of collection

### Urine Cytology

Freely voided, catheter, ileal conduit specimens or bladder/ureteric washings may be collected. It is essential that the mode of specimen collection is documented on the request form. The sample can be put in the refrigerator if there is a short delay. A maximum of 20ml of fresh sample is required.

The first urine passed in the morning should be avoided. A midstream specimen is suboptimal. For voided urine an aliquot of the whole voided sample should be submitted. Samples may be taken from the upper urinary tract by clinicians specialised in the technique and should be handled in the same way as the other urine specimens.

### Fine Needle Aspirates (FNA) of Solid Lesions and brushings

Fine needle aspirates are generally taken by physicians, surgeons and radiologists. Palpable masses may be aspirated without image guidance. Ultrasound guided aspirates generally give better samples. FNA kits are available in the laboratory or by contacting pathology supplies on ext 85793The FNA kit comprises slides, slide box, fixative, normal saline, instructions and a request card. Needles and syringes are not included.

Links to educational material on how to take an FNA are available on several web sites, e.g. [(www.pathlab.org)](http://www.pathlab.org/)

Generally 22 to 25 gauge needles are used for aspiration of solid organs ([www.pathlab.org](http://www.pathlab.org/)).

A 25 gauge needle with a 10cc syringe can be used for thyroid aspiration. A 3/4 inch long needle is usually sufficient; a 1.5 inch long needle can be used for large and deep lesions. A Cameco syringe pistol can be attached to the syringe to facilitate the process (Douglas P. Clark and William C. Faquin. How to Perform and Process a Thyroid FNA. Thyroid Cytopathology 2005:12). A larger bore needle is associated with increased risk of bleeding.

[(www.pathlab.org)](http://www.pathlab.org)

Smear the sample directly onto a slide and label with patient’s name and MRN or date of birth at the frosted end. Please label in PENCIL as ink is dissolved by subsequent staining process.

The smears must be very thinly spread and rapidly air-dried or fixed immediately (the latter smears must not be allowed to dry before applying the fixative). Rapid air dry is very important, and a hairdryer can be used on a cool setting for this purpose.

**The type of smear required depends on the type of specimen:**

**Breast**: Air dried smears.

Breast fluid could be submitted if it is haemorrhagic or suspicious in a pot.

**Thyroid**: Two or three air dried smears and one fixed smear is enough.

Thyroid cyst fluid could be sent in a pot.

**Lymph node:** Air dried smears are critical and should always is done. If metastases is suspected include one or two fixed smears in addition to the air dried smears. Needle washings could also be sent.

**Bronchial/oesophageal brushings**: Wet fixed.

Please mark all slides as “fixed” or “dried” as it is not always possible to identify how the smears have been prepared once received in the laboratory.

The remaining contents of the syringe and needle could be washed into the pot of normal saline and sent to the lab along with the slides. The slides must be placed into the slide boxes provided for transport to the laboratory. Do not place air dried and fixed smears in the same box. The sample sides of the smears should not touch each other.

An adequacy assessment service of one session per week is provided to the **EBUS** clinic.

### Joint Fluid Cytology

Fluid should be taken into a 20ml universal container and reach the laboratory as soon as possible. Minimum amount required for testing is 1ml.

### CSF Cytology

Fluid should be taken into a plain universal container and reach the laboratory as soon as possible as the specimens can degenerate rapidly.

### Bronchial Lavage/ Washings

Lavage specimens may be sent to the laboratory in the original collection container or transferred to a plain universal.

### Factors affecting tests on the above samples.

It is important to put cytology samples in the correct containers as stated above and to transport them to the laboratory as soon as possible. This is especially important with FNA samples – slides sent to us in the wrong fixative or smeared too thickly may be rendered undiagnosable

## Reports

All authorised reports are available on ICE and eCare and are sent electronically to GP systems as soon as they are authorised. (Please be aware that if the request was made on a hand written blue request card the request will not be present on eCare/ICE until final authorisation of the report.)

Hard copies of all reports are sent via the internal mail to the requesting clinician.

## Turnaround Times

Diagnostic samples identified as urgent or 2WW are prioritised in the laboratory. The reports will typically be available within 2–10 days. The department has a KPI target to report 90% of all 2WW requests within 7 days. Our performance against this target is monitored monthly and reported to the Core Clinical and Support Services Divisional Meetings.

The laboratory communicates with MDT co-ordinators, Cancer Services and Patient Pathway co-ordinators to identify priority cases and to ensure that these cases are reported in time for the relevant MDT Meeting.

Routine specimen reports will typically be available within 10-14 days; however, some may require extensive further laboratory techniques. In such cases, the reporting process may be extended.

Tissue (paraffin embedded blocks) may be referred to specialist centres to perform further tests to aid diagnosis and to provide information for treatment options and prognosis. These include specialist Immunohistochemical testing (including Her2, PD-L1,) and genetic/molecular tests (including ALK, EGFR, ROS-1, KRAS, BRAF and MMR). The expected TAT for these tests is 2–14 days.

A small number of specimen types are sent directly to specialist centres for reporting:

Bone marrow trephine biopsies are sent to The John Radcliffe Hospital, Oxford. The expected TAT for these cases is 10–14 days.

Conjunctival biopsies are sent to The Royal Hallamshire Hospital, Sheffield. The expected TAT for these cases is 5–7 days.

Skin biopsies querying alopecia are sent to Source BioScience, Nottingham. The expected TAT for these specimens is 3–4 weeks.

Skin biopsies for Direct Immunofluorescence are sent to The John Radcliffe Hospital, Oxford. The expected TAT for these specimens is 5–7 days.

On occasions it is necessary to operate with a backlog for the reporting of non-urgent specimens. At such times the reporting turnaround times will increase. The clinical information provided on the request form is used to decide which cases will be placed

into the backlog. During particularly busy times slides from the backlog may be sent off-site to Source BioScience for reporting. This outsourcing will entail an additional turnaround time of approximately 7-10 days.

## Second Opinions

Occasional samples may require a second/expert opinion. The blocks and slides from these cases are sent, as necessary, to an agreed list of nominated specialist pathologists. The full text of the second opinion together with the name of the reporting pathologist is included in the final supplementary report. The reports of second opinions will typically be available within 14 days but may be longer.

## Further Information

Further information may always be obtained from the laboratory (Ext.85819) during normal working hours (Mon - Fri 0900 to 1700hrs)

**POST MORTEMS**

When a patient dies, the **Medical Certificate of Cause of Death** should be issued as soon as possible, and the case reviewed by the Medical Examiner before the certificate is given to the family. Where a hospital post mortem is required for medical reasons, the medical certificate should be issued to relatives and their consent for the procedure obtained in writing on the appropriate consent form. The Bereavement Officer should be contacted in the first instance (extn. 86155 / bleep 1917) who will ensure that the Consent Team, consisting of clinicians and pathologists amongst others, will be available to answer any questions that the bereaved may have. Trust policies and consent forms, available on the intranet, must be followed in this regard.

## Cremation Certificates

Should the family opt to have the body cremated, then under the *Cremation Acts* *1902 & 1952, Statutory Rules and Orders 1930 & 1952*, the doctors in charge of an in-patient will be required to sign Form 4 (first part) of the Cremation Form and the medical examiner will complete the form 5 (second part). These forms are private certificates and a fee is payable upon completion via finance. These are issued and completed in the Mortuary or the Medical Examiner’s office, located in Oak House where the patient’s notes will be available. The doctor will also need to examine the deceased if they have not done so already after death. These forms must be approved by the crematorium medical referee a full 48 hrs before the cremation can take place. This can extend to 72 hrs if the family have requested to see the form). There is often considerable urgency as a funeral cannot go ahead without the completion of the forms thus it is important that any doctor asked to fill in a form should do so **as soon as conveniently possible**.

**Further Information**

Mortuary Technicians are available during normal hours on Ext. 85828, or by Direct Dial on (01908) 995828. Out of hours, contact switchboard and ask them to contact the on call Mortuary Technician.

## Referral to the Coroner

The coroner is an officer of the Crown, whose duty it is to inquire into all violent, unnatural or sudden deaths of which the cause is doubtful or unknown. Should the death of the patient fall into any of the criteria listed below, the death should be reported through the coroners officer on 01908 254326. Not all deaths reported to the coroner will require a post-mortem.

All deaths from the following should be reported:

1. The cause of death is unknown.

2. It cannot be readily certified as being natural causes.

3. The deceased was not attended by the doctor during his last illness or was not seen within 14 days or viewed after death.

4. There are any suspicious circumstances or a history of violence.

5. The death may be linked to an accident (whenever it occurred).

6. There is a question of self-neglect or neglect by others.

7. The death has occurred, or the illness arisen during or shortly after detention in police or prison custody (including voluntary attendance at a police station).

8. The deceased was detained under the mental health act.

9. The death is linked with an abortion

10. The death might have been contributed to by the actions of the deceased (such as history of drug or solvent abuse, self-injury or overdose)

11. The death could be due to industrial disease, or related in any way to the deceased’s employment

12. The death occurred during an operation or before full recovery from the effects of an anaesthetic or was in any way related to the anaesthetic (in any event a death within 24 hours should normally be referred).

