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	Revision Number:	25
	Issue Date:	30/05/2023
	Next Review:	05/2026
National Orthopaedic Hospital Cappagh, Finglas, Dublin 11.		PATHOLOGY LABORATORY USER MANUAL


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INTRODUCTION

The Laboratory in National Orthopaedic Hospital Cappagh (NOHC) consists of 4 sections: Biochemistry, Blood Transfusion, Haematology and Microbiology/Molecular. NOHC ensures patient confidentiality is maintained at all times.

The laboratory is accredited to ISO 15189:2012 standards (Registration Number 318MT). Refer to <https://www.inab.ie/FileUpload/Medical-Testing/National-Orthopaedic-Hospital-Cappagh-318MT.pdf> for a list of accredited tests. Under the terms of the EU Tissue Directive, NOHC is a designated Tissue Establishment and requires licensing by the Health Products Regulatory Authority.

POSITION	PHONE
Laboratory Manager	(01) 814 0386
Quality Systems Manager	(01) 814 0456
Chief /Senior Medical Scientist (Haematology/Blood Transfusion)	(01) 814 0413 (01) 814 0377
Chief/Senior Medical Scientist (Biochemistry)	(01) 814 0472
Chief/Senior Medical Scientists (Microbiology)	(01) 814 0305
Haemovigilance Officer	(01) 814 0330
Senior Phlebotomist	(01) 814 0365

Location:

The Laboratory is located beside the Professorial Unit opposite the Out- Patient's entrance.


Address:

Pathology Department,
National Orthopaedic Hospital Cappagh,
Finglas, Dublin 11
D11 EV29

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Hours of service:

Monday – Friday 8.00am to -8.00pm, 6.00 pm to 8.00 pm for urgent specimens including COVID- 19

Monday – Friday 7.00am to 8.00am the on call medical scientist is on site to process pre-op bloods for Same Day Admission (SDA) Patients

There is a limited emergency on call service available for Biochemistry, Blood Transfusion, Haematology and Microbiology outside of these hours.

Sample requirements:


All sample tubes must be filled to the marker on the bottle. The samples are placed into the plastic biohazard bag attached to a blue NOHC request form specifically for Biochemistry, Haematology and Microbiology/Molecular (RF-LAB-50) or a red NOHC request form specifically for Blood Transfusion (RF-BT-78).

SAMPLES		
DISCIPLINE	ESSENTIAL	DESIREABLE
Biochemistry and Haematology	Patient's Full Name	Patient's consultant
	Date of Birth	Patient location/destination for report
	Hospital Number (MRN)	Signature of Venepuncturist
		Date &, where relevant, time of primary Specimen Collection
Blood Transfusion	Refer to Biochemistry and Haematology essential requirements. Handwritten or BloodTrack PDA labels accepted ONLY. There is zero tolerance on any discrepancies in patient demographics. Additionally: Signature of venepuncturist or	Patient's consultant
		Patient location/destination for report
		Clinical Details including previous Transfusion history and/or pregnancies.

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	BloodTrack PDA label. Date & Time of Specimen Collection. Special Requirements i.e. CMVNEG/IRR	
Microbiology and Histology	Refer to Biochemistry and Haematology essential requirements	Refer to Biochemistry and Haematology desirable requirements
	Site of Sample	
	<u>Specimen Description</u> If more than 1 Microbiology sample is taken then label the individual samples 1. 2, 3 etc. (with full details on each). If more than 1 Histology sample is taken then label the individual samples A. B, C etc. (with full details on each).	

Request Form Requirements:


Addressographs are accepted on all NOHC request forms. Please affix addressograph labels to ALL copies. The request form should contain the following information

REQUEST FORM		
DISCIPLINE	ESSENTIAL	DESIRABLE
Biochemistry and Haematology Pathology Laboratory Request Form (RF-LAB-50) Rheumatology Request Form (RF-LAB-63)	Patient's Full Name	Clinically relevant information
	Date of Birth	Patient's address
	Hospital Number (MRN)	
	Gender	

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	Patient's consultant	
	Patient location/destination for report	
	Signature of Venepuncturist or BloodTrack PDA label.	
	Date &, where relevant, time of primary Specimen Collection	
	Required tests*	
Microbiology (RF-LAB-50) Histology (RF-LAB-54) Request Forms	Refer to Biochemistry and Haematology essential requirements	Relevant clinical details and antimicrobial therapy
	Specimen Type	
	Specimen Site	
	Specimen Description If more than 1 Microbiology sample is taken then label the individual samples 1. 2, 3 etc. (with full details on each). If more than 1 Histology sample is taken then label the individual samples A. B, C etc. (with full details on each).	
Blood Transfusion: Blood Transfusion Request for Blood Grouping and Crossmatching (RF-BT-78)	Refer to Biochemistry and Haematology essential requirements. There is <u>zero tolerance</u> on any discrepancies in patient demographics.	Refer to Biochemistry and Haematology desirable requirements
	Requesting Doctor *	
	CMV/IRR requirements	

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** If additional requests are required for Biochemistry, Haematology or Microbiology/Molecular, the requesting clinician MUST complete the Pathology Laboratory Request Form (RF-LAB-50) and forward to the laboratory. Please note that the laboratory will accept an emailed copy of this completed request form to labrequests@nohc.ie*

*** If Blood/blood products are required, the requesting clinician MUST complete the Blood Transfusion Request form (RF-BT-78) and forward to the Blood Transfusion laboratory. Blood/blood products will be prepared but not issued, until receipt of this form. Please note that the blood transfusion laboratory will accept an emailed copy of this completed request form to labrequests@nohc.ie*

Requirements for samples referred to the Mater Misericordiae University Hospital (MMUH), St James Hospital, IBTS and NVRL are listed from **page 52.**

Specimen Acceptance/Rejection Criteria

It is the policy of the Pathology Laboratory within the NOHC to only accept hospital specimens if both the specimen and the request form match and have the following minimum three key identifiers:

- *Patient's name*
- *Patients' Date of Birth*
- *Patient's Hospital number*


While every effort is made to minimise rejection of samples, some samples will be deemed unsuitable for analysis or will yield inaccurate results. These include:

- Grossly haemolysed samples
- Clotted Samples
- Underfilled samples
- Contaminated Samples due to incorrect order of draw. If an EDTA sample is taken before the biochemistry lithium heparin the potassium and calcium measurement can be affected.

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- Samples taken from an arm with an infusion or a line may yield falsely elevated or decreased results.
- Sample/Analyte that may have deteriorated due to prolong transit time e.g. APTT >4 HRS.
- Leaking Specimens
- Aged samples

Samples received into the laboratory, which are rejected for analysis due to sample quality, etc must be recorded in the LIS and then credited with the appropriate comment and requestor notified.

In the case of samples that cannot be easily repeated, the sample and request form are amended by the originator in the laboratory. A laboratory Disclaimer (AP-LAB-5) is completed and these samples are processed as per normal. The nature of the disclaimer is indicated on the laboratory report.

Additional/Spare Samples

Any additional/spare samples received with no apparent test request, are also given a unique laboratory number. The requester, if known, is informed. The sample type will be indicated on the final laboratory report. Samples are stored for up to 7 days if viable. This is not related to spare Blood Transfusion samples.

Within Hours Sample Storage

All samples are stored at Ward level in the specimen Transport Box provided by the Laboratory.
All SSC samples are prepared by SSC staff for transport to NOHC in line with ADR Regulations


Out of Hours Sample Storage

All NOHC out of hours microbiology/molecular samples (with the exception of blood cultures and bone harvest samples) are to be stored at 4°C in the out of hours laboratory specimen fridge (which is located in the old recovery room) for up to 48h. Urine samples that are retained at room temperature for > 2 hours will not be processed.

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NOHC Blood cultures must immediately go to the NOHC Microbiology Laboratory and be placed onto the Bact/ALERT Blood Culture analyser. **DO NOT REFRIGERATE.**

Bone Bank samples must immediately go to the NOHC Microbiology Laboratory and be placed into the designated incubators. If this is not possible, store at room temperature but **DO NOT REFRIGERATE.**

For any out of hours NOHC Biochemistry, Haematology and Blood Transfusion samples contact the **on call medical scientist ASAP as per table below. DO NOT REFRIGERATE.**

Contacting the On Call Medical Scientist:

On Call Medical Scientists for Biochemistry, Blood Transfusion and Haematology can be contacted through the Laboratory mobile phones on:

Laboratory On call	Telephone Numbers
Weekday (8pm-7am)	Mobile: 087 2301505
Weekend (8am-8am)	Mobile: 087 6607955
Bank Holidays (8am-8am)	Mobile: 087 2301505

In the event that the rostered medical scientist cannot be contacted **please contact NOHC switchboard on (01) 834 1211.**

Specimen collection times:


Porters collect and deliver NOHC samples to the lab at intervals throughout the day during routine working hours. Phlebotomists will also deliver NOHC samples if not collected by the porter.

Porters will collect and deliver NOHC samples to the lab on Saturday morning. The on call Medical Scientist will collect samples as required, outside this time.

SSC will have a twice daily specimen collection from NOHC Ambulance service. SSC will arrange **Authorised by the Quality Systems Manager – see Q-Pulse Record for details**

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transport of SSC specimens outside the set collections. All request forms are date stamped at specimen reception.

Routine samples must arrive at the lab before **5pm** to be analysed on the same day. Please see Microbiology Department Section for exceptions to these cut off times. Samples to be referred to other hospitals must arrive in the lab before **2:30pm** to be sent on the same day.

Report delivery times:

Reports are placed into sealed envelopes (to maintain confidentiality) and placed into specific pigeonholes in the laboratory and collected by porters throughout the day. The reports are delivered to their source location as indicated on the request form.

SSC reports will be delivered twice daily by NOHC Ambulance service, Monday to Friday. SSC reports will be delivered throughout the weekend/bank holiday by taxi.

The reports are also available on ward enquiry.

Turnaround times are calculated from the time the sample is RECEIVED by the laboratory until the report is available on Ward lookup

Additional Examinations:

In compliance with PP-NOHC-22 (Effective Clinical Communication), verbal requests for additional examinations (add-on requests for tests) for all disciplines will require a completed request form for the relevant department to be sent via email (labrequests@nohc.ie) or hard copy to the laboratory. Viability of additional tests will be confirmed prior to testing by the Laboratory staff.


Repeat examination:

Repeat examination due to analytical failure, further examination of primary specimen or delayed processing is recorded on the LIS by way of a report comment. Discarded specimens and additional examinations are captured on the LIS.

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Specimen Transport: *P650 Packaging Instructions (Extract from ADR 2017) AP-LAB-2*

P650	PACKAGING INSTRUCTION	P650
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This packing instruction applies to UN 3373 (Diagnostic Specimens).