13. The death may be related to a medical procedure or treatment, whether invasive or not.

14. The death may be due to a lack of medical care.

15. There are any other unusual or disturbing features to the case.

16. The death occurs within 24 hours of admission to hospital.

17. It may be wise to report any death where there is an allegation of medical mismanagement.

18. Any maternal death relating to pregnancy or childbirth.

19. Any stillbirth.

20. The deceased had *Clostridium difficile* or *Legionella.*

21. A Deprivation of Liberty is in place.

If there is any doubt, the Coroner, his officers or one of the consultant pathologists should be contacted without delay. This list may be amended occasionally so if there is any doubt, please contact the mortuary staff or bereavement office. If the case is referred to the Coroner, and he requests a post mortem, the doctor is not required to issue a Death Certificate. The post mortem report or the cause of death is available from the coroner’s office. Telephone number 01908 254326

**CHEMICAL PATHOLOGY**

**INTRODUCTION**

Chemical Pathology offers both an analytical and advisory service. Most routine tests are performed and reported the same day. Critically abnormal results are telephoned. Urgent requests for the common tests are performed within one hour of receipt. Some requests are always treated as urgent requests and are processed as soon as possible, e.g. Blood gas analysis is processed within ten minutes.

Details of available tests, normal ranges, sample requirements and turnaround times are in the attached test repertoire list.

## Out of Hours Emergency service

Tests routinely available ‘out of hours’ are: -

* Renal profiles
* Liver profiles
* Bone profiles
* Troponin
* Glucose
* CRP
* Paracetamol & Salicylate
* Serum & Urine Osmolality
* CSF Glucose & Protein
* Amylase
* BNP
* Blood Gases

Other tests may be available depending on clinical need and should be discussed with the ‘on-call’ BMS or the Consultant Chemical Pathologist.

Results are available from eCare/ ICE and are not routinely telephoned unless critically abnormal.

## Sample Containers

### Routine blood samples

Sample requirements are detailed individually in the test repertoire. Usually one gold bottle filled to the mark is sufficient for a full range of routine tests including hormone profiles.

### Glucose samples

Where the sample for glucose estimation is likely to be greater than 3 hours old before analysis, the use of a grey fluoride oxalate bottle is strongly recommended. Delay in receipt of sample for testing will produce low results unless taken into fluoride oxalate

### ‘Special’ containers/collection conditions

Some of the tests in the repertoire require special containers or collection conditions; these vary depending on the referral centre and are subject to change. Please contact Pathology Support Unit (Ext. 85842) who will be pleased to supply the current container and collection conditions.

### Urine samples

(i) Random urines should be sent in a white top universal bottle. Samples for reducing substances MUST reach Pathology within 2 hours of collection.

(ii) Timed/24 hour urines - containers are available from Pathology on request. Please state which investigation is required as the preservative added will vary with investigations. An instruction sheet detailing the collection conditions is issued with each set of 24 hour urine containers.

### Faecal samples

Random samples in blue top universal for Calprotectin analysis

Samples for FIT analysis should use the appropriate FIT collection device.

### CSF samples

In white top universals labelled CSF ONLY.

### Sweat samples

A specialised collection system is used by the Paediatric Department.

### Other fluids/calculi

Fluids should be collected into a gold top gel tube and calculi into a white top

universal.

### Blood gas samples

(i) Warn the laboratory that the sample is about to be taken.

(ii) Use a special heparinised blood gas syringe.

(iii) Collect the sample - Adults at least 2.0 mL - Paediatrics at least 0.5 mL.

1. Label sample syringe with patient’s eCare request
2. Dispose of the needle into a ‘sharps’ box
3. Purge the syringe of air.

**Never send a syringe with the needle attached. This is a major hazard to those transporting the sample and to those analysing the sample. Samples with needles still attached will not be analysed and will be discarded.**

**Do not send blood gas samples via air tube**

1. Fit a blanking hub to syringe.

(vii) Mix the sample by rolling the syringe barrel between both hands.

(viii) Place the syringe in a separate plastic bag containing **crushed** ice and seal.

(ix) Complete the eCare request and place the sample in a separate plastic bag in the plastic bag attached to the card and seal.

(x) Send to Pathology via the porters to arrive within 15 minutes of collection.

Capillaries are available which are ‘triple-heparinised’ and do not require mixing but need sealing with the plastic end caps supplied and should be taped to a wooden spatula for support and protection before being transported.

## Antibiotic Assays

The Antibiotic assays Gentamicin and Vancomycin are analysed by Chemical Pathology; advice on times of collection and interpretation of results is available from the Consultant Microbiologist.

## Downs Syndrome Screening

The Down’s Screening portion of the Antenatal request card MUST be fully completed for analysis and interpretation this test is now referred to Oxford.

## Dynamic Function Tests

Protocols for the following dynamic functions tests are held by the Pathology Support Unit and are available on request:

* Investigation of Primary Aldosteronism
* Cryoglobulin collection
* Dexamethasone (overnight suppression) test (screening for Cushing’s)
* Dexamethasone (high dose) test (cause of Cushing’s)
* Oral Glucose Tolerance test
* Growth Hormone (GTT suppression) test
* Investigation of Unexplained Hyperkalaemia
* Porphyria - investigation and sample collection
* Synacthen (short) test
* Synacthen (long) test
* TRH test
* Water Deprivation test
* Xylose Absorption test
* Urine Metanephrines - issued with collection bottles
* Urine 5HIAA - issued with collection bottles if 24hr urine required following abnormal urine spot test.

## Therapeutic Drug Monitoring

Typically, samples should be taken as trough levels i.e. pre-dose.

Digoxin should be taken 6 - 8 hours post dose, please state on card.

For Antibiotics see above.

## Analytical Instruments in Clinical Areas

The Blood Gas analysers on DOCC, A&E, MAU, NNU Lung Function, W15 and Maternity may be used ONLY by staff trained and approved by Pathology **Point of Care Staff.**

## Xanthochromia

One mL of CSF is the minimum required for this test.

CSF for Xanthochromia should be the last specimen taken – ideally it should be protected from the light and brought to the laboratory as soon as possible.

Samples **must not** be sent via the vacuum tube transport system.

The initial processing of the specimen in Biochemistry needs to be done **within one hour** of collection.

The Xanthochromia service operates between 09:00 and 17:00 during the week and between 09:00 and 12.30 on Saturdays.,

Useful clinical details to aid interpretation of results:

1. Time of onset of symptoms.

2. Time LP taken – should be no less than 12 hours after onset of

symptoms, otherwise false negative results can occur.

3. Does differential diagnosis include meningitis?

4. Requests should clearly indicate the name and number of the

person requiring the results.

## Key factors Which May Affect results in Chemical Pathology

Several non-analytical factors may affect the performance of individual chemistry tests, these include the type of collection bottle used, the time taken for a sample to arrive in the laboratory, interfering substances such as high levels of lipids (lipaemia) and bilirubin (icterus), haemolysis and drug interference. Some common examples of these factors are listed below:

**Collection Bottle:**

**Serum only Lithium heparin plasma Fluoride oxalate plasma**

*Lithium, Tumour Ammonia Lactate*

*Markers, Digoxin Glucose(optional)*

*Protein electrophoresis.*

**Sample Separation Delay:**

Unsuitable on arrival if:

>30 mins old > 3 hrs old > 1 day old >2 days old

*Lactate, Troponin Alcohol Tumour markers*

*Ammonia, Glucose (if not Downs*

*Blood Gases in a fluoride oxalate*

*Bottle) Magnesium*

*Phosphate*

*Potassium*

**Lipaemia:**

Iron, Glucose, IgM Progesterone

**Icterus:**

Paracetamol, Cholesterol, Cortisol, FSH, Lactate, Progesterone, Testosterone, Triglyceride, Urea

**Haemolysis:**

AST, Ammonia, Bile Acids, Cholesterol, CK, Iron, LDH, Magnesium, Osmolality, Phosphate, Potassium, Total Protein, Vitamin D.

**Drug Interference:**

Many drugs interfere with Biochemistry tests, either through their interaction with the analyte to be measured or through interference with the method of analysis used. Some common examples include:

Digibind interferes with Digoxin estimation; Steroid therapy will affect the Cortisol level detected. It is therefore important to consider the drug therapy when interpreting results.

## Biochemistry Results from ‘Cardiac Arrest’ Patients

These samples are of course given absolute priority over **all** other samples, by

all members of Pathology staff.

The following, if observed by all concerned, should reduce any possible delays:

1. Requests must be made for U&E, Creatinine, Calcium & Glucose ONLY and sent in a GREEN (Lithium Heparin Gel)bottle. This facilitates the rapid centrifugation of samples. Gold (Clotted Blood Gel) bottle MUST NOT BE USED, as the analysis of these tests cannot be performed until the sample has clotted, typically 10 - 20 minutes. Request for Blood Gases may also be made and the appropriate sample sent. A paper copy of the request should be sent with the sample.

2. Requests for any other tests on these patients MUST be requested on a separate number accompanied by a separate sample, this again will remove any possible delay caused by longer processing time required for these additional tests. These requests will be dealt with as ‘URGENT’ samples and given high priority.

3. The Pathology Support Unit MUST be aware of these very urgent samples, either by phone or directly by a person handing the sample to reception staff, i.e. giving the sample to a member of Pathology staff and identifying it as being from an ‘arrest’ patient, rather than leaving the sample on the reception counter or in the ‘on call’ box for someone to find. Of course, should the lab be expecting the sample, (via a phoned warning) then lab staff will be waiting for it. Merely writing ‘arrest’ on the request form **CANNOT** be considered as adequate notification.