General Requirements

(1) The packaging shall be of good quality, strong enough to withstand the shocks and loadings normally encountered during carriage, including transshipment between vehicles or containers and between vehicles or containers and warehouses as well as any removal from a pallet or overpack for subsequent manual or mechanical handling. Packaging shall be constructed and closed to prevent any loss of contents that might be caused under normal conditions of carriage by vibration or by changes in temperature, humidity or pressure.

(2) The packaging shall consist of at least three components:


- i) a leak-proof primary receptacle(s)
- ii) a leak-proof secondary packaging
- iii) an outer packaging of adequate strength for its capacity, mass and intended use, and with at least one surface
 - having minimum dimensions of 100 mm × 100 mm;
 - Either the secondary or outer packaging should be rigid.

(3) Primary receptacles shall be packed in secondary packaging in such a way that, under normal conditions of carriage, they cannot break, be punctured or leak their contents into the secondary packaging. Secondary packagings shall be secured in outer packagings with suitable cushioning material. Any leakage of the contents shall not compromise the integrity of the cushioning material or of the outer packaging. Enough absorbent material is placed

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between layers so that in the case of leakage there is enough absorbent material to absorb all fluid.

- (4) For carriage, the mark illustrated below shall be displayed on the external surface of the outer packaging on a background of a contrasting colour and shall be clearly visible and legible. The mark shall be in the form of a square set at an angle of 45° (diamond shaped) with minimum dimensions of 50 mm by 50 mm; the width of the line shall be at least 2 mm and the letters and numbers shall be at least 6 mm high. The proper shipping name "Biological Substance, Category B" in letters at least 6 mm high shall be marked on the outer package adjacent to the diamond-shaped mark.



- (5) At least one surface of the outer packaging shall have a minimum dimension of 100mm x 100mm.
- (6) The completed package shall be capable of successfully passing the drop test in 6.3.5.3 as specified in 6.3.5.2 at a height of 1.2 m. Following the appropriate drop sequence, there shall be no leakage from the primary receptacle(s) which shall remain protected by absorbent material, when required, in the secondary packaging.


(7) For liquid substances:

- (a) The primary receptacle(s) shall be leakproof;

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- (b) The secondary packaging shall be leakproof;
- (c) If multiple fragile primary receptacles are placed in a single secondary packaging, they shall be either individually wrapped or separated to prevent contact between them;
- (d) Absorbent material shall be placed between the primary receptacle(s) and the secondary packaging. The absorbent material shall be in quantity sufficient to absorb the entire contents of the primary receptacle(s) so that any release of the liquid substance will not compromise the integrity of the cushioning material or of the outer packaging;
- (e) The primary receptacle or the secondary packaging shall be capable of withstanding, without leakage, an internal pressure of 95 kPa (0.95 bar).

(8) For solid substances:

- (a) The primary receptacle(s) shall be sift proof;
- (b) The secondary packaging shall be sift proof;
- (c) If multiple fragile primary receptacles are placed in a single secondary packaging, they shall be either individually wrapped or separated to prevent contact between them.
- (d) If there is any doubt as to whether or not residual liquids may be present in the primary receptacle during carriage, then a packaging suitable for liquids, including absorbent materials, shall be used.


(9) Refrigerated or frozen specimens: Ice:

- (a) Ice shall be placed outside the secondary packagings or in the outer packaging or an overpack. Interior supports shall be provided to secure the secondary packagings in the original position. If ice is used, the outside packaging or overpack shall be leakproof.
- (b) The primary receptacle and the secondary packaging shall maintain their integrity at the temperature of the refrigerant used as well as the temperatures and the pressures which could result if refrigeration were lost.

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- (10) When packages are placed in an overpack, the package markings required by this packing instruction shall be clearly visible or reproduced on the outside of the overpack.
- (11) Infectious substances assigned to UN No. 3373 which are packed and packages which are marked in accordance with this packing instruction are not subject to any other requirement in ADR.
- (12) Clear instructions on filling and closing such packages shall be provided by packaging manufacturers and subsequent distributors to the consignor or to the person who prepares the package (e.g. patient) to enable the package to be correctly prepared for carriage.
- (13) Other diagnostic goods shall not be packed in the same packaging as Class 6.2 infectious substances unless they are necessary for maintaining the viability, stabilizing or preventing degradation or neutralization the hazards of the infectious substances. A quantity of 30 mls or less of dangerous goods included in Classes 3,8 or 9 may be packed in each primary receptacle containing infectious substances. When these small quantities of dangerous goods are packed with infectious substances in accordance with this packing instruction no other requirements of ADR need be met.
- (14) If any substance has leaked and has been spilled in a vehicle or container, it may not be reused until after it has been thoroughly cleaned and, if necessary, disinfected or decontaminated. Any other goods and articles carried in the same vehicle or container shall be examined for possible contamination.

For packaging of limbs see "Procedure for Specimen and Amputated Limb Collection and Processing" (PP-THE-1).

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PHLEBOTOMY SERVICE

Phlebotomy provides a service for NOHC In-Patients, Out-Patients and staff members (GP referral letters, hospital referrals, occupational referrals and inoculation screening) SSC will maintain an onsite Phlebotomy service.

Location:

NOHC Phlebotomy is located on the ground floor opposite the Day Ward. Bloods are taken outside of the area on the wards or in the Out-Patient Department.

Hours of service:

Monday – Thursday 07.00am – 15.00pm (Last Call 14.45pm)
Friday 07.00am – 14.15pm (Last Call 14.00pm)
Saturday 08.00am – 10.00am


Staff:

The phlebotomists can be reached on bleep numbers: 8867, 8868 and 8869.








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Order of Draw: (AP-PHL-3)





COLOUR CODE	ORDER	INVESTIGATION
 1 WHITE	SERUM 4.9ml 1 STAND UPRIGHT ONCE DRAWN	Serology, Testosterone, Viral Screens, Vitamin D (always do Bone Profile with Vitamin D). IGF-1 Allergen Screening (4.9 ml x2) (Specify each allergen)
 2 GREEN	Sodium Citrate (COAG) 2.7/10ml	PT/INR, APTT, Fibrinogen and D-Dimers Factor V Leiden (2.7ml x 2 plus 2.7ml x 1 EDTA grey top) Thrombophilia Screen (2.7ml x 6 plus 2.7ml x 1 EDTA grey top) Von Willebrand's Screen (2.7ml x 4) UNDER FILLED SAMPLES WILL BE REJECTED
 3 BROWN	SERUM GEL 4.9ml STAND UPRIGHT ONCE DRAWN	LDH, ACE, Therapeutic Drug levels, Lithium (4.9 ml x1) for Biochemistry only All Immunology : Electrophoresis, RH Factors, CCP, Ds DNA, C3, C4, ANA, ENA, ANCA, Coeliac/TTG (4.9 ml x1)
 4 ORANGE	METAL FREE LITHIUM HEPARIN 7.5 ml	Metal Ions including ASR's – Chromium/Cobalt Use special Metal Free Lithium Heparin tube and special metal free needle Extra metal free tube required for Titanium
 5 ORANGE	LITHIUM HEPARIN GEL 4.9ml	Tests done by NOHC require (4.9ml x1) Renal, Liver, Bone Profiles (Calcium & Phosphate), Magnesium, CRP, Vancomycin, Haematinics (Iron, Transferrin, B12, Folate, Ferritin, TSAT) Troponin I (Extra Sample bottle to be Taken for Troponin I) Tests referred to MMUH Lab require separate (4.9ml x1) Lipid Profile, Urate (Uric Acid), CK (CPK), Amylase, PSA, TFT, TPO Ab, FSH/LH, Progesterone, Prolactin, Oestradiol, Gentamicin, Osmolality, Cortisol. Homocysteine (contact Lab) Tumour markers (CEA, CA 125, CA 19.9)
 6 RED	BLOOD TRANSFUSION 7.5ml	Group, Crossmatch HLA B27 (full tube x1)
 7 RED Wide 2.7ml tube only	EDTA 2.7ml	HBA1c (separate 2.7ml tube) PTH (Always do Bone Profile with PTH), NT-Pro BNP (Contact lab prior to collection 4-hour window)

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	8 GREY	EDTA 2.7ml	FBC, IM Screen, Sickledex/Hb electrophoresis, Haemochromatosis, Lymphocyte Subsets.
	9 DARK PINK	ThromboExact 2.7ml	Blood sample tubes kept in the laboratory
	10 YELLOW	SODIUM FLUORIDE 2.7ml	Glucose, Glucose Tolerance Test (GTT)
	11 PURPLE		ESR UNDER FILLED SAMPLES WILL BE REJECTED

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Recommendations for blood culture collection

a summary of good practice

Using winged blood collection set

1> Prepare blood collection kit

Gather all materials before beginning the procedure. Ensure the blood culture bottles are within date. Do not use bottles which show any signs of damage, deterioration or contamination.



2> Prepare bottles for inoculation

Wash hands with soap and water then dry, or apply an alcohol hand rub. Remove the plastic "flip-cap" from the blood culture bottles and disinfect the septum using an appropriate disinfectant, such as 2% chlorhexidine in 70% isopropyl alcohol, 70% isopropyl alcohol, or iodine in swab or applicator form. Use a fresh swab/applicator for each bottle. **Allow bottle tops to dry in order to fully disinfect.**



3> Prepare venipuncture site

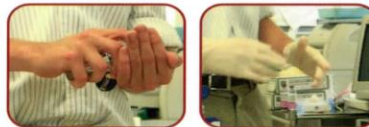
Confirm the patient's identity. If skin is visibly soiled, clean with soap and water. Apply a disposable tourniquet. Palpate to identify the vein and cleanse using an appropriate disinfectant, such as 2% chlorhexidine in 70% isopropyl alcohol, 70% isopropyl alcohol, or iodine in swab or applicator form.

The venipuncture site is not fully clean until the disinfectant has fully evaporated.



4> Wash hands. Wear gloves.

Wash hands again or use an alcohol hand-rub and apply clean examination gloves. Sterile gloves are not necessary.



5> Venipuncture

Attach a winged blood collection set to a collection adapter cap. **To prevent contaminating the puncture site, do not re-palpate the prepared vein before inserting the needle.** Insert the needle into the prepared site.



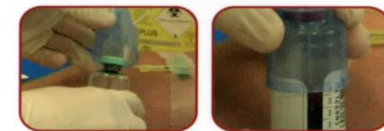
6> Culture bottle inoculation

Place the adapter cap over the aerobic bottle and press down to pierce the septum. Hold the bottle below the patient's arm with the bottle in an upright position, and use the graduation lines to accurately gauge sample volume.

Do not allow the culture bottle contents to touch the stopper or the end of the needle during the collection. Monitor the process closely to ensure proper flow is obtained and that there is no backflow of bottle contents in the adapter tubing.

Add up to 10 ml of blood per adult bottle and up to 4 ml per pediatric bottle. Once the aerobic bottle has been inoculated, remove the adapter cap and repeat the procedure for the anaerobic bottle. Release the tourniquet as soon as the blood starts to flow into the bottle, or within 2 minutes of application.

The use of blood collection adapters without blood collection sets is not recommended.