4. All ‘arrest’ samples are given absolute priority by reception staff; they will number the samples if necessary and take them directly to the appropriate analytical area.

5. The scientific staff will centrifuge (where appropriate) and analyse for blood gases, U&E, Creatinine, Calcium & Glucose as a priority over **all** other samples awaiting analysis.

6. The results will be relayed to the appropriate ward or department immediately.

**LIST OF TESTS AND INFORMATION**

The sample stability indicates the time limit the sample will be valid for the test to be added if requested at a later date. If additional tests are required, please check with the laboratory on availability and sample volume. Urgent requests for routine tests will be processed within 1 hour of receipt or less. Inpatient requests for routine testing will be processed the same day. Those tests with short sample stability such as Ammonia or Blood gas analysis will be processed within the sample stability time displayed.

Expected turnaround times are valid during normal working hours 9am to 5pm Monday to Friday unless the samples is designated as urgent and agreed with the laboratory. In these cases, the turnaround time is valid regardless of sample receipt date and time.

The list below contains those tests performed on site by the Biochemistry department and includes the container requirements. Most tests can all be done on one 3.5 ml gold bottle if full. If in doubt, please contact the laboratory for advice.

| **TEST** | **UNITS** | **BOTTLE / CONTAINER** | **ANALYSIS** | **SAMPLE STABILITY** | **EXPECTED TURNAROUND TIME FROM RECEIPT OF SAMPLE** | **NORMAL / THERAPEUTIC RANGE** | | |
| --- | --- | --- | --- | --- | --- | --- | --- | --- |
| **123** |  |  |  |  |  |  |  |  |
| % OXYGEN SATURATION | % | BLOOD GAS SYRINGE (On ice) | On Demand | 30 mins | 20 mins – always urgent | 95 | - | 98 |
| **A** |  |  |  |  |  |  |  |  |
| ACTUAL BICARBONATE | mmol/L | BLOOD GAS SYRINGE (On ice) | On Demand | 30 mins | 20 mins – always urgent | 20.0 | - | 26.0 |
|  |  |  |  |  |  |  |  |  |
| AFP-SERUM-TUMOUR MARKER\* | kU/L | GOLD | Daily | 48 hrs | 48 hours | less than 9 |  |  |
| ALBUMIN | g/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | 35 | - | 50 |
| ALK PHOSPHATASE | iU/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | –  30 |  | –  130 |
| ALT\* | iU/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | Female <35 | - | Male <50 |
| AMMONIA | umol/L | GREEN (Lith Hep) on ICE | On Demand | 30 mins | 30 mins – always urgent | 6 | - | 47 |
| AMYLASE-SERUM\* | iU/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | 28 | - | 100 |
| ANGIOTENSIN CONVERTING ENZYME\* | nmol/min/mL | GOLD | Daily | 48 hrs | 3 hours | 8 | - | 52 |
| AST\* (needs updating) | iU/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | Female <35 |  | Male <50 |
| **B** |  |  |  |  |  |  |  |  |
| B-HCG-PREGNANCY | iU/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine |  | | |
| B-HCG-SERUM-TUMOUR MARKER | iU/L | GOLD | Daily | 48 hrs | 48 hours | less than 5 |  |  |
| B2MICROGLOBULIN\* | g/L | GOLD / GREEN | Daily | 48 hrs | 48 hours | 0.8 | - | 2.4 < 60yrs  <3.1 > 59yrs |
| BASE EXCESS | mmol/L | BLOOD GAS SYRINGE (On ice) | On Demand | 30 mins | 20 mins – always urgent | -2.0 | - | 2.0 |
| BICARBONATE | mmol/L | GOLD / GREEN | Daily | 24 hrs | 1 hour – urgent  3 hours - routine | 22 | - | 29 |
| BILIRUBIN-CONJUGATED \* | umol/l | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | 0 |  | 3.4 |
| BILIRUBIN-TOTAL | umol/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | 3 | - | 21 |
| BILIRUBIN-UNCONJUGATED | umol/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | 3 | - | 14 |
| BILE ACIDS\* | umol/L | GOLD | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | 0 | - | 12 |
| **BNP\* (available for cardiology/medicine consultants only)** | pg/ml | LAVENDER | Daily | 4 hrs | 1 hour – urgent  3 hours - routine |  |  | <100 |
| **C** |  |  |  |  |  |  |  |  |
| C.S.F LACTATE | mmol/L | GREY (Fluoride) | On Demand | 1 hr | 1 hour | Less than 2.8 |  |  |
| C.S.F. GLUCOSE | mmol/L | WHITE UNIVERSAL | On Demand | 1 hr | 1 hour | 2.5 | - | 4.5 |
| C.S.F. PROTEIN | g/L | WHITE UNIVERSAL | On Demand | 1 hr | 1 hour | 0.15 | - | 0.45 |
| CA 19-9\* | U/ml | GOLD | Daily | 48 hrs | 48 hours | 0 | - | 35 |
| CA-125\* | U/ml | GOLD | Daily | 48 hrs | 48 hours | 0 | - | 35 |
| CA-153\* | U/ml | GOLD | Daily | 48 hrs | 48 hours | 0 | - | 23 |
| CALCIUM | mmol/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | 2.20 | - | 2.65 |
| CALCIUM CORRECTED | mmol/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | 2.20 | - | 2.60 |
| CALCIUM OUTPUT-URINE | mmol/24hr | 24HR URINE | Within 4 days | 1 week | 5 days | 2.5 | - | 7.5 |
| CALPROTECTIN (FAECAL) \*1 | ug/g | BLUE UNIVERSAL | Within 4 days | 3 days before extraction, 6 days post. | 5 days | Normal  Equivocal  Positive |  | <50  50 – 200  >200 |
| CARBAMAZEPINE | mg/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | 4 |  | 12 |
| CARBOXYHAEMOGLOBIN | % | BLOOD GAS SYRINGE/EDTA | On Demand | 30 mins | 20 mins – always urgent | Interpretation on screen with results | | |
| CEA\* | ug/L | GOLD | Daily | 48 hrs | 48 hours | less than 6 |  |  |
| CHLORIDE | mmol/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | 95 | - | 108 |
| CHOLESTEROL-TOTAL\* | mmol/L | GOLD / GREEN | Daily | 48 hrs | 3 hours | 0 |  | 5.2 |
| CHOLESTEROL TOTAL/HDL RATIO |  | GOLD / GREEN | Daily | 48 hrs | 3 hours |  |  |  |
| CHOLESTEROL-HDL | mmol/L | GOLD / GREEN | Daily | 48 hrs | 3 hours |  |  |  |
| CHOLESTEROL-LDL (Calculated) | mmol/L | GOLD / GREEN | Daily | 48 hrs | 3 hours |  |  |  |
| CK | iU/L | GOLD / GREEN | Daily | 12hrs | 1 hour – urgent  3 hours - routine | Female25-200 | - | Male 40-320 |
| CORTISOL-SERUM\* | nmol/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  48 hours - routine | 185 | - | 624 at 09:00 am |