7> Other blood tests

If blood is being collected for other tests, place an insert into the adapter cap. The insert is used to guide blood collection tubes onto the needle. **If other blood tests are required, always collect the blood culture first.**



8> Finish the procedure

Discard the winged collection set into a sharps container and cover the puncture site with an appropriate dressing. Remove gloves and wash hands before recording the procedure, including indication for culture, time, site of venipuncture, and any complications. **Ensure additional labels do not cover the bottle barcodes and that the tear-off barcode labels are not removed.**





**NATIONAL
ORTHOPAEDIC
HOSPITAL**
CAPPAGH

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**National Orthopaedic Hospital Cappagh,
Finglas, Dublin 11.**

PATHOLOGY LABORATORY USER MANUAL

Recommendations for blood culture collection

a summary of good practice

Using needle and syringe

1> Prepare blood collection kit

Gather all materials before beginning the procedure. Ensure the blood culture bottles are within date. Do not use bottles which show any signs of damage, deterioration or contamination.



2> Prepare bottles for inoculation

Wash hands with soap and water then dry, or apply an alcohol hand rub. Remove the plastic "flip-cap" from the blood culture bottles and disinfect the septum using an appropriate disinfectant, such as 2% chlorhexidine in 70% isopropyl alcohol, 70% isopropyl alcohol, or iodine in swab or applicator form. Use a fresh swab/applicator for each bottle. Allow bottle tops to dry in order to fully disinfect.



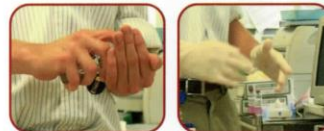
3> Prepare venipuncture site

Confirm the patient's identity. If skin is visibly soiled, clean with soap and water. Apply a disposable tourniquet. Palpate to identify the vein and cleanse using an appropriate disinfectant, such as 2% chlorhexidine in 70% isopropyl alcohol, 70% isopropyl alcohol, or iodine in swab or applicator form. **The venipuncture site is not fully clean until the disinfectant has fully evaporated.**



4> Wash hands. Wear gloves.

Wash hands again or use an alcohol hand-rub and apply clean examination gloves. Sterile gloves are not necessary.



5> Venipuncture

Attach the needle to a syringe. To prevent contaminating the puncture site, **do not re-palpate the prepared vein before inserting the needle.** Carefully insert the needle into the vein.



6> Culture bottle inoculation


Collect the sample. Transfer the blood into the culture bottles, starting with the anaerobic bottle. Hold the bottle upright and use the graduation lines to accurately gauge sample volume. Add up to 10 ml of blood per adult bottle and up to 4 ml per pediatric bottle.



7> Finish the procedure

Discard the needle and syringe into a sharps container and cover the puncture site with an appropriate dressing. Remove gloves and wash hands before recording the procedure, including indication for culture, time, site of venipuncture, and any complications. **Ensure additional labels do not cover the bottle barcodes and that the tear-off barcode labels are not removed.**



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BIOCHEMISTRY

Clinical Biochemistry deals with the biochemical basis of disease and the use of biochemical tests for its diagnosis, prognosis and management. Renal (sodium, potassium, Chloride, urea, creatinine, calculated eGFR), Liver (total protein, albumin, calculated globulins, bilirubin, alkaline phosphatase, ALT and Gamma GT), bone profile (albumin, calcium, phosphate, calcium adjusted for albumin and alkaline phosphatase) and Iron profile (Iron, Transferrin, calculated Transferrin Saturation, Ferritin, Folate and Vitamin B12) are carried out on a daily basis as requested. Blood Gases, which includes: pH, pCO₂, pO₂ std bicarbonate and Lactate are measured shortly after collection due to the rapid rate of change of the values in both the specimen and patient. Other tests include High Sensitivity Troponin I, Glucose, C- Reactive Protein (CRP) and Magnesium. A Blood Gas Analyser is located in NOHC HDU, which measures Blood Gases, Electrolytes (Sodium and Potassium ion concentration), Total Haemoglobin and Metabolites (Lactate and Glucose concentration). Some low volume tests are referred to other hospitals and these are listed from page 39. Routine chemistry samples may be processed on the Beckman Coulter AU chemistry platform and/or the Beckman Coulter Access 2 Immunoassay analyser, depending on the test required.

Note: One lithium heparin 4.9ml is sufficient to measure renal, liver, bone profiles, magnesium and CRP.

Turn Around Times

Urgent specimens: 1 hour

HDU/Fast track specimens: 2 hours (priority)

Routine specimens: 8 hours (same day)

Iron studies 24 hours (Weekdays)

Hours of service:


Monday – Friday 8.00 am to 8.00pm, 6.00 pm to 8.00 pm for urgent specimens only.

Monday – Friday 7.00am to 8.00am, the on call medical scientist is on site to process pre-op bloods for Same Day Admission (SDA) Patients.

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A limited off site on call emergency service is available outside these hours.


All Biochemistry tests as listed in the below tables, are available on call, with the exception of iron, transferrin, Ferritin, Folate and Vitamin B12.

	Test	Specimen Bottle	Units	Reference Range	Whole Blood Sample Viability at Room Temp **	Stability after separation (Room temp)	Storage Requirements
RENAL	Sodium	Li Hep 4.9 ml	mmol/l	135-145	24 Hours	7 days	Room Temp
	Potassium	Li Hep 4.9 ml	mmol/l	3.5-5.3	4 Hours	7 days	
	Chloride	Li Hep 4.9 ml	mmol/l	95-108	24 Hours	7 days	
	Urea	Li Hep 4.9 ml	mmol/l	2.6-6.8 (F) P<17yrs 2.6-7.5 (M) P<17yrs 2.5-7.8 Adult	24 Hours	7 days	
	Creatinine	Li Hep 4.9 ml	umol/l	22-41 P<5yrs 30-57 P<12yrs 43-75 P<15yrs 46-77(F)P<18yrs 58-99(M)P<18yrs 49-90 (F) Adult 64-104(M) Adult	24 Hours	7 days	
	eGFR	Calculated	mL/min	See below	N/A	N/A	N/A
LIVER	Total Protein	Li Hep 4.9 ml	g/l	59-72 P<6yrs 62-74 P<9yrs 63-77 P<18yrs 60-80 Adult	24 Hours	6 days	Room Temp
	Albumin	Li Hep 4.9 ml	g/l	38-47 P<8yrs 39-49 P<15yrs 38-51(F) P<18yrs 40-52(M)P<18yrs 35-50 Adult	24 Hours	7 days	
	Total Bilirubin	Li Hep 4.9 ml	umol/l	2.0-8.0 P<9yrs 2.0-10 P<12yrs 3.0-13 P<15yrs 3-15 P<18yrs <21 Adult	24 Hours	1 day (7days-In fridge)	

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	Globulin	Calculated (TP-ALB)	g/l	18-36	N/A	N/A	
	Alkaline Phosphatase	Li Hep 4.9 ml	IU/l	160-381 P<10yrs 144-475 P<13yrs 62-288(F) P<15yrs 130-534(M) P<15yrs 54-131 (F) P<17yrs 90-377 (M) P<17yrs 48-96(F) P<18yrs 59-168(M) P<18yrs 30-130 Adult	24 Hours	7 days	
	ALT	Li Hep 4.9 ml	IU/l	9-23 P<13yrs 8-22 P<18yrs <50 (M) Adult <35 (F) Adult	24 Hours	3 days (7daysIn fridge)	
	Gamma GT	Li Hep 4.9 ml	IU/l	5-13 P<11yrs 6-17 P<18yrs 3-38 (F) Adult 3-55 (M) Adult	24 Hours	7 days	Room Temp
BONE	Albumin	Li Hep 4.9 ml		38-47 P<8yrs 39-49 P<15yrs 38-51(F) P<18yrs 40-52(M)P<18yrs 35-50 Adult	24 Hours	7 days	Room Temp
	Calcium	Li Hep 4.9 ml	mmol/l	2.20-2.70 P<13yrs 2.20-2.65 Adult	24 Hours	7days	
	Adjusted Calcium	Calculated (corrected for Albumin)	mmol/l	2.20-2.60	N/A	N/A	
	Inorganic Phosphate	Li Hep 4.9 ml	mmol/l	1.4-2.33 P<5yrs 1.28-1.98 P<13yrs 0.93-1.87(F) P<16yrs 1.05-2.10(M) P<16yrs 0.93-1.63 P<18yrs 0.8-1.5 Adult	4 Hours	1 day (4 days in the fridge)	
Iron Profile	Iron	Li Hep 4.9 ml	µmol/l	3.0-23 P<14yrs 4.0-29(F) P<18yrs 6.0-30(M) P<18yrs 10.7-32 (F) Adult 12.5-32(M) Adult	4 Hours	7 days	Room Temp
	Transferrin	Li Hep 4.9 ml	g/l	2.14-3.30 P<18yrs	24 Hours	7 days	Room Temp

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				2.0-3.6 Adult			
	Transferrin Saturation	Calculated	%	10-50% (F) 17-55% (M)	N/A	N/A	N/A
	Ferritin	Li Hep 4.9 ml	ug/l	11-306.8 (F) 23.9-336.2 (M)	4 Hours	2 Days if in the Fridge	Freeze if testing after 2 days
	Folate	Li Hep 4.9 ml	ug/l	5.9-22.3	4 Hours	2 Days if in the fridge	Freeze if testing after 2 days
	Vitamin B12	Li Hep 4.9 ml	ng/l	122-626	4 Hours	24hrs if in the fridge	Freeze if testing after 1 day


The laboratory must be notified prior to ABG specimens being taken.

	Test	Specimen Bottle	Units	Reference Range	Whole Blood Sample Viability at Room Temp	Stability after separation (Room temp)	Storage Requirements
Blood Gas*	pH	Heparinised Syringe safe Pico self-fill with needle		7.35-7.45	30 minutes	N/A	Room Temp
	PCO2		kPa	4.5-6.0		N/A	
	PO2		kPa	11-14.0		N/A	
	Base Excess		mmol/l	-2.0 - +3.0		N/A	
	STD Bicarbonate		mmol/l	22.4-25.8		N/A	
	Oxygen Saturation		percentage	95-98%		N/A	
	cLactate		mmol/l	0.5 – 2.0		N/A	
	cGlucose		mmol/l	3.5 – 6.0		N/A	
	ctHb		g/dL	M 13.0-18.0 F 11.5-16.5		N/A	
	Magnesium	Li Hep 4.9 ml	mmol/l	0.83-1.13 P<18yrs 0.7-1.0 Adult	4 Hours	7 days	Room Temp
	Glucose	Fluoride Oxalate 2.7ml	mmol/l	3.9-5.5 (Fasting) WHO	4 Hours	2 days (7 days in the fridge)	Room Temp
	C-Reactive Protein	Li Hep 4.9 ml	mg/l	0 – 6	24 Hours	7 days	Room Temp

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	High Sensitivity Troponin I	Li Hep 4.9 ml	ng/L	M <20, F <12 Refer-to algorithm ED-BIO-50	8 Hours	8 Hours	Room Temp
Antibiotic Level in blood: (Vancomycin)	Vancomycin	Li Hep with gel 4.9 ml	mg/l	Once daily/BD dosing-desirable level (Predose/Trough) 10-20 mg/l Complicated Infection-desirable level (predose/trough) 15-20 mg/l Continuous Infusion-desirable level (random) 20-25 mg/l For children aged 1 month to 17 years: (predose/trough) 10-15 mg/l (15-20 mg/L for less sensitive strains of MRSA	24 Hours	7 days	Room Temperature

**** Stability references reviewed in accordance with Consultant Clinical Biochemist:**

“Stability study of 81 analytes in human whole blood, in serum and in plasma

Christiane Oddoze , Elise Lombard, Henri Portugal”

<https://www.sciencedirect.com/science/article/abs/pii/S0009912012000197?via%3Dihub>


Information is taken from the manufacturer’s kit insert unless otherwise stated.