| CREATININE\* | umol/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | Female 49 – 90 | - | Male 64 - 104 |
| CREATININE CLEARANCE | ml/min | 24HR URINE+GOLD | 48 hrs | 1 week | 5 days | Interpretation on screen |  |  |
| CREATININE OUTPUT-URINE | umol/24hr | 24HR URINE | 48 hrs | 1 week | 5 days | Female 7000 - 13000 |  | Male 13000 - 18000 |
| CRP\* | mg/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine |  | - | <5 |
| **D** |  |  |  |  |  |  |  |  |
| DIGOXIN | ug/L | GOLD | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | 0.5 – 1.0 |  | if >6hrs and <24hrs post dose |
| **E** |  |  |  |  |  |  |  |  |
| ELECTROPHORESIS-SERUM |  | GOLD | Within 1 week | 10 days | 10 days |  |  |  |
| ELECTROPHORESIS-URINE |  | WHITE UNIVERSAL | Within 1 week | 10 days | 10 days |  |  |  |
| ETHANOL-SERUM | mg/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | Interpretation on screen with results | | |
| ETHANOL-URINE |  | WHITE UNIVERSAL | Daily | 48 hrs | 1 hour – urgent  3 hours - routine |  |  |  |
|  |  |  |  |  |  |  |  |  |
| **F** |  |  |  |  |  |  |  |  |
| F.S.H. | U/L | GOLD / GREEN | Daily | 48 hrs | 48 hours | –Interpretation on screen |  |  |
| FAECAL IMMUNOCHEMICAL TESTING (FIT)\*2 | ug/g | FIT COLLECTION DEVICE | Within 4 days | 14 days | 7 days |  |  | <10 |
|  |  |  |  |  |  |  |  |  |
| FLUID PROTEIN/GLUCOSE |  | GOLD | Daily | 48 hrs | 5 hours |  |  |  |
| FREE T3\* | pmol/L | GOLD / GREEN | Daily | 48 hrs | 48 hours | 3.8 | - | 6.0 |
| FREE T4\* | pmol/L | GOLD / GREEN | Daily | 48 hrs | 48 hours | 7.9 | - | 14.4 |
|  |  |  |  |  |  | 1st trimester 6.67 | - | 14.12 |
|  |  |  |  |  |  | 2nd trimester 5.79 | - | 12.7 |
|  |  |  |  |  |  | 3rd trimester 6.11 | - | 12.2 |
| **G** |  |  |  |  |  |  |  |  |
| GAMMA GT\* | iU/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | Female <38 |  | Male <55 |
| GENTAMICIN | mg/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | See Trust Antibiotic Policy on the Intranet |  |  |
| GLOBULIN | g/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | 20 | - | 42 |
| GLUCOSE\*3 | mmol/L | GOLD / GREEN / GREY | Daily | 48 hrs if fluoride oxalate | 1 hour – urgent  3 hours - routine | 3.5 | - | 7.7 |
| GROWTH HORMONE\* | ug/L | GOLD / GREEN | Daily | 48 hrs | 48 hours | Male 0 | - | 1.0 |
|  |  |  |  |  |  | Female 0 | **-** | 3.6 |
| GLYCOSALATED HAEMGLOBIN\* | mmol/mol | LAVENDER | 48 hrs | 1 week | 4 days | 20 |  | 42 |
| **H** |  |  |  |  |  |  |  |  |
| HAEMOGLOBIN A1c\* | % | LAVENDER | 48 hrs | 1 week | 4 days | 4.0 | - | 6.0 |
| **I, J, K** |  |  |  |  |  |  |  |  |
| IgA\*4 | g/L | GOLD / GREEN | Within 4 days | 1 week | 5 days | Aged<2wks |  | 0.01 - 0.08 |
|  |  |  |  |  |  | Aged 2 - 6wks |  | 0.02 - 0.15 |
|  |  |  |  |  |  | Aged 2 – 3mths |  | 0.05 - 0.4 |
|  |  |  |  |  |  | Aged 3 - 6mths |  | 0.10 - 0.5 |
|  |  |  |  |  |  | Aged 6 - 9mths |  | 0.15 - 0.7 |
|  |  |  |  |  |  | Aged 9 - 12mths |  | 0.2 - 0.7 |
|  |  |  |  |  |  | Aged 1 - 2yrs |  | 0.3 - 1.2 |
|  |  |  |  |  |  | Aged 2 - 3yrs |  | 0.3 - 1.3 |
|  |  |  |  |  |  | Aged 3 - 6yrs |  | 0.4 - 2.0 |
|  |  |  |  |  |  | Aged 6 - 9yrs |  | 0.5 - 2.4 |
|  |  |  |  |  |  | Aged 9 - 12yrs |  | 0.7 - 2.5 |
|  |  |  |  |  |  | Aged 12 - 45yrs |  | 0.8 - 2.8 |
|  |  |  |  |  |  | Aged >45yrs |  | 0.8 - 4.0 |
| IgG\*4 | g/L | GOLD / GREEN | Within 4 days | 1 week | 5 days | Aged <2wks |  | 5.0 - 17.0 |
|  |  |  |  |  |  | Aged 2 - 6wks |  | 3.9 - 13.0 |
|  |  |  |  |  |  | Aged 2 – 3mths |  | 2.1 - 7.7 |
|  |  |  |  |  |  | Aged 3 - 6mths |  | 2.4 - 8.8 |
|  |  |  |  |  |  | Aged 6 - 9mths |  | 3.0 - 9.0 |
|  |  |  |  |  |  | Aged 9 - 12mths |  | 3.0 - 10.9 |
|  |  |  |  |  |  | Aged 1 - 2yrs |  | 3.1 - 13.8 |
|  |  |  |  |  |  | Aged 2 - 3yrs |  | 3.7 - 15.8 |
|  |  |  |  |  |  | Aged 3 - 6yrs |  | 4.9 - 16.1 |
|  |  |  |  |  |  | Aged 6 - 45yrs |  | 5.4 - 16.1 |
|  |  |  |  |  |  | Aged >45yrs |  | 6.0 - 16.0 |
| IgM\*4 | g/L | GOLD / GREEN | Within 4 days | 1 week | 5 days | Aged<2wks |  | 0.05 - 0.2 |
|  |  |  |  |  |  | Aged 2 - 6wks |  | 0.08 - 0.4 |
|  |  |  |  |  |  | Aged 2 – 3mths |  | 0.15 - 0.7 |
|  |  |  |  |  |  | Aged 3 - 6mths |  | 0.2 - 1.0 |
|  |  |  |  |  |  | Aged 6 - 9mths |  | 0.4 - 1.6 |
|  |  |  |  |  |  | Aged 9 - 12mths |  | 0.6 - 2.1 |
|  |  |  |  |  |  | Aged 1 - 3yrs |  | 0.5 - 2.2 |
|  |  |  |  |  |  | Aged 3 - 6yrs |  | 0.5 - 2.0 |
|  |  |  |  |  |  | Aged 6 - 12yrs |  | 0.5 - 1.8 |
|  |  |  |  |  |  | Aged 12 - 45yrs |  | 0.5 - 1.9 |
|  |  |  |  |  |  | Aged >45yrs |  | 0.5 - 2.0 |
| IGE\*4 | kU/L | GOLD / GREEN | Within 4 days | 1 week | 5 days | Aged <1wks |  | 0-5 |
|  |  |  |  |  |  | Aged 1 - 14wks |  | 0-11 |
|  |  |  |  |  |  | Aged 14wks - 1yr |  | 0-29 |
|  |  |  |  |  |  | Aged 1 - 5yrs |  | 0-52 |
|  |  |  |  |  |  | Aged 5 - 10yrs |  | 0-63 |
|  |  |  |  |  |  | Aged 10 -15yrs |  | 0-75 |
|  |  |  |  |  |  | Adult |  | 0-100 |
| IMMUNOFIXATION-SERUM |  | GOLD | Within 1 week | 2 weeks | 15 days from request |  |  |  |
| IMMUNOFIXATION-URINE |  | WHITE UNIVERSAL | Within 1 week | 2 weeks | 15 days from request |  |  |  |
| IRON-SERUM\* | umol/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | Female 10.7 – 32.2 | - | Male 12.5 – 32.2 |
|  |  |  |  |  |  |  |  |  |
| **L** |  |  |  |  |  |  |  |  |
| L.H. | U/L | GOLD / GREEN | Daily | 48 hrs | 48 hours | Interpretation on screen | - |  |
| LACTATE | mmol/L | GREY on ICE | On Demand | 30 minutes | 30 mins – always urgent | 0.6 | - | 2.5 |
| LDH\* | iU/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | 135 | - | 360 |
| LITHIUM | mmol/L | GOLD | Daily | 48 hrs | 1 hour – urgent  24 hours - routine |  | - | Interpretation on screen with results |
| **M** |  |  |  |  |  |  |  |  |
| MAGNESIUM | mmol/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | 0.7 | - | 1.0 |
| MET-HAEMOGLOBIN | % | BLOOD GAS SYRINGE (On ice) | On Demand | 30 mins | 30 mins – always urgent | 0 | - | 1 |