Paediatric reference ranges taken from Adeli K,Higgins V, Trajcevski K, Al Habeeb (2017). “The Canadian Laboratory initiative on pediatric reference intervals: A Caliper white paper”.

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Adult reference ranges taken from the Pathology Harmony project were available and the Kit inserts.
Blood gas * reference ranges taken from MMUH in-house study.

Add on requests

Samples are stored in the lab for 7 days. Additional testing can be requested on samples if within the test stability stated above. If requesting an add on request, send a form with add on request and the test(s) required to the lab.

Stratification of Chronic Kidney Disease by eGFR

Stage Description	GFR (ml/min/1.73m²)
1. Kidney damage with normal	GFR >90 with other evidence of chronic kidney damage*
2. Mild reduction in GFR	GFR 60-89 with other evidence of chronic kidney damage*
3. Moderate reduction in GFR	GFR 30-59
4. Severe reduction in GFR	GFR 15-29
5. Kidney Failure	GFR <15 or RRT

- Estimated GFR is calculated using the 4v-MDRD Formula
- Estimates of GFR are unreliable in acute renal failure due to the kinetics of creatinine accumulation.
- GFR estimates between 60 and 89 mL/min/1.73m² do not indicate CKD unless there is other laboratory/clinical evidence of disease.
- Estimated GFR should be multiplied by 1.212 for patients of African ethnicity.
- The formula is applicable to adult patients (i.e. patients >18 years old).
- Values will not be given for results of eGFR >60ml/min/1.73m²
- The reason is that the MDRD equation was derived in patients with CKD and the formula has proven to be inaccurate in that it underestimates the GFR at higher values.


Interfering Substances

Many tests are subject to interference. This may be biological, where the offending substance alters the true concentration within the body, or analytical, where the method is not specific. The report will outline the more common interfering substances such as haemolysis, Icteria (bilirubin interference) and Lipaemia(lipoprotein interference). Depending on the degree of interference, some assays may not be reportable.

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Abbreviations:

M – Male

F – Female

P < Paediatric range less than

A - Adult

Li Hep – Lithium Heparin

ALP – Alkaline Phosphatase

ALT – Alanine Aminotransferase

cCa²⁺ - corrected calcium

eGFR Estimated Glomerular Filtration Rate

TSat Transferrin Saturation

CRP – C-reactive protein

ctHb – Total Haemoglobin (concentration)

* Non-accredited test

NOTE* Reference Ranges quoted are both adult and paediatric unless otherwise indicated by letter P

HAEMATOLOGY

The Haematology Laboratory deals with investigations into general haematological diseases including anaemia, and monitoring the blood coagulation status of the patients.


The following tests are carried out in Haematology:

- Full blood counts which include red cell counts, white cell counts and differential, haemoglobin, haematocrit, red cell indices such as mean cell volume (MCV), mean cell haemoglobin (MCH) and mean cell haemoglobin concentration (MCHC), and platelet counts
- Coagulation studies which include prothrombin time, activated partial thromboplastin time and fibrinogen
- Erythrocyte sedimentation rate
- Blood Films
- B12, Folate and Ferritin

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Some low volume tests are referred to other hospitals and these are listed from Page 37. Samples may be processed on the CELL DYN, ACL TOP 550 and Access 2 Immunoassay (test dependant)

Please note all samples run on the ACL TOP 550 are accompanied on the report denoting they are outside the scope of accreditation.

Turn Around Times

Urgent specimens: 1 hour

HDU/Fast track specimens: 2 hours (priority)

Routine specimens: 8 hours (same day)

Blood Films: 24 hours

Hours of service:

Monday – Friday 8.00 am to 8.00pm, 6.00 pm to 8.00 pm for urgent specimens only

Monday – Friday 7.00 am to 8.00am, the on call medical scientist is on site to process pre-op bloods for Same Day Admission (SDA) Patients.

A limited off site on call emergency service is available outside these hours.

All Haematology tests as listed in the below tables, are available on call, with the exception of B12, Folate, Ferritin and blood films. These are measured in batches routinely only.


Sample Requirements for Haematology

	Test	Specimen Bottle	Units	Reference Range	Specimen Viability	Storage Requirements
FBC	WBC	2.7ml EDTA	x10 ⁹ /L	Male and Female 4.0-11.0 (≥14yrs)	8 Hours	Room Temperature
				≤1 year 6.0-18.0		
				≤8 years 5.0-15		

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				9-13 years 4.5-13.5		
	RBC	2.7ml EDTA	x10 ¹² /L	Male 4.5-5.3 13- 18 years 4.5-6.5 (≥19yrs)		
				Female 4.1-5.1 13-18 years 3.8-5.8 (≥19yrs)		
				91 days 3.1 - 4.5 2 years 3.7-5.3 3-6years 3.9-5.3 7-12 years 4.0-5.2		
	HB	2.7ml EDTA	g/dl	Male 13.0-18.0 (≥14yrs)		
				Female 11.5-16.5 (≥14yrs)		
				63 day: 9.0- 14.0 18 month: 10.5-13.5 ≥3 years: 10.5-13.5 4 to 7 years: 11.5-14.5		

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				8 to 13years: 11.5-15.5		
	HCT	2.7ml EDTA	%	Male: 40.0-54.0 (≥14yrs)		
				Female: 37.0-47.0 (≥14yrs)		
				98 days: 29.0-41.0 ≥3 years: 33.0-39.0 4 to 13years: 35.0-45.0		
	MCV	2.7ml EDTA	fl	M/F 80-95 (≥14yrs)		
				98 days: 74-118 ≥3 years 70-86 5 to 6 years 75-87 7 to 13years 77-96		
	MCH	2.7ml EDTA	pg	M/F 28.0-32.0 (≥14yrs)		
				95 day: 25-35 ≥3 years 23-31 4 to 7years 24-30 8 to 13years 25-33		

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	MCHC	2.7ml EDTA	g/dl	M/F 32.0-36.0 (≥7yrs) 1 year 32.1-36.5 ≥6years 31-37	8 Hours	Room Temperature
	RDW	2.7ml EDTA	%	M/F/Paeds 11.0-16.0		
	PLATELET	2.7ml EDTA	x10 ⁹ /L	M/F/Paeds 150-400		
	MPV	2.7ml EDTA	fl	M/F/Paeds 7.5-10.5		
	Neutrophils	2.7ml EDTA	x10 ⁹ /L	M/F 2.0-7.5 ≥17years ≤ 3 days 1.5 – 7.0 ≤ 2 year: 1.0 – 8.5 3-6 years 1.5-8.5 7-12 years 1.5-8.0 13-16years 1.8-8.0		
	Lymphocytes	2.7ml EDTA	x10 ⁹ /L	M/F 1.5-4.0 ≥17years ≤ 3 days 2.0 – 5.0 ≤ 2 year: 3.0 – 13.5 3-6 years 2.0-9.5 7-12 years 1.5-7.0		

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				13-16years 1.2-5.2		
	Monocytes	2.7ml EDTA	$\times 10^9/L$	M/F 0.2-0.8 ≥ 17 years ≤ 3 days 0.3– 1.1 ≤ 2 year: 0.3 – 1.5 3-6 years 0.3-1.5 7-12 years 0.1-1.8 13-16years 0.1-0.8		
	Eosinophils	2.7ml EDTA	$\times 10^9/L$	M/F 0-0.4 ≥ 17 years ≤ 3 days 0.2 – 2.0 ≤ 2 year: 0.1- 0.3 3-6 years 0.3-0.8 7-16 years 0.1-		
	Basophils	2.7ml EDTA	$\times 10^9/L$	M/F 0-0.1 ≥ 17 years ≤ 3 days 0.02 – 0.1 ≤ 2 year: 0.02 – 0.1 3-6 years 0.02-0.1 7-16 years 0.0-0.2		

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Coag	PT	3.0ml Citrate	Seconds	10.8-13.4 (>17years)	4 Hours	Room Temperature
				1-5 years (9.9-13.4)		
				6-10 years (10.0-14.6)		
				11-17 years (10.0-14.1)		
	APTT	3.0ml Citrate	Seconds	25 – 36.5 (>17years)		
				1-5 years 24.0-39.2		
				6-10 years 26.9-38.7		
				11-17 years 24.6-38.4		
	Fibrinogen	3.0ml Citrate	g/l	1.5-4.0 (>17years)		
				1-5 years 1.64-4.97		
				6-10 years 1.71-5.37		
				11-17years 1.68-5.29		
	ESR	Sedivette ESR/Citrate	mm/hr	Male 0-10	24 Hours	Refrigerate

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		Buffer 3.5ml		(≥14 years)		
				Female 0-20 (≥14 years)		
				Neonate – Puberty: 3-13		
	Blood Film	2.7ml EDTA			8 Hours	Room Temperature
				Female/Paeds 11 – 306.8		


Abbreviations:

WBC	– White Blood Cell	PT	– Prothrombin Time
RBC	– Red Blood Cell	APTT	– Activated Partial Thromboplastin Time
HB	– Haemoglobin	ESR	– Erythrocyte Sedimentation Rate
HCT	– Haematocrit	EDTA	– Ethylenediaminetetraacetic acid
MCV	– Mean Cell Volume	MPV	– Mean Platelet Volume
MCH	– Mean Cell Haemoglobin	RDW	– Red Cell Distribution Width
MCHC	– Mean Cell Haemoglobin Concentration		
VB12	- Vitamin B12	FOLW	– Folate
FER	- Ferritin		
Paeds	- Paediatrics		

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LABORATORY CRITICAL VALUES: (AP-LAB-4)