| METHOTREXATE | umol/L | GOLD / GREEN | On Demand | 48 hrs | 2 hours – urgent  2 days routine | **For high dose methotrexate testing 2 days prior notice is required by the laboratory. It may be possible to process samples urgently following discussion with the laboratory** | | |
| MICROALBUMIN - URINE | mg/L | WHITE UNIVERSAL | Daily | 48 hrs | 2 days |  | | |
| MICROALBUMIN CREATININE RATIO | mg/mmol | WHITE UNIVERSAL | Daily | 48 hrs | 2 days | Female 0 – 3.4 |  | Male 0 – 2.4 |
| MYOGLOBIN-URINE |  | WHITE UNIVERSAL | On Demand | 48 hrs | 2 days |  |  |  |
|  |  |  |  |  |  |  |  |  |
| **N** |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |
| **O** |  |  |  |  |  |  |  |  |
| OESTRADIOL | pmol/L | GOLD / GREEN | Daily | 48 hrs | 48 hours |  |  | Interpretation on screen with results |
| OSMOLALITY-SERUM | mosmol/kg | GOLD / GREEN | Daily | 48 hrs | 48 hours | 275 | - | 295 |
| OSMOLALITY-URINE | mosmol/kg | WHITE UNIVERSAL | Daily | 48 hrs | 48 hours | 300 | - | 900 |
| **P, Q** |  |  |  |  |  |  |  |  |
| P.S.A. (Must be spun within 24 hrs)\* | ug/L | GOLD | Daily | 48 hrs | 48 hours | <4.0 |  |  |
| p02 | kPa | BLOOD GAS SYRINGE (On ice) | On Demand | 30 mins | 20 mins – always urgent | 10.0 | - | 13.0 |
| PARACETAMOL | mg/L | GOLD / GREEN | On Demand | 48 hrs | 1 hour – urgent  3 hours - routine | Follow national guidelines | | |
| PARAQUAT SCREEN |  | WHITE UNIVERSAL | On Demand | 1 week | 3 days, sample sent to referral lab |  |  |  |
| pC02 | kPa | BLOOD GAS SYRINGE (On ice) | On Demand | 30 mins | 20 mins – always urgent | 4.7 | - | 6.0 |
| pH - BLOOD |  | BLOOD GAS SYRINGE (On ice) | On Demand | 30 mins | 20 mins – always urgent | 7.35 | - | 7.45 |
| PHENOBARBITONE | mg/L | GOLD / GREEN | Daily | 24 hrs | 1 hour – urgent  3 hours – routine | 10- 40 | | |
| PHENYTOIN | mg/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | 5 - 20 | | |
| PHOSPHATE | mmol/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | 0.8 | - | 1.5 |
| PHOSPHATE OUTPUT-URINE | mmol/24hr | 24HR URINE | 48 hrs | 1 week | 5 days | 16 | - | 48 |
| POTASSIUM | mmol/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | 3.5 | - | 5.3 |
| POTASSIUM OUTPUT-URINE | mmol/24hr | 24HR URINE | 48 hrs | 1 week | 5 days | 25 | - | 125 |
| PROGESTERONE | nmol/L | GOLD / GREEN | Daily | 48 hrs | 48 hours |  |  |  |
| PROLACTIN | mU/L | GOLD / GREEN | Daily | 48 hrs | 48 hours |  |  | –  Interpretation on screen with results |
| PROTEIN CONCENTRATION-URINE | g/L | WHITE UNIVERSAL | Daily | 48 hrs | 1 day | 0.05 | - | 0.08 |
| PROTEIN OUTPUT-URINE | g/24hr | 24HR URINE | 48 hrs | 1 week | 5 days | less than 0.15 |  |  |
| PROTEIN- TOTAL | g/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | 60 | - | 80 |
| PTH\* | pmol/l | GOLD/ GREEN | Daily | 48 hrs | 48 hours | 1.3 | - | 9.3 |
| **R** |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |
| **S** |  |  |  |  |  |  |  |  |
| SALICYLATE | mg/L | GOLD / GREEN | On Demand | 48 hrs | 1 hour – urgent  3 hours - routine | Interpretation on screen with results | | |
| SEX HORMONE BINDING GLOBULIN\* |  | GOLD / GREEN | Daily | 48 hrs | 48 hours | Female 16.8 – 135.6 | - | Male 13.3 – 89.5 |
| SODIUM | mmol/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | 133 | - | 146 |
| SODIUM OUTPUT-URINE | mmol/24hr | 24HR URINE | 48 hrs | 1 week | 5 days | 40 | - | 220 |
| SWEAT CHLORIDE AND SODIUM | mmol/l | SWEAT COLLECTOR | On Demand | 48 hrs | 2 days | Interpretation on screen |  |  |
| **T** |  |  |  |  |  |  |  |  |
| T.S.H.\* | mU/L | GOLD / GREEN | Daily | 48 hrs | 48 hours | 0.38 | - | 5.33 |
|  |  |  |  |  |  | TSH Pregnancy Related Reference Ranges: | | |
|  |  |  |  |  |  | 1st Trimester 0.05 | - | 3.70 |
|  |  |  |  |  |  | 2nd Trimester 0.31 | - | 4.35 |
|  |  |  |  |  |  | 3rd Trimester 0.41 | - | 5.18 |
| TESTOSTERONE | nmol/L | GOLD / GREEN | Daily | 48 hrs | 48 hours | Male (Adult) 6.1 | - | 27.1 |
|  |  |  |  |  |  | Male (18-30) 9.0 | - | 28.3 |
|  |  |  |  |  |  | Male (31-44) 6.9 | - | 23.6 |
|  |  |  |  |  |  | Male (45-66) 5.2 | - | 23.7 |
|  |  |  |  |  |  | Female 0 – 3.0 |  |  |
| THEOPHYLLINE | mg/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | Aged <7wks |  | 5-10 i.e. neonatal apnoea |
|  |  |  |  |  |  | Aged >7wks |  | 10-20 i.e. asthma etc |
| THYROID MICROSOMAL ANTIBODIES |  | GOLD / GREEN | Daily | 48 hrs | 48 hours |  |  |  |
| TRANSFERRIN\* | g/L | GOLD / GREEN | Daily | 48 hrs | 48 hours | 2.0 | - | 3.6 |
| TRIGLYCERIDES | mmol/L | GOLD / GREEN | Daily | 48 hrs | 48 hours |  |  |  |
| TROPONIN I\* (Not practical from H.C.) must be received within 3 hrs | ng/L | GOLD / GREEN | On Demand | 48 hrs once spun | 1 hour –always urgent | Female<11.7 |  | Male <19.9 |
| **U** |  |  |  |  |  |  |  |  |
| UREA | mmol/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – always urgent | 2.5 | - | 7.8 |
| UREA OUTPUT-URINE | mmol/24hr | 24HR URINE | 48 hrs | 1 week | 5 days | 170 | - | 580 |
| URIC ACID | umol/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | Female 140 - 360 | - | Male 200 - 430 |
| URINE ELECTROLYTES |  | WHITE UNIVERSAL | Daily | 48 hrs | 2 days |  |  |  |
| URINE GLUCOSE |  | WHITE UNIVERSAL | Daily | 48 hrs | 2 days |  |  |  |
| URINE KETONES |  | WHITE UNIVERSAL | Daily | 48 hrs | 2 days |  |  |  |
| URINE PH |  | WHITE UNIVERSAL | Daily | 48 hrs | 2 days |  |  |  |
| URINE BILIRUBIN |  | WHITE UNIVERSAL | Daily | 48 hrs | 2 days |  |  |  |
| URINE UROBILINOGEN |  | WHITE UNIVERSAL | Daily | 48 hrs | 2 days |  |  |  |
| **V** |  |  |  |  |  |  |  |  |
| VALPROIC ACID | mg/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | 50 |  | 100 |
| VANCOMYCIN | mg/L | GOLD / GREEN | Daily | 48 hrs | 1 hour – urgent  3 hours - routine | See Trust Antibiotic Policy on Intranet |  |  |
| VITAMIN D | nmol/L | GOLD / GREEN | Daily | 48 hrs | 48 hours | Interpretation on screen |  |  |