BIOCHEMISTRY					
Test	Sex	Age (Yrs.)	Code	Less than or equal to (\leq)	Greater than or equal to (\geq)
Urea (plasma)	M/F		URE		25.0 mmol/L
Sodium (plasma)	M/F		NA	120 mmol/L	155 mmol/L
Potassium (plasma)	M/F		K	2.8 mmol/L	5.5 mmol/L
Chloride (plasma)	M/F		CL	70 mmol/L	128 mmol/L
Creatinine (plasma)	M/F		CREA		354 μ mol/L and/or 3x baseline if known as per KDIGO
Calcium (adjusted) (plasma)	M/F		CA	1.8 mmol/L	2.90 mmol/L
Glucose (Fasting/Random) (plasma)	M/F		GLU	2.5 mmol/L	25 mmol/L
C-Reactive Protein (plasma)	M/F		CRP		300 mg/L
hsTroponin I (plasma)	M/F		TNI		M 20 ng/L F 12 ng/L
ALT (plasma)	M/F		ALT		600 IU/l
Phosphate (plasma)	M/F		PO4	0.30 mmol/L	2.50 mmol/L
Magnesium (plasma)	M/F		Mg	0.40 mmol/L	1.50 mmol/L
Lactate (whole blood)	M/F		LAC		4.0 mmol/L
Vancomycin (plasma)	M/F		VANC		25.0 mg/L

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ED-HAEM-69 Critical Values/Panic Values in Haematology in accordance with Temple Street Hospital
Please note NOHC reports Hb and MCHC in g/dl Critical Values/Panic Values in Haematology
Primary critical results are telephoned ASAP to the clinical team

Parameter	Value	Comment
Haemoglobin	<66 g/l >170 g/l	Supply of oxygen to the myocardium inadequate. Corresponds to Haematocrit of 51% and leads to hyper viscosity syndrome
Haematocrit	55%	N/A
Platelet count	< 20 x 10 ⁹ /l If known patient < 100 if first time patient > 1000 x10 ⁹ /l	Risk of Haemorrhage. Exclude EDTA induced thrombocytopenia, New Leukaemia Risk of thrombosis
White cell count	< 2 x10 ⁹ /l >50 x10 ⁹ /l (1 st time) Blood film Review: <ul style="list-style-type: none"> • Suspected leukaemia • Suspected Aplastic crisis • Sickle cells • Malarial parasites 	High risk of infection if the granulocyte count is <0.5 x10 ⁹ /l Indicative of leukemoid reaction e.g. in sepsis or of leukaemia.
Prothrombin Time (PT)	>20 secs	Decrease in the vitamin K-dependent factors II, VII and X or in factor V. Since all these factors are synthesized in the liver, a decrease in the PT to values below the specified level indicates a considerable disturbance of synthesis.
Activated Partial Thromboplastin Time (APTT)	≥ 75 secs	Deficiency or inactivity of factor VIII, IX, XI or XII with risk of Haemorrhage. In patients receiving heparin therapy there is a risk of haemorrhage if the APTT is more than 2.5 times higher than the upper reference limit.
Fibrinogen	<1 g/l	Risk of Haemorrhage and hypo coagulopathy

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D-Dimers	Positive	Positive may be suggestive of thrombosis
Malaria Screen	Positive	Delay in diagnosis and treatment is a leading cause of death
Neonatal quantitative limits of Laboratory results which, after confirmation through repeat measurement in the same sample, need urgent notification to the Clinician.		
Haemoglobin	< 85 g/l >230 g/l	Risk of multi-organ failure, especially if the patient has a combination of ischemia and hypoxia Abnormal flow kinetics (hyper viscosity) with increased circulatory resistance and an increased load on the heart.
Hematocrit	< 33% (l/l) > 71% (l/l)	Indicative of marked anaemia with an inadequate supply of oxygen to tissue. Hyper viscosity of the blood with increased circulatory resistance.
Platelet Count	<100 x10 ⁹ /l >1000 x10 ⁹ /l	Risk of Haemorrhage. Exclude EDTA induced thrombocytopenia Risk of thrombosis
White Cell Count	< 5 x 10 ⁹ /L > 25 x 10 ⁹ /L Blood film Review: <ul style="list-style-type: none"> • Suspected leukaemia • Suspected Aplastic crisis • Sickle cells • Malarial parasites 	Values below and above these limits can be indicative of neonatal sepsis.
Prothrombin Time (PT)	>18 secs	N/A
Activated Partial Thromboplastin Time (APTT)	≥ 50 secs	N/A
Fibrinogen	<1.0 g/dl	N/A

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HAEMATOLOGY					
Test	Sex	Age(Yrs)	Code		
	Pre-Operatively				
White Blood Cell	M/F/U	0-12	WBC	<3.0 x 10 ⁹ /L	>20.0 x 10 ⁹ /L
	M/F/U	13-121		<3.0 x 10 ⁹ /L	>20.0 x 10 ⁹ /L
Neutrophils	M/F/U	13-121	NEUT	<1.0 x 10 ⁹ /L	>20.0 x 10 ⁹ /L
Lymphocytes	M/F/U	13-121	LYMP	-	>6.0 x 10 ⁹ /L
Haemoglobin	M/F/U	0-12	HB	<9.5 g/dl	>16.0 g/dl
	M/F/U	13-121		≤6.0 g/dl	>20.0 g/dl
				≤8.0g/dl-phone for transfusion purposes	
				≤7.0g/dl-phone to request repeat Or unexpected drop in Hb	
Platelets	M/F/U	0-12	PLT	≤50 x 10 ⁹ /L	>800 x 10 ⁹ /L
	M/F/U	13-121		≤80 x 10 ⁹ /L	>800 x 10 ⁹ /L
				<100 Phone result following film check	
				≤80 phone to request repeat	
	Post Operatively				
White Blood Cell	M/F/U	0-12	WBC	<3.0 x 10 ⁹ /L	>20.0 x 10 ⁹ /L
	M/F/U	13-121		<3.0 x 10 ⁹ /L	>20.0 x 10 ⁹ /L
Haemoglobin	M/F/U	0-12	HB	<9.5 g/dl	<16.0 g/dl
	M/F/U	13-121		≤6.0 g/dl	>20.0 g/dl
				≤8.0g/dl-phone for transfusion purposes	
				≤7.0g/dl-phone to request repeat Or unexpected drop in Hb	
Platelets	M/F/U		PLT	≤50 x 10 ⁹ /L	≥800 x 10 ⁹ /L
				<100 Phone result following film check	
				≤80 phone to request repeat	
Preoperative/Post Operative					
PT (INR)	M/F		PT		>4.5 phone to request repeat
					>6.0
APTT	M/F		APTT		>40 phone to request repeat
					≥120 Sec
Fibrinogen	M/F		FIBR	<1.5g/l	

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BLOOD TRANSFUSION

The Blood Transfusion Laboratory performs a range of tests including blood grouping, antibody screening and compatibility testing. This testing is to ensure the right blood gets to the right patient in a timely manner. A range of blood products are available from the blood transfusion laboratory- red cells, plasma, platelets, prothrombin complex and fibrinogen.

Turn Around Times:

Routine specimens: Same Day

Urgent group and screen: 1 hr (antibody screen negative)

Crossmatch: 60 minutes*

*In the case of crossmatching for Sports Surgery Clinic, additional logistics (transfer by taxi) may result in increased turnaround times.

Hours of service:

Monday – Friday 8.00am to 8.00pm, 6.00 pm to 8.00 pm for urgent specimens only

Monday – Friday 7.00am to 8.00am, the on call medical scientist is on site to process pre-op bloods for Same Day Admission (SDA) Patients.

A limited off site on call emergency service is available outside these hours.

The following tests are available on call:

Group and hold, crossmatching and issuing of blood products (Red Cell, Platelets, Plasma, Prothrombin Complex Concentrate (**PCC**) and Fibrinogen)


Samples may be processed manually (using tube technology or gel technology) or automatically (using gel technology on the IH-500).

Samples for HLAB27 are referred to the IBTS. Refer to page 93.

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In line with the BSH Guidelines, Appendix 7, two (2) Group and Hold Specimens are required prior to the issue of blood group specific red cell units. This requires two (2) separate Group and Hold specimens to be taken on two separate episodes.

Guide to pre-transfusion collection of samples:

- If the patient **has not** been pregnant or transfused in the previous three (3) months, the patient's sample is valid for **seven (7) days**.
- If the patient **has** been pregnant or transfused in the previous three (3) months the patient's sample is valid for **three days (72 hours)**.
- This testing frequency is essential because of the possibility of antibody development in patients exposed to red cell antigens from transfusion or pregnancy during the referred intervals. This will ensure compatible blood is available for patients.

If 72 hours have elapsed since the patient was transfused and further transfusion is contemplated, then a new sample must be sent for group and antibody screen and crossmatching. This is required for the confirmation of antibody status, ensuring the correct blood is issued at all times. EDTA samples are retained in the laboratory for 7 days after being processed and are then discarded. Samples that have been used for crossmatch are retained in the laboratory for 14 days after the crossmatch.


Crossmatched units for NOHC patients are stored in the 08-BT-003 Recovery Room Blood Issue Fridge, unless otherwise stated (Please check Blood Track or phone the Blood Transfusion laboratory to confirm the availability of blood products). Crossmatched units for SSC patients are stored in the Blood Transfusion laboratory until they are required for transfusion. Crossmatched units and blood products are issued to the patient for 48 hours before being returned to stock in the laboratory.

Four emergency O Rhesus D Negative, Kell Negative, CMV Negative and group confirmed blood units are held on standby in the Blood Transfusion Department at all times. This is to cater for emergency situations.

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A notice period is required for specific requests regarding ordering of blood products:

- CMV negative and irradiated blood units Routine (at least 24 hours for SSC)
- Rhesus/ Kell typed blood units At least 24hrs Notice
- Other Antibodies or multiple antibodies At least 48hrs Notice


Sample Requirements for Blood Transfusion

Test Profile	Specimen Bottle	Profile Includes	Results available/Comments	Sample viability	Storage
Group and Hold	7.5ml/4.5ml EDTA (Red Writing)	Blood Group Antibody Screen	Specimens are analysed on the Biorad IH-500 analyser and reported same day. Automated blood group and antibody screen – 60 minutes. If an urgent blood group and antibody screen is required, the group and screen may be carried out manually and a result may be available within 35 – 40 minutes, (if antibody screen is negative).	7 Days (72 hours if patient has been transfused /pregnant in last 3 months)	4°C
			If negative, the results will be available on the same day. Positive results will necessitate further investigation.		
Issuing of Blood Products:	7.5ml/4.5ml EDTA (Red Writing)	Red cells Octaplas, PCC & Fibrinogen Platelets	For an urgent Immediate spin cross match -20 minutes . FULL crossmatch – 60 minutes Octaplas (Frozen Plasma) are defrosted and issued in 25- 30 minutes . Fibrinogen and PCC are stored on site and issued in 10 minutes . Platelets are not stored on site, so will have to be ordered in from the IBTS, which may take between 1.5–2 hours to be delivered and issued from ordering	7 Days (72 hours if patient has been transfused /pregnant in last 3 months)	4°C
Antibody Panels	7.5ml/4.5ml EDTA (Red Writing)		If a positive antibody screen is detected, extensive panels are carried out to	7 Days (72 hours)	4°C

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	Writing) by 2		ascertain antibody identification. This assists in provision of antigen negative blood. Turnaround dependant on the complexity of the antibodies present. If the antibody cannot be identified in-house; the sample is referred to the IBTS. (Turnaround time (TAT) usually 10 working days)	if patient has been transfused /pregnant in last 3 months)	
Direct Coombs Test (DAT/DCT)	7.5ml/4.5ml/2.7ml EDTA		The DAT/DCT is a test used to determine if red cells have been coated in vivo with immunoglobulin, complement or both. A positive DAT may or may not be associated with immune-mediated haemolysis. The DAT is used primarily for the investigation of haemolytic transfusion reactions (HTR's), auto-immune haemolytic anaemia (AIHA), drug-induced haemolysis and haemolytic disease of the newborn. Occasionally a DAT is performed on a donor unit if the crossmatch is unexpectedly positive due to the donor having a positive DAT.	7 Days	4°C
Red Cell Phenotyping	7.5ml/4.5ml EDTA (Red Writing)		Red cell Units are occasionally typed for antigen specificity. This is carried out in house rather than at the IBTS. Patient samples are also phenotyped to assist in antibody identification. TAT One day.	7 Days (72 hours if patient has been transfused /pregnant in last 3 months)	4°C
Transfusion Reaction	7.5ml EDTA/4.5ml (Red Writing) x2, 2.7ml EDTA sample (red lid), Lith. Heparin (orange lid), Coag sample (green lid),	Various tests FBC (Blood Film) Coagulation DAT (DCT) Full Screen Biochemistry. Microbiology IgA levels	A thorough investigation of a transfusion reaction may take some time. If blood units are required before the investigation is complete, either the haematology consultant is contacted and the case is discussed, or the requesting physician takes full responsibility for the issue and administration of the unit(s)	Until a Medical Scientist who works routinely in Blood Transfusion is able to process the Transfusion	4°C


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	First MSU, Blood Cultures Residue from any units transfused (including all units crossmatch ed and the giving set of the suspected unit.)			n reactions	
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Maximum Surgical Blood Ordering Schedules at NOHC: (AP-HVO-8)

UPPER LIMB SURGERY	
Shoulder Replacement	Group & Hold
Revision Shoulder Replacement	Group & Hold
Elbow Replacement	Group & Hold
Revision Elbow Replacement	Group & Hold
SPINAL SURGERY	
Removal of Spinal Instrumentation	Group & Hold
Spinal Decompression	Group & Hold
Adolescent Posterior Spinal Fusion (For Scoliosis)	Group & Hold
All other spinal fusions	Group & Hold
LOWER LIMB SURGERY	
THR	Group & Hold
Hip Resurfacing	Group & Hold
Conversion THR	Group & Hold
Revision THR	Group & Hold
Infected Revision THR/1 st Stage and 2 nd Stage	Group & Hold
Hip Osteotomy	Group & Hold
Hip Arthrodesis	Group & Hold
Disarticulation	X-Match 4 Units
Removal of Femoral Plate	Group & Hold
Amputation	Group & Hold
Enbloc Resection	Group & Hold
TKR	Group & Hold
Revision TKR	Group & Hold
Knee Arthrodesis	Group & Hold
Patellar Resurfacing	Group & Hold
Total Ankle Replacement	Group & Hold
Revision Total Ankle Replacement	Group & Hold
Ankle Arthrodesis	Group & Hold


1 unit of blood will be X-matched for all hip and knee surgical patients with multiple or unusual red cell antibodies as notified by the Transfusion Laboratory.

X-matching will be carried for any patient, as requested by a Surgeon or Anaesthetist.

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MICROBIOLOGY

The Microbiology Department provides a range of diagnostic services in routine microbiology. Referral for mycobacteriology, parasitology and virology serology (including bone bank and post Hepatitis B vaccination levels) and any specialised microbiology can be arranged through the Department.

Some low volume tests are referred to other hospitals and these are listed from Page 39.

Please note that SSC Microbiology samples excluding MRSA screens, CPE screens & Urine samples (which are processed in-house) are referred to the Mater Private Hospital directly by SSC. The Microbiology Department supports the National and NOHC infection control policies. Theatre air sampling and environmental monitoring is carried out when sanctioned by the Consultant Microbiologist.

It is necessary to discuss processing of urgent samples including direct Gram staining of fluid samples and tissue samples with a member of staff in Microbiology when taking the samples. Please provide as much information as possible on the request form including site of sample, clinical details and antibiotic therapy, to assist with the interpretation of the results.

The blood culture analyser is monitored during routine hours, Monday – Friday (8.00 am to 6.00 pm) and weekends/Bank Holidays (9.00 am to 12:30pm only). The Medical Scientist in Microbiology will process the blood culture and will liaise with the Consultant Microbiologist/ Microbiology Registrar during these hours.

Hours of service:

Monday – Friday 8.00am to 6.00pm,

Saturday - 9.00 am to 12.30pm;

Sunday/Bank Holiday - 9.00 am to 12.30pm on standby for positive blood culture / COVID-19 processing only.


An urgent COVID-19 service for **symptomatic patients only** is provided outside of the above hours until midnight. Samples must reach the lab by 10pm.

Any other requests have to be sanctioned by the Consultant Microbiologist.

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Tests performed in NOHC Microbiology Department:

Microscopy, Culture and Antimicrobial Susceptibility testing of isolates from specimens including Blood Cultures, Aspirates, Tissue, Bone, Intraoperative swabs, Intravascular Cannulae, Urines and Superficial swabs.

Screening swabs for HCAI - MRSA, CPE and VRE.

SARS-CoV-2, Influenza & RSV testing.

Urinary hCG testing, and Bone Bank Sterility Testing.

Tests referred to other hospital laboratories:

CSF for analysis - cell count, culture and susceptibility testing.

Fluids for Crystal Determination and differential cell counts.

Faeces for enteric pathogen analysis, *C. difficile*, Norovirus and ova, cysts and parasites. (Please send individual samples for each test).

Sputum for culture and susceptibility testing. Sputum, Fluids and Tissues for AFB and Mycobacterium culture and susceptibility testing. Blood for Quantiferon testing.

Blood for virology testing and Bone Bank serology.


Sample Requirements for Microbiology:

Test	Container/ Swab	Results Available ^{1,2}	Comments	Sample viability	Storage
Aspirates: 1. Gram Stain 2. C/S 3. Crystal Determination 4. Cell count and differential	1, 2 and 3. Sterile double wrapped 30mL clear PS Universal Tube 4. EDTA Sample bottle	1. Urgent – 30-60 mins, Routine – 24hrs. 2. Negative culture report: 7 days, interim report released after 48hrs. Positive culture report: As soon as	If Gram stain is urgent, please contact the Microbiology laboratory prior to sending the sample. *Do not use wide MSU containers* Please transport to the laboratory ASAP	Up to 48 hrs if stored at 4°C	Processed ASAP if not refrigerate

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		available 3 & 4. Referred to MMUH			
Tissue/Bone: 1. Gram Stain 2. C/S	Sterile Double wrapped 30mL clear PS Universal Tube	1. 30-60 minutes if urgent or 24 hrs if routine 2. Negative culture report: 7 days. Positive culture report: As soon as available	If Gram stain is urgent, contact the Microbiology laboratory in advance of taking the specimen. Large tissue/Bone samples should be divided into Marble sized representative samples in the clean air environment. These should then be submitted in individual containers to the Microbiology Laboratory Do NOT add any fluid to tissue samples Please transport to the laboratory ASAP	Up to 48 hrs if stored at 4°C	Processed ASAP if not Refrigerate
Intraoperative swabs	Sterile Double wrapped Transport Swab with Amies medium – no charcoal	Negative culture report: 5 days. Positive culture report: As soon as available	N/A	Up to 48 hrs if stored at 4°C	Processed ASAP if not Refrigerate
Blood cultures*	BacT/ALERT Blood culture bottles – aerobic and anaerobic	Negative culture report: after 5 days or as soon as available if positive	Bottles kept on each ward, replenish stocks from microbiology. Please note expiry date. Immediately bring bottles to the Microbiology Laboratory for loading onto the BacT/ALERT blood culture instrument	24 hrs store at room temp	Processed ASAP. If not store at Room Temp Do NOT Refrigerate
Intravascular Cannulae - Central /PICC	Place in Sterile Urine or Universal Container	Negative culture - 48hrs Positive culture as soon as	N.B. Cannulae specimens will <u>only</u> be cultured if a blood culture sample is received simultaneously	Up to 48 hrs if stored at 4°C	Processed ASAP if not Refrigerate

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		available			
MSU	Sterile Urine Container	Microscopy - same day, Negative culture - 24hrs Positive culture – as soon as available	Mid-stream clean catch. Please close container lid tightly, leaking specimens will not be processed. Urine samples that are retained at room temperature for > 2 hrs may not be processed	Up to 48 hrs if stored at 4°C	Processed ASAP if not Refrigerate
CSU	Sterile Urine Container	Microscopy - same day, Negative culture - 24hrs Positive culture– As soon as available	Urine should be collected from tube leading into catheter. Urine samples that are retained at room temperature for > 2 hrs may not be processed.	Up to 48 hrs if stored at 4°C	Processed ASAP if not Refrigerate
Superficial Swabs	Transport Swab with Amies medium – no charcoal	Negative culture report - 48 hrs, positive culture -as soon as available	Please record site of swab on request form and any relevant clinical details and antimicrobial therapy	Up to 48 hrs if stored at 4°C	Processed ASAP if not Refrigerate
MRSA Screens	<u>Culture:</u> Transport Swab with Amies medium – no charcoal	Negative culture at 24 hrs, positive culture as soon as available may take 72 hrs	Minimum screening swabs of Nasal and Groin.	Up to 48 hrs if stored at 4°C	Processed ASAP if not Refrigerate
	<u>Molecular MRSA</u> For samples requiring same day MRSA results <u>only.</u> Stuart Dual Swab (900-	3-3.5 hours or earlier depending on workload	To be used for urgent MRSA requests only. Nasal and Groin swabs only Not suitable for patients <2years old.	24hrs at room temp. Up to 7 days if stored at 4°C	Processed ASAP if not Refrigerate


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	0370) Available from the Microbiology Department				
VRE Screens	Rectal swab taken in a Transport Swab with Amies medium – no charcoal	Negative culture - 48hrs Positive culture as soon as available	Taking of VRE screens is carried out in accordance with Infection Prevention and Control Guidelines. Screening to identify VRE colonised patients is recommended during outbreaks only and on the instructions of the Clinical Microbiologist	Up to 48 hrs if stored at 4°C	Processed ASAP if not Refrigerate
CPE Screens	Culture: Rectal swab taken in a Transport Swab with Amies medium – no charcoal. Additional samples from other sites (e.g. urine, swabs from skin breaks may also be suitable for surveillance purposes	Negative culture – up to 24 hrs Positive culture as soon as available	Taking of CPE screens is carried out in accordance with Infection Prevention and Control Guidelines. Other CPE Screening is recommended only on the instructions of the Clinical Microbiologist	Up to 48 hrs if stored at 4°C	Processed ASAP if not Refrigerate