These tests are referred to other hospitals for analysis. They have variable turnaround times, and some can take up to 4 weeks to return. Contact the laboratory if you have a specific query relating to turnaround times and please check for outstanding results before requesting. These tests are always costly - consider other in house assays if possible. Unless otherwise stated the samples are stable for 48 hrs post collection. Reporting units and Normal / Therapeutic ranges are reported in accordance with referral laboratory protocols and are included in any eCare/ ICE / paper reports

|  |  |  |  |
| --- | --- | --- | --- |
| **TEST** | **BOTTLE / CONTAINER** | **ANALYSIS** | **SAMPLE STABILITY (if applicable)** |
| **LOCATION** |
| **1 2 3 etc** |  |  |  |
| 11-DEOXY CORTISOL | GOLD | ST THOMAS |  |
| 17-OH PROGESTERONE | GOLD | LEEDS |  |
|  |  |  |  |
|  |  |  |  |
|  |  |  |  |
| 17-OH PROGESTERONE - Saliva | universal | LEEDS |  |
| Saliva 17HP (pre-supper) | universal | LEEDS |  |
| Saliva 17HP (pre-breakfast) | universal | LEEDS |  |
| Saliva 17HP (pre-lunch) | universal | LEEDS |  |
| 17-OH PROGESTERONE Blood spots | Blood spots card | CARDIFF |  |
| 5.H.I.A.A OUTPUT | 24HR URINE - Acid container | OXFORD |  |
| **A** |  |  |  |
| ACETYLCHOLINE RECEPTOR ABS | GOLD | OXFORD |  |
| ACTH | LAVENDER (On ice) | BARTS | 15 mins |
| ACYLCARNITINE | BLOOD SPOT | SHEFFIELD |  |
| ADRENAL ANTIBODIES | GOLD | OXFORD |  |
| ALDOSTERONE - only done with RENIN | 10mL LITH HEP bottle | CHARING CROSS |  |
| ALK PHOS ISOENZYMES | GOLD | ROYAL FREE |  |
| ALPHA-1-ANTI TRYPSIN | GOLD | SHEFFIELD |  |
| ALPHA-1-ANTI TRYPSIN GENOTYPE | LAVENDER | SHEFFIELD |  |
| ALPHA-1-ANTI TRYPSIN IN FAECES | BLUE FAECES CONTAINER | ST GEORGES | 3Hrs |
| ALPHA-1-ACID GLYCOPROTEIN | GOLD | SHEFFIELD |  |
| ALPHA AMINO ADIPIC SEMIALDEHYDE | WHITE UNIVERSAL (urine sample) | ICH |  |
| ALUMINIUM | ROYAL BLUE (Trace metal bottle) | CARDIFF |  |
| AMINO ACIDS-PLASMA | ORANGE (Lith Hep) | SHEFFIELD |  |
| AMINO ACIDS – URINE | WHITE UNIVERSAL (urine) | SHEFFIELD |  |
| AMNIOTIC FLUID | WHITE UNIVERSAL | OXFORD |  |
| AMPHETAMINE (urine drug screen) | WHITE UNIVERSAL (urine) | KINGS |  |
| AMYLASE ISOENZYMES | SERUM | GREAT ORMOND ST |  |
| AMA - SUBTYPE 2 | SERUM | OXFORD |  |
| AMYLOID A | SERUM | SHEFFIELD |  |
| ANCA | GOLD | NORTHAMPTON |  |
| ANDROSTENEDIONE | GOLD | LEEDS |  |
| ANTI BASAL GANGLIA AB | GOLD | QUEENS SQUARE |  |
| ANTI CARDIOLIPIN ABS | GOLD | NORTHAMPTON |  |
| ANTI GAD ABS | GOLD | OXFORD |  |
| ANTI GLOMERULAR ABS | GOLD | OXFORD |  |
| ANTI GANGLIOSIDE AB (GQ1B) | GOLD | OXFORD |  |
| ANTI GLYCINE RECEPTOR AB | GOLD | OXFORD |  |
| ANTI MAG ATOX | GOLD | OXFORD |  |
| ANTI MULLERIAN HORMONE | GOLD | GLASGOW |  |
| ANTI NEURONAL AB (anti HU and anti RI) | GOLD | OXFORD |  |
| ANTI NUCLEAR ANTIBODIES | GOLD | NGH | 7 days |
| ANTI MUSK ANTIBODIES | GOLD | OXFORD |  |
| ANTI PURKINJE CELL Ab (anti YO) | GOLD | OXFORD |  |
| ANTI VOLTAGE GATED CHANNEL (potassium and calcium) | GOLD | OXFORD |  |
| APOLIPOPROTEIN A | GOLD | ROYAL FREE |  |
| APOLIPOPROTEIN B | GOLD | ROYAL FREE |  |
| APOLIPOPROTEIN C | GOLD or LAVENDER | GLASGOW |  |
| APOLIPOPROTEIN E -Genotyping | LAVENDER | EDINBURGH |  |
| APOC III | LAVENDER | GLASGOW |  |
| AQUA PORINE ANTIBODIES (NMO) | GOLD | OXFORD |  |
| ARSENIC | WHITE UNIVERSAL (urine) + LAVEDER (BLOOD) | BIRMINGHAM |  |
| ARIPIPRAZOLE | LAVENDER | KINGS |  |
| ASCORBIC ACID (VITAMIN C) | GREEN (protected from light) | ST THOMAS |  |
| ATENOLOL | GOLD | PENARTH |  |
| ATRIAL NATRIURETIC PEPTIDE | NOT ROUTINELY AVAILABLE | | |
| AUTOANTIBODY SCREEN | GOLD | NORTHAMPTON | 7 days |
| AZATHIOPRINE SENSITIVITY | LAVENDER | BIRMINGHAM |  |
| **B** |  |  |  |
| B-HCG (Molar pregnancy) | GOLD | CHARING CROSS |  |
| B2 MICROGLOBULIN-CSF |  |  | NOT ROUTINELY AVAILABLE SUGGEST SERUM B2M |
|  |  |  |  |
| BCR-ABL | 20mls LAVENDER | HAMMERSMITH & OXFORD |  |
| B2 GLYCOPROTEIN | GOLD | SHEFFIELD |  |
| BETA HYDROXYBUTYRATE | GREY FLU OX (paed yellow) | SHEFFIELD |  |
| BIOPTERINS | BLOOD SPOTS | BIRMINGHAM |  |
| BIOTINIDASE | GREEN | SHEFFIELD |  |
| BONE ALKALINE PHOSPHATASE | GOLD | LIVERPOOL |  |
| BONE MARKERS – URINE | WHITE TOP UNIVERSAL (URINE) | LIVERPOOL |  |
| BONE MARKERS – BLOOD |  |  | NOT ROUTINELY AVAILABLE |
| BRCA2 ONCOGENE | LAVENDER X 2 | CHURCHILL |  |
| **C** |  |  |  |
| C1 ESTERASE INHIBITOR | LAVENDER | OXFORD |  |
| C3 NEPHRITIC FACTOR | GOLD | SHEFFIELD | 20 mins |
| C-PEPTIDE | GOLD | OXFORD | 20 mins |
| CH 50 | GOLD | OXFORD | 20 mins |
| CH 100 | GOLD | OXFORD | 20 mins |
| C1Q | GOLD | SHEFFIELD | 20 mins |
| C.S.F. OLIGOCLONAL BANDS | CSF-WHITE (Uni)+Blood-GOLD (Gel) | SHEFFIELD |  |
| CALCITONIN | GOLD (on ice) | CHARING CROSS | 15 mins |
| CALCULI COMPOSITION | WHITE UNIVERSAL | UCL |  |
| CARBOHYDRATE DEFICIENT TRANSFERRIN | GOLD | KINGS |  |
| CARDIAC MUSCLE ANTIBODIES | GOLD | SHEFFIELD | 7 days |
| CARNITINES-PLASMA | GREEN | SHEFFIELD |  |
| CAROTENE (VITAMIN A) | DARK GREEN GEL FREE (protect from light) | ST THOMAS |  |
| CATECHOLAMINES – PLASMA |  |  | NOT ROUTINELY AVAILABLE SUGGEST PLASMA METANEPHRINES |
| CASPR2 ANTIBODIES | GOLD | OXFORD |  |
| CATHINONE (KHAT) |  |  | ASSAY WITHDRAWN |
| CD TRANSFERRIN - CSF | WHITE UNIVERSAL | SHEFFIELD |  |
| CD4/CD8 | LAVENDER | NORTHAMPTON | Monday – Thursday only. To arrive in pathology by 4pm. |
| CELL MARKER STUDIES | LAVENDER | OXFORD |  |
| CERULOPLASMIN | GOLD | CARDIFF |  |
| CIRCULATING EPIDERMAL Abs (Pemphigus and pemphigoid) | GOLD | NORTHAMPTON |  |
| CHLORAMPHENICOL | GOLD (Gel Bottle) | BRISTOL | Pre and Post wrapped in foil |
| CHOLINESTERASE | GOLD | MANCHESTER |  |
| CHROMIUM | White (universal)/ royal blue - blood | CARDIFF |  |
| CHROMOSOMES | GREEN (Lith Hep) | OXFORD |  |
| CHROMOGRANIN A | GOLD | SHEFFIELD |  |
| CHROMOGRANIN B | LAVENDER (on ice) | CHARING CROSS | 5 mins |
| CITRULLINATED PEPTIDE AB | GOLD | SHEFFIELD | 3 days |
| CLOBAZAM | GOLD | PENARTH |  |
| CLOMIPRAMINE |  |  | NOT ROUTINELY AVAILABLE |
| CLONAZEPAM | RED TOP GEL FREE | CHALFONT |  |
| CLOZAPINE/CLOZARIL | LAVENDER | PENARTH |  |
| COBALT | ROYAL BLUE | CARDIFF |  |
| COLISTIN | GOLD | BRISTOL | Pre + 1hr post dose |
| COMMON α SUBUNITS | GOLD | BIRMINGHAM |  |
| COPPER | GOLD | CARDIFF |  |
| CORTISONE | GOLD | SOUTHAMPTON |  |
| CORTISOL OUTPUT-URINE | 24HR URINE/ WHITE UNIVERSAL FOR PAEDS | LEEDS |  |