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	<u>Molecular:</u> LQ Stuart Dual Rectal Swab (900- 0370) Available from the Microbiology Laboratory.	Negative: 2 hours Positive: As soon as available	Transport to the laboratory as soon as possible Swabs overly soiled with stool may not be suitable for analysis. Same day testing available if the sample is received into the laboratory by 4pm Monday to Friday	Up to 5 days if stored at 15-28°C	Processed ASAP, do not refrigerate
Influenza/RSV*	Nasopharyngeal Sample Collection Kit for Viruses Available from the Microbiology Laboratory.	Negative: 2 hours Positive: 2 Hours	Transport to the laboratory as soon as possible Same day testing available if the sample is received into the laboratory by 4pm Monday to Friday	Up to 7 days if stored at 2-8°C	Processed ASAP if not Refrigerate
SARS-CoV-2 (Covid-19) *	Nasopharyngeal Sample Collection Kit for Viruses. Available from the Microbiology Laboratory.	Urgent samples: Negative: 2 hours Positive: 2 Hours.	Transport to the laboratory as soon as possible Same day testing of urgent samples available if the sample is received into the lab by 22:00	Up to 7 days if stored at 2-8°C	Processed ASAP if not Refrigerate
*Pregnancy Test	Sterile Urine Container	Same day	Please phone lab for urgent requests	48hrs	Processed ASAP if not Refrigerate
Bone Bank Sterility Testing	Direct inoculation into 20ml Tryptone Soy Broth and Thioglycolate broth	Negative culture 16 days, positive culture as soon as available	Transport immediately to laboratory for incubation	48hrs	Incubated ASAP, if not do NOT Refrigerate

* Non-accredited tests


Please note that samples which do not meet the requirements specified in the above table, will be rejected.

Faeces for culture and sensitivity will only be processed if the patient has been hospitalised for < 3 days.

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¹These times are a typical guide and due to the nature of microbiological testing they may vary.

²The TAT times outlined above are bound by the hours of service and may be delayed accordingly due to limited microbiology weekend service or increased workloads at busy periods

CYTOLOGY/HISTOLOGY

NOHC Pathology Specimen Reception provides the transport service for NOHC Cytology/ Histology specimens to the Histology Laboratory at the MMUH.

The majority of surgical specimens should be sent to the NOHC Laboratory in the appropriate preservative (10% Neutral Buffered formalin).

Avoid squeezing specimens into small jars.

The body of the container must be labelled with the patients' details (Name, DOB and hospital number) and the specimen details. If more than 1 sample is taken then label the individual samples A, B, C etc. (with full details on each).

For NOHC Cytology/ Histology specimens complete the NOHC Histology Request form (RF-LAB-54) and provide a contact name and number.

For NOHC Cytology/ Histology specimens the appropriate specimen Log (RF-LAB-17 & RF-LAB-18) should be completed at source (Theatre / X-ray). Signed for specimens are to be transported by porters and signed into the Laboratory by a member of the laboratory staff.


SPECIMEN REQUIREMENTS FOR CYTOLOGY:

- Fluid samples such as joint fluids for query crystal analysis should always to be sent **fresh**, as well as fluids for query mucin, ground substance, colloid, collagen etc. These fluid samples **MUST NOT BE PLACED INTO FORMALIN**. Please notify the NOHC laboratory prior to taking these samples.

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- For cytology fluid samples **which do not require the former**, Cytolyt ® fixative should be used (1: 1 ratio) instead of 10% buffered formalin.
- Ganglion cysts for aspiration: if no crystal analysis is required, aspirated fluid must be placed directly into Cytolyt ® fixative. See Histology below for Ganglion cyst samples.

SPECIMEN REQUIREMENTS FOR HISTOLOGY

- Ganglion cyst samples: if no crystal analysis is required, the cyst must be placed directly into 10% Buffered Formalin. Any fluid aspirated from the Ganglion cyst must be treated as for Ganglion cysts for aspiration (see Cytology section above) prior to placing sample in Formalin.
- Frozen Section Analysis A frozen section service is provided in MMUH from 9.00 – 17.30, Monday to Friday. Frozen sections must be booked at least 24 hours in advance of elective surgery by phoning the MMUH Histology Laboratory at 01 803 2116.
- Specimen: Fresh tissue in an appropriately labelled sealed container. Delivered immediately to MMUH Histology laboratory.

Turnaround time: Same day

Special precautions: Specimens from patients who have/ suspected to have TB are NOT appropriate for frozen section. These specimens must be placed into appropriate fixative immediately. If a patient is suspected of having such an infection, the clinical staff must inform MMUH Histology laboratory


- Please refer to
 - PP-THE-1 (Procedure for Specimen and Amputated Limb Collection and Processing) on the Q-Pulse system.
 - PP-RAD-21 (Management of Biological Specimen Taken During an Interventional Radiological Procedure)
 - WI-LAB-18 Cytology/Histology Specimen Requirements

***** If in doubt about how to treat a Cytology or Histology specimen, please contact a member of the MMUH Histology staff before putting the specimen into a fixative on (01) 803 2116***


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Cytology/ Histology reports are received by NOHC Pathology Specimen Reception and forwarded to the NOHC oncology nurse. Copies are not stored in the laboratory or on the LIS.

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REFERRAL LABORATORIES

It may be necessary to refer specimens to referral /external laboratories, where:

- the referral laboratory provides a unique or unusual service
- the test requested is not routinely processed in the Biochemistry, Haematology or Microbiology/Molecular laboratories in NOHC (low volume test). Only clinically significant tests will be referred out of hours e.g. D-Dimer and Gentamycin. Needle stick serology is referred to NVRL by the medical scientist on call if the source is deemed as high risk by the doctor in charge or Infection Control Nurse.
- Confirmation of initial or unusual findings or address uncertainty of measurement.
- Backup service in the event of an unplanned interruption of the service at NOHC

For packaging instructions of laboratory specimens in compliance with ADR regulations Ref to AP-LAB-2


TURNAROUND TIMES (TATs)-Important note

TATs stated for tests referred to MMUH, indicate the Total Testing Process, from **time of receipt by MMUH laboratory**. Please allow additional time to refer the sample from the NOHC Laboratory. Routine times should be interpreted as minimum of 8 hours for all Mater referrals (same working day).

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MATER MISERICORDIAE UNIVERSITY HOSPITAL (MMUH) UNIVERSITY HOSPITAL (Refer to ED-LAB-7)


Clinical Chemistry and Diagnostic Endocrinology Tests

ACR (Albumin to Creatinine Ratio)	
Specimen:	Urine (Spot sample) analysed for "Microalbumin" and Creatinine.
Turnaround time:	3 Hours
Reference range:	0 – 2.5 mg/mmol Creatinine (Microalbuminuria: 2.5 – 25.0 mg/mmol Creatinine)
AFP (Alpha Feto Protein)	
Specimen:	4.9ml blood in WHITE tube (no anticoagulant).
Turnaround time:	1 working day
Reference range:	0 – 9 µg/L
Amylase	
Specimen:	4.9ml blood in LITHIUM HEPARIN GEL tube. Urine (spot sample) may also be tested.
Turnaround time:	Routine: 3 hours. Emergency: 1 hour
Reference range:	28 – 97 IU/L (Plasma); 0 – 470 IU/L (Urine)
Anti-TPO	
Specimen:	4.9ml blood in LITHIUM HEPARIN GEL tube.
Turnaround time:	1 – 3 days
Reference range:	<5.6 IU/ml
AST	
Specimen:	4.9ml blood in LITHIUM HEPARIN GEL tube.
Turnaround time:	3 Hours
Reference range:	19-42 IU/L
CA125	
Specimen:	4.9ml blood in LITHIUM HEPARIN GEL tube.
Turnaround time:	1 – 3 days
Reference range:	0 – 35 kU/L

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CA15.3	
Specimen:	4.9ml blood in LITHIUM HEPARIN GEL tube.
Turnaround time:	1 – 3 days
Reference range:	0 – 31.3 kU/L
CA19.9	
Specimen:	4.9ml blood in LITHIUM HEPARIN GEL tube.
Turnaround time:	1 – 3 days
Reference range:	2 – 23 U/ml
Catecholamines	
Specimen:	24-hour urine collection. (Acid container; pH <3) Referred to Beaumont Hospital. Catecholamines plus Metanephrines afford the optimal sensitivity in diagnosis of
Phaeochromocytoma.	
Turnaround time:	2 weeks
Therapeutic range:	See Report form
Carbamazepine	
Specimen:	4.9ml blood in BROWN SERUM GEL tube for clotted blood. Trough sample should be taken immediately
next	dose.
Turnaround time:	Routine: 3 hours. Emergency: 45 mins.
Therapeutic range:	4 – 12 mg/L
Carcinembryonic Antigen (CEA)	
Specimen:	4.9ml blood in LITHIUM HEPARIN GEL tube.
Turnaround time:	1 – 3 days
Reference range:	0 - 52 µg/L
Cholesterol	
Specimen:	4.9ml blood in LITHIUM HEPARIN GEL tube.

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Random or fasting sample. Reported as part of Lipid Screen (min 12 hour fast)	
Turnaround time:	3 hours
Reference range:	Desirable level: 3.1 – 5.0 mmol/L
CK (Creatine Kinase)	
Specimen:	4.9ml blood in LITHIUM HEPARIN GEL tube.
is preferred	Add-on allowed up to 8 hours if to establish baseline level, but if for query – MI a fresh (later) sample
Turnaround time:	Routine: 3 hours. Emergency: 1 hour
Reference range:	33 – 208 IU/L (Female), 44 – 272 (Male) Reference range for Caucasians: for Afro-Caribbean's twice these
Cortisol	
Specimen:	4.9ml blood in LITHIUM HEPARIN GEL tube. Note time and date of collection on tube to facilitate
appropriate	interpretation of result. Morning sample preferred.
Turnaround time:	Routine: 3 hours.
Reference range:	Morning: 150 – 455 nmol/L
Creatinine Clearance	
Specimen:	24-hour urine collection plus 4.9ml blood in LITHIUM HEPARIN GEL tube, the blood collected during
the 24 hour	collection period. Urine and plasma must be received by the laboratory together. NB: ensure only
one early	morning void included in 24-hour urine collection.
Turnaround time:	Calculation = $(UCr [\mu\text{mol/L}]/PCr[\mu\text{mol/L}]) \times (\text{Vol}[\text{mL}/24\text{hr}]/1440)$
Reference range:	3 hours 100 – 130 mL/min (Female), 110 – 150 mL/min (Male)
Digoxin	
Specimen:	4.9ml blood in BROWN SERUM GEL tube.
	Sample must be taken before dose or at least 6 hours post-dose.