| COTININE | GOLD | MANCHESTER |  |
| CSF AMINO ACIDS | CSF + MATCHING GREEN TOP BLOOD | SHEFFIELD |  |
| CSF GLYCINE | CSF SAMPLE White universal | SHEFFIELD |  |
| CSF B2 MICROGLOBULIN |  |  | NOT AVAILABLE SUGGEST SERUM B2M |
| CTX | GOLD | NOTTINGHAM |  |
| CYCLOSPORIN | LAVENDER (EDTA) |  | Collection varies with each transplant centre |
| CYCLOSERINE | GOLD | BRISTOL | **PRE + POST DOSE** |
| CYSTIC FIBROSIS | LAVENDER | OXFORD |  |
| CYTOKINES |  |  | **NOT ROUTINELY AVAILABLE** |
| CYRFA 21-1 | GOLD |  |  |
| SHEFFIELD |
| **D** |  |  |  |
| 7 DEHYDROCHOLESTEROL | GOLD | ICH |  |
| 11 DEOXYCORTISOL | GOLD | ST THOMAS |  |
| DEHYDROEPIANDROSTERONE | GOLD | LEEDS |  |
| DIAZEPAM | LAVENDER | PENARTH |  |
| DIHYDRO TESTOSTERONE | GOLD or GREEN | LEEDS |  |
| DILTIAZEM | WHITE UNIVERSAL (URINE) |  |  |
| BIRMINGHAM |
| DNA STUDIES – MUSCULAR DYSTROPHY | LAVENDER | OXFORD |  |
| DNA STUDIES – FRAGILE X | LAVENDER | OXFORD |  |
| DNA STUDIES - PRADER WILLI & ANGLEMANS SYNDROME | LAVENDER | OXFORD |  |
| DNA STUDIES - MYOTONIC DYSTROPHY/MITOCHONDRIAL DISEASE/CYSTIC FIBROSIS/HUNTINDONS DISEASE | LAVENDER | OXFORD |  |
| DOWN'S RISK | GOLD (Gel Bottle) | OXFORD |  |
| DRUG SCREEN – BLOOD |  |  | **NOT ROUTINELY AVAILABLE SUGGEST URINE DRUG SCREEN** |
| DRUG SCREEN – URINE | WHITE UNIVERSAL (urine) | KINGS |  |
| dsDNA | GOLD | Northampton | 3 days |
| **E** |  |  |  |
| ENA | GOLD | NORTHAMPTON |  |
| ENDOMYSIAL ANTIBODY | GOLD | NORTHAMPTON | 7 days |
| ENGRAFTMENT STUDIES | LAVENDER | GOSH |  |
| ERYTHROPOIETIN | GOLD | KINGS |  |
| ETHOSUXAMIDE | LAVENDER | PENARTH |  |
| ETYLENE GLYCOL | GOLD or GREY or GREEN | BIRMINGHAM |  |
| **F** |  |  |  |
|  |  |  |  |
| FAECAL ELASTASE | BLUE UNIVERSAL WITH SPOON | CARDIFF | 10hrs |
| FAECAL FAT GLOBULES | BLUE UNIVERSAL WITH SPOON | OXFORD |  |
| FAMILIAL MEDITERRANEAN FEVER | LAVENDER | ROYAL FREE |  |
| FIBROBALST GROWTH FACTOR 23 | LAVENDER | NORWICH |  |
| FLECAINIDE | GOLD | PENARTH |  |
| FLIP1L1 - PDGFRα | 2X EDTA - MARROW | SALISBURY |  |
| FLUCYTOSINE | GOLD | BRISTOL | Pre and 1hr Post samples required |
| FLUOXETINE | LAVENDER | PENARTH |  |
| FRAGILE X | LAVENDER | OXFORD |  |
| FREE FATTY ACIDS | GREY/YELLOW (fluoride) | SHEFFIELD |  |
| FREE PHENYTOIN | GOLD | CHALFONT ST PETER |  |
| FREE SERUM LIGHT CHAINS | GOLD | OXFORD |  |
| FRUCTOSAMINE | GOLD | BIRMINGHAM |  |
| **G** |  |  |  |
| G6PD | LAVENDER (+ normal control) | OXFORD |  |
| GABAPENTIN | LAVENDER | PENARTH |  |
| α GALACTOSIDASE - (not Friday must be in ref lab within 24hrs) | GREEN | GOSH |  |
| Β GALACTOSIDASE - (not Friday must be in ref lab within 24hrs) | GREEN | GOSH |  |
| GAL-1-PUT - (not Friday must be in ref lab within 24hrs) | GREEN or ORANGE (Lith Hep) | INSTITUTE CHILD HEALTH (ICH) |  |
| GASTRIN (fasting sample) | LAVENDER on ice | CHARING CROSS | 5 mins |
| GENE PROBES | LAVENDER | OXFORD |  |
| GILBERTS DISEASE | LAVENDER | NINEWELLS |  |
| GLOMERULAR BASEMENT MEMBRANE | GOLD | NORTHAMPTON |  |
| GUT HORMONES | LAVENDER (on ice) | CHARING CROSS | 5 mins |
| GLUTATHIONE PEROXIDASE | ORANGE or LAVENDER | GLASGOW |  |
| GLYCOSAMINOGLYCANS | WHITE UNIVERSAL (urine) | SHEFFIELD |  |
| GLYCINE – BLOOD OR CSF | GREEN or WHITE UNIVERSAL | SHEFFIELD |  |
| Β2 GLYCOPROTEIN | GOLD | SHEFFIELD |  |
| GOLD (BLOOD OR URINE) | GOLD or WHITE UNIVERSAL | GUILFORD |  |
| **H** |  |  |  |
| 5HIAA | 24hr URINE (ACID) dietary restrictions | OXFORD |  |
| HAEMOCHROMATOSIS GENE TEST | LAVENDER | OXFORD |  |
| HAPTOGLOBIN | GOLD | SHEFFIELD |  |
|  |  |  |  |
| HLA TISSUE TYPING | LAVENDER | OXFORD |  |
| HOMOCYSTEINE - FASTING | LAVENDER | SHEFFIELD | 1 hour |
|  |  |  |  |
| HUNTINGDONS CHOREA | LAVENDER | OXFORD |  |
| HOMOVANILLIC ACID | 24 HR URINE (ACID) | SHEFFIELD |  |
| HEXOSAMINIDASE LEVELS (not Friday must be in ref lab within 24hrs) | DARK GREEN | GUYS |  |
| **I, J, K** |  |  |  |
| IGD | GOLD | SHEFFIELD |  |
| IGFBP3 | GOLD | ROYAL SURREY |  |
| 68KD INNER EAR PROTEIN | RED TOP | CAMBRIDGE |  |
| IMMUNOGLOBULIN SUBCLASSES | GOLD | SHEFFIELD |  |
| IMMUNOPHENOTYPING | LAVENDER X2 BONE MARROW | OXFORD | Consultant request only. |
| IMMUNOREACTIVE TRYPSIN |  |  | NO LONGER AVAILABLE |
| IMMUNOSELECTION ELECTROPHORESIS | GOLD or WHITE UNIVERSAL (urine) | SHEFFIELD |  |
| TOTAL INHIBIN | GOLD | SHEFFIELD |  |
| INSULIN (Not practical from H.C.) | GOLD | CHARING CROSS | 80 mins |
| INSULIN ANTIBODIES | GOLD | SHEFFIELD |  |
| INSULIN-LIKE GROWTH FACTOR 1 | GOLD | OXFORD |  |
| INTERLEUKIN 6 |  |  | NOT ROUTINELY AVAILABLE |
| ITRACONAZOLE | GOLD | BRISTOL |  |
| JAK - 2 | LAVENDER | OXFORD |  |
| KALETRA |  |  | **NOLONGER AVAILABLE** |
| **L** |  |  |  |
| LAMOTRIGINE (LAMICTAL) | LAVENDER | PENARTH |  |
| LEAD | LAVENDER | CARDIFF |  |
| LDH ISOENZYMES | GOLD OR GREEN | GOSH |  |
| LEBER’S HEREDITARY OPTIC NEUROPATHY | LAVENDER | OXFORD |  |
| LEPTIN ASSAY | GOLD or GREEN | ADDENBROOKS |  |
| LEVETIRACETAM | 2 ml EDTA | PENARTH |  |
| LGI1 ANTIBODIES | GOLD | OXFORD |  |
| LONG CHAIN FATTY ACIDS | GOLD or GREEN or LAVENDER | SHEFFIELD |  |
| LYMPHOCYTE CELL MARKERS | 2X LAVENDER | OXFORD |  |
| LYSOSOMAL ENZYMES | GREEN or ORANGE | GOSH |  |
| **M** |  |  |  |
| MACROAMYLASE | GOLD | KINGS |  |
| MAGNESIUM | 24HR URINE (PLAIN) | GUILFORD |  |
| MANGANESE | EDTA OR DARK BLUE | CARDIFF |  |
| MAST CELL TRYPTASE (mast cell syndrome) | GOLD or LAVENDER | SHEFFIELD |  |
| MCAD DEFICIENCY | GUTHRIE CARD (neonates) WHITE UNIVERSAL URINE (older children) | SHEFFIELD |  |
| MERCURY | LAVENDER+WHITE UNIVERSAL (urine) | CARDIFF |  |
| METALLOPROTEASE | BLUE (CITRATE) | OXFORD |  |
| METANEPHRINES - PLASMA | LAVENDER ON ICE | MANCHESTER | Strict protocol contact 85768 for Info |
| METANEPHRINES - URINE | 24hr URINE - PLAIN | OXFORD |  |
| METHYL MALONIC ACID | WHITE UNIVERSAL (urine) | SHEFFIELD |  |
| NEUTROPHIL FUNCTION TEST | LAVENDER | OXFORD | 1hr – prior arrangement with pathology required before sample taken. |
| MIDAZOLAM | GOLD | PENARTH |  |
| MIRTAZEPINE | EDTA | PENARTH |  |
| MYCOPHENOLATE MOSETIL | LAVENDER | KINGS |  |
| MUCOPOLYSACCHARIDES | WHITE UNIVERSAL (urine) | SHEFFIELD |  |
| MYOCARDIAL ANTIBODIES | GOLD | SHEFFIELD |  |
|  |  |  |  |
| **N** |  |  |  |
| N-ACETYLGLUCOSAMINIDASE | WHITE UNIVERSAL (urine) | GOSH |  |
| NEORAL LEVELS | LAVENDER | OXFORD |  |
| NEURODEGENERATIVE ENZYME SCREEN | GREEN | GOSH | 2hrs. Monday – Thursday only. |
| NEUROMYELITIS OPTICA Ab | GOLD | OXFORD |  |
| NEURO SPECIFIC ENOLASE | GOLD | SHEFFIELD |  |
| NEUTROPHIL FUNCTION TEST | LAVENDER | OXFORD | 2hrs. Monday - Thursday am only |
| NITRAZEPAM | LAVENDER – FOIL WRAPPED | PENARTH |  |
| NMDA | GOLD | OXFORD |  |
| NORADRENALINE | GREEN | BARTS | Strict protocol contact 85768 for Info |
| NTX | WHITE UNIVERSAL | SHEFFIELD | 2nd morning void |
| **O** |  |  |  |
| OLANZAPINE | LAVENDER | PENARTH |  |
| OLIGOCLONAL BANDS | CSF AND GOLD (blood sample) | SHEFFIELD |  |
| OLIGOSACCHARIDES | WHITE UNIVERSAL (urine) | LEEDS |  |
| ORGANIC ACIDS | WHITE UNIVERSAL (urine) | SHEFFIELD |  |
| OROSOMUCOID (α-1 acid glycoprotein) | GOLD | SHEFFIELD |  |
| OROTIC ACID QUANTITATION | WHITE UNIVERSAL (urine) | SHEFFIELD |  |
| OSTEOCALCIN | LAVENDER | NORWICH |  |
| OVARIAN ANTIBODIES | GOLD | CARDIFF |  |
| OXALATE OUTPUT-URINE | 24HR URINE | UCL |  |
| **P** |  |  |  |
| P1NP | GOLD | LIVERPOOL |  |
|  |  |  |  |
| P3NP | RED | LIVERPOOL |  |
| P50 | LAVENDER | BIRMINGHAM | Before 12 noon Mon-Thur only |
| PARANEOPLASTIC ANTIBODIES (anti HO, YU, RI) | GOLD | OXFORD |  |
| PERIODIC FEVER SYNDROME | LAVENDER | ROYAL FREE |  |
| PETHIDINE | WHITE UNIVERSAL (urine) | BIRMINGHAM |  |
| PHENCYCLIDINE (Angle Dust) (urine drug screen) | WHITE UNIVERSAL (urine) | BIRMINGHAM |  |
| PHENYLALANINE | ORANGE or BLOOD SPOT | SHEFFIELD |  |
| PHOSPHOETHANOLAMINE | WHITE UNIVERSAL (urine) | GOSH |  |
| PHYTAMIC ACID | GREEN | SHEFFIELD |  |
| PIPECOLIC ACID | GREEN OR LAVENDER (BLOOD) | SHEFFIELD |  |
|  | WHITE UNIVERSAL (CSF) | SHEFFIELD |  |
|  | WHITE UNIVERSAL (URINE) | SHEFFIELD |  |
| PITUITARY POLYPEPTIDES | GOLD | BIRMINGHAM |  |
| PLACENTAL ALK PHOS | GOLD | CHARING CROSS |  |
| PLASMA METANEPHRINES | LAVENDER ON ICE | MANCHESTER | strict protocol contact 85768 for info |
|
|
|
| PLASMA PHYTOSTEROLS | GREEN | INSTITUTE OF CHILD HEALTH |  |
| PNH STUDIES | LAVENDER | OXFORD | Mon - Thur am only |
| PORPHYRINS – BLOOD | LAVENDER or GREEN (in foil) | KINGS |  |
| PORPHYRINS - FAECES | BLUE UNIVERSAL (in foil) | KINGS |  |
| PORPHYRINS-URINE | 24 HR URINE (in foil) | KINGS |  |
|
| PSA (FREE) | GOLD | CHARING CROSS | Arrive within 1 hr |
| PROCOLLAGEN TYPE III | RED TOP – GEL FREE | SHEFFIELD |  |
| PROINSULIN | GOLD or GREEN | GUILDFORD | 15 mins |
| PSEUDOCHOLINESTERASE | GOLD | MANCHESTER |  |
| PURINE STUDIES | 24HR URINE + LAVENDER | ST THOMAS |  |
| PURKINJE CELL ANTIBODIES (anti YO) | GOLD | OXFORD |  |
|  |  |  |  |
| PYRUVATE | SPECIAL BOTTLE CONTACT PATHOLOGY | SHEFFIELD | Only with prior arrangement with Sheffield Childrens Hospital and Milton Keynes Pathology. |
| PYRUVATE KINASE | LAVENDER + NORMAL CONTROL | HAMMERSMITH |  |
|
|
| **Q** |  |  |  |
| QUETIAPINE | EDTA | PENARTH |  |
| QUININE | EDTA | KINGS |  |
| **R** |  |  |  |
| RAST (specify individual rast required) | GOLD | SHEFFIELD |  |
| RED CELL ENZYMES | GREEN | GOSH | 2hr to pathology Mon – Thur only |
| RENIN | GREEN | CHARING CROSS | 30 min |
| RETINOL BINDING PROTEIN | WHITE UNIVERSAL (freeze immediately) | GOSH | 5 mins |
| RISPERIDONE (BLOOD) | LAVENDER | PENARTH |  |
| RISPERIDONE (URINE) | WHITE UNIVERSAL (urine) | ST THOMAS |  |
| **S** |  |  |  |
| SABRIL (VAGABATRIN) |  |  | NO LONGER AVAILABLE |
| SELENIUM | GOLD or DARK BLUE | CARDIFF |  |
|  |  |  |  |
| SIALIC ACID | WHITE UNIVERSAL (urine) | MANCHESTER |  |
| SOFT TISSUE TRANSGLUTAMINASE | GOLD | SHEFFIELD |  |
| SOTALOL | LAVENDER | PENARTH |  |
| STEROID PROFILE-URINE | 24 HR URINE or UNIVERSAL | UCL |  |
| STONES | STONE | UCL |  |
| STRIATED MUSCLE ANTIBODIES | GOLD | OXFORD |  |
| SULPHITE | WHITE UNIVERSAL (urine) | GOSH |  |
| SULPHONYL UREA | GOLD | GUILFORD |  |
| **T** |  |  |  |
| TAY-SACHS (CARRIER TESTING) | PREGNANT – 2 GREEN + 1 LAVENDER | ST THOMAS |  |
|  | NON PREGNANT OR MALE – 2 GREEN + 1 GOLD | ST THOMAS |  |
| TACROLIMUS | LAVENDER | KINGS |  |
| T-CELL REARRANGEMENT STUDIES | LAVENDER | SURREY |  |
| T-CELL SUBSETS | LAVENDER | NORTHAMPTON | Monday – Thursday only. To arrive in pathology by 4pm |
| THALLIUM | WHITE UNIVERSAL (urine) + DARK BLUE | CARDIFF |  |
| THIOGUANINE NUCLEOTIDE | LAVENDER | ST THOMAS' |  |
| THIOSULPHATE | WHITE UNIVERSAL (urine) | SHEFFIELD |  |
| THYROGLOBULIN | GOLD |  |  |
| SHEFFIELD |
| THYROGLOBULIN ANTIBODIES | GOLD | SHEFFIELD |  |
| THYROID HORMONE BINDING PROTEIN | GOLD | ADDENBROOKES |  |
| THYROID HORMONE RESISTANCE SYNDROME | LAVENDER | ADDENBROOKES |  |
| THYROTROPIN RECEPTOR Abs | GOLD | SHEFFIELD |  |
| TIAGABINE LEVEL | EDTA | PENARTH |  |
| TITANIUM | ROYAL BLUE | CHARING CROSS |  |
| TPMT | LAVENDER | ST THOMAS |  |
| TRANSFERRIN GLYCOFORMS (aka isoforms + iso electric) | GOLD | QUEENS SQUARE |  |
|  |  |  |  |
| TRAXODONE | RED TOP | PENARTH |  |
| TRICYCLIC ANTIDEPRESSANTS | RED TOP | BIRMINGHAM |  |
| TRYPTASE – ALLERGIC REACTION | GOLD or LAVENDER or GREEN (<1hr, 3hr,24hr post event) | SHEFFIELD |  |
| TRYPTASE – NON URGENT REACTION | GOLD or LAVENDER | SHEFFIELD |  |
| TSH RECEPTOR ANTIBODIES | GOLD | SHEFFIELD |  |
| tTGs | GOLD | NORTHAMPTON | 7 days |
| **U** |  |  |  |
| URATE/CREATININE RATIO | WHITE UNIVERSAL (urine) | SHEFFIELD |  |
| URINE AMINO ACIDS | WHITE UNIVERSAL (urine) | SHEFFIELD |  |
| URINARY CITRATE | WHITE UNIVERSAL (urine) | UCH |  |
| URINARY COPPER | 24hr URINE (PLAIN) | CARDIFF |  |
| URINARY CYSTEINE/CITRATE | WHITE UNIVERSAL (urine) | UCH |  |
| URINARY IRON | 24hr URINE (PLAIN) | SOUTHAMPTON |  |
|  |  |  |  |
| **V** |  |  |  |
| 1,25 OH VITAMIN D | GOLD | GLASGOW |  |
| VIGABATRIN | LAVENDER | PENARTH |  |
| VIP | LAVENDER (on ice) | CHARING CROSS | 5 mins |
| VISCOSITY – PLASMA | LAVENDER | OXFORD |  |
| VITAMIN A | DARK GREEN GEL FREE | ST THOMAS |  |
| VITAMIN B1 | LAVENDER (in foil) | ST THOMAS |  |
| VITAMIN B2 | LAVENDER (in foil) | ST THOMAS |  |
| VITAMIN B6 | LAVENDER (in foil) | ST THOMAS |  |
| VITAMIN C | GREEN (in foil) | ST THOMAS |  |
| VITAMIN E | DARK GREENGEL FREE (in foil) | ST THOMAS |  |
| VITAMIN K | GOLD (in foil) | ST THOMAS |  |
| VORICONAZOLE | GOLD | BRISTOL |  |
| **W, X, Y, Z** |  |  |  |
| WARFARIN | GOLD OR LAVENDER | ST THOMAS |  |
| WHITE CELL ENZYMES | GREEN | ICH | 2hrs to pathology |
| Y DELETION | LAVENDER | BRISTOL |  |
| ZARONTIN (ETHOSUXIMIDE) | GOLD | PENARTH |  |
| ZINC | ROYAL BLUE (Trace metal bottle) | CARDIFF | 2hrs |
| ZINC PROTOPORPHYRIN | LAVENDER | CARDIFF |  |

**MILTON KEYNES UNIVERSITY HOSPITAL SITE MAP:**