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Turnaround time:	Routine: 1 working day. Emergency: 45 minutes.
Therapeutic range:	0.5 – 1.0 µg/L
Epanutin	See Phenytoin
Epilim	See Valproate
Free T4	
Specimen:	4.9ml blood in LITHIUM HEPARIN GEL tube. Included as part of Thyroid Function Test (TFT)
Turnaround time:	1 – 3 days
Reference range:	8-20 pmol/l
Free T3	
Specimen:	4.9ml blood in LITHIUM HEPARIN GEL tube. Included as part of Thyroid Function Test (TFT)
Turnaround time:	1 – 3 days
Reference range:	2.6-6.0 pmol/l
FSH	
Specimen:	4.9ml blood in LITHIUM HEPARIN GEL tube for clotted blood.
Turnaround time:	1 – 3 days
Reference range:	1.4 – 10.8 mIU/ml (Males) Female: Follicular phase: 3.0 – 8.1 IU/L. Mid-cycle peak: 4.9 – 16.7 IU/l, Luteal phase: 1.4 – 5.5 IU/l
Growth Hormone (GH)	
Specimen:	4.9ml blood in WHITE tube. (no anticoagulant)
Turnaround time:	2 weeks
Reference range:	Not applicable. Please evaluate GH response in relation to the relevant Dynamic Function Test protocol. As a guide to the initial investigation of hypoglycaemia, a rise to >5 µg/L, may be considered adequate.
HbA1c (Glycosylated Haemoglobin)	
Specimen:	2.7 ml=L PINKISH EDTA tube.
Turnaround time:	1 – 2 days

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Reference range:	IFCC: 20 – 42 mmol/mol; DCCT: 4.0 – 6.0%
HDL (HDL – Cholesterol)	
Specimen:	4.9ml blood in LITHIUM HEPARIN GEL tube. Included as part of lipid profile. Fasting 12 hours + is required
Turnaround time:	3 hours
Reference range:	> 1.5 mmol/L
Homocysteine	
Specimen:	2.7ml PINKISH EDTA tube. (Contact lab; must be spun and separated within 10 minutes of taking blood)
Vincent's Hospital.	Referred to Temple Street Hospital for Metabolic investigations, all other Homocysteine's for to St
Turnaround time:	6 weeks
Reference range:	See report form
LDH (Lactate Dehydrogenase)	
Specimen:	4.9ml blood in BROWN SERUM GEL tube for clotted blood.
Turnaround time:	Routine: 3 hours, Emergency: 45 minutes
Reference range:	120 – 220IU/L
LDL (LDL – Cholesterol)	
Specimen:	4.9ml blood in LITHIUM HEPARIN GEL tube. Fasting for 12 hours or more is required. LDL is a calculated parameter. $LDL = Chol - HDL - C - [TG/2.2]$ Tested as part of lipid profile/lipid screen.
Turnaround time:	2 hours
Reference range:	Desirable range: < 4.9 mmol/L (lower levels are recommended for secondary prevention)
LH	
Specimen:	4.9ml blood in LITHIUM HEPARIN GEL tube.
Turnaround time:	1 – 3 days
Reference range:	1.4 – 6.5 mIU/ml (Male) Female: Follicular phase: 1.8 – 11.8 IU/l, Mid-cycle peak: 7.6 – 89.1 IU/l, Luteal phase: 0.6 – 14.0 IU/l

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Lipid Screen / Lipid Profile	
Specimen:	4.9ml blood in LITHIUM HEPARIN GEL tube. Lipid profile comprises Total Cholesterol, Triglycerides, LDL-Cholesterol and HDL – Cholesterol. Fasting for 12 hours or more is required.
Turnaround time:	3 hours
Reference range:	See individual test
Lithium	
Specimen:	4.9ml blood in BROWN SERUM GEL tube for clotted blood. Trough sample should be taken immediately before
Turnaround time:	Routine: 3 days, Emergency: 3 hours
Therapeutic range:	0.6 – 1.2 mmol/L
Macroprolactin	
Specimen:	4.9ml blood in LITHIUM HEPARIN GEL tube. Automatically added to first time prolactin's >550 mIU/L
(Males),	>700 mIU/L (Females)
Turnaround time:	14 days
Reference range:	Males: 72 – 229 Miu/L, Females: 79 – 347 mIU/l
Osmolality	
Specimen:	4.9ml blood in LITHIUM HEPARIN GEL tube or urine (spot)
Turnaround time:	3 hours
Reference range:	Plasma: 285 – 295 mmol/Kg
investigation.	Assess urine osmolality relative to plasma osmolality and electrolytes and in the context of disease
	Calculated Osmolality = (1.87 x [Na]) + Glu + Urea + 9 (all in mmol/L)
Phenytoin	
Specimen:	4.9ml blood in BROWN SERUM GEL tube for clotted blood. Trough samples should be taken

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Turnaround time:	immediately before next dose
Therapeutic range:	Routine: 3 hours, Emergency: 45 minutes 10 – 20 mg/L
Prolactin	
Specimen:	4.9ml blood in LITHIUM HEPARIN GEL tube.
Turnaround time:	1 – 3 days
Reference range:	Males: 103 – 460 mIU/l; Females: 116 – 585 mIU/l. Macroprolactin screening is added if prolactin is >550 mIU/l in males, >700 mIU/l in females.
Progesterone	
Specimen:	4.9ml blood in LITHIUM HEPARIN GEL tube. Day 21 sample required (or 7 days before onset of menses if not a 28-day cycle)
Turnaround time:	1 – 3 days
Reference range:	Assuming correctly timed sample: <10nmol/l: ovulation unlikely, 10 – 30 nmol/l: equivocal, >60 nmol/l: ovulation likely
PSA (Prostate Specific Antigen)	
Specimen:	4.9ml blood in LITHIUM HEPARIN GEL tube. PSA test is available for diagnostic and monitoring purposes only, not for screening.
Turnaround time:	1 working day
Reference range:	0 – 4.0 µg/L
PTH (Parathyroid Hormone)	
Specimen:	2.7 ml blood in PINKISH EDTA tube.
Turnaround time:	1 – 3 days
Reference range:	1.89 – 7.61 pmol/l
Red Cell Folate	
Specimen:	2.7 ml blood in PINKISH EDTA FBC tube.
Turnaround time:	4 days
Reference range:	126 – 651 µg/l

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Tegretol	See Carbamazepine
Testosterone	
Specimen:	4.9ml blood in WHITE tube. (no anticoagulant)
Turnaround time:	10 days
Reference range:	Males: 6.3 – 24.7 nmol/l, Females: 0.7 – 1.9 nmol/l
Note:	SHBG is added to male testosterone >0.7, <10.0 nmol/l, and females if testosterone is >1.5 nmol/l. The free testosterone is also calculated in these cases. Refer to reports for reference ranges.
Theophylline	
Specimen:	4.9 mL blood in BROWN SERUM GEL tube for clotted blood. Dry tube (no anticoagulant) may be used.
Turnaround time:	Trough sample should be taken immediately before next dose
Therapeutic range:	Routine: 3 hours. Emergency: 1 hr.
	10 – 20 mg/L
TSH (Thyroid Stimulating Hormone)	
Specimen:	4.9 mL blood in LITHIUM HEPARIN GEL tube. Included as part of Thyroid Function Test (TFT)
Turnaround time:	1 to 3 days
Reference range:	0.35-4.94 mIU/l
Thyroid Function Test	
Specimen:	4.9 mL blood in LITHIUM HEPARIN GEL tube. Includes fT4 and TSH
Turnaround time:	1 to 3 days
Reference range:	See individual tests
Troponin I	
Specimen:	4.9mL LITHIUM HEPARIN tube.
Turnaround time:	Routine: 3 hours. Emergency: 1 hour
Reference range:	16 ng/L (F), 34 ng/L (M)

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MI should not be ruled out on the basis of a single negative early Troponin I value post onset of pain.

Urate (Uric Acid)

Specimen: 4.9 mL blood in LITHIUM HEPARIN GEL tube. Urine may also be tested; 24hour urine collected in plain container if delivered fresh to lab.

Turnaround time: Routine: 3 hours. Emergency: 1 hour

Reference range: Plasma: 177-465 µmol/L. Urine: 1.2 – 4.0 mmol/L /24hrs

Valproate (Epilim)

Specimen: 4.9 mL blood in BROWN SERUM GEL tube for clotted blood. Dry tube (no anticoagulant) may be used. Trough

sample should be taken immediately before next dose

Turnaround time: Routine: 3 hours. Emergency 1 hour

Vitamin D

Specimen: 4.9 mL blood in WHITE tube (no anticoagulant). Must be a separate specimen

Turnaround time: 2 weeks

Reference range: Desirable range greater than 50 nmol/L

VMA **VMA is now correctly called HMMA. See Catecholamines. HMMA (VMA) only measured if Catecholamines are abnormal**

Haematology

Blood Film

Specimen: 2.7 ml blood in EDTA tube (Pink Cap FBC tube).


Blood film will be examined, if requested, with relevant clinical information or if indicated by the F.B.C. In the presence of a normal F.B.C., there are few indications for routine film examination, e.g. possible infectious mononucleosis, malaria. Film must be made within 12 hrs. of venipuncture

Turnaround time: 96 hours

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Reference range:	See report form
*** Note min 96 hrs (4 working days and max 7 days for referred Blood films as Consultant Haematologist will review on site at weekly meetings)	
D-Dimers (DD)	
Specimen:	3ml Coagulation tube (green cap) Blood exactly aspirated to the mark and gently mixed. Specimen must be assayed within 4 hours of venipuncture
Turnaround time:	3 hours
Reference range:	0 - 0.5mg/L
Flow Cytometry	
Specimen:	2.7 ml blood in EDTA PINK FBC tube or RPMI Heparinized sample for bone marrow available in laboratory. Must be arranged in advance with Haematology Medical "team" and Haematology Laboratory.
Turnaround time:	7 Days
Reference range:	See Report Form
Haemoglobin Electrophoresis	
Specimen:	2.7 ml blood in EDTA tube (Pink Cap/FBC tube). Analysis carried out using alkaline electrophoresis and HPCL to quantify HbA2 and HbF and to identify/quantify haemoglobin variants. Acid electrophoresis is used to confirm/identify rarer variant haemoglobins
Turnaround time:	10 days
Reference range:	Adult HbA2 1.5 - 3.4 % HbF < 1.0%
Lupus Anticoagulant	
Specimen:	Three x 3ml Coagulation tubes (green cap). Blood exactly aspirated to the mark and gently mixed. Tests performed include PT, APTT, PTTLA, DVV, and Lupus confirmatory tests.
Turnaround time:	14 days
Reference range:	PTTLA Please refer to reference range stated with result DVV Please refer to reference range stated with result
Sickle Cell Screen	

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Specimen:	2.7 ml blood in EDTA tube (Pink Cap FBC tube).
Turnaround time:	24 hours
Reference range:	Positive or negative result BUT all results must be verified on Haemoglobin electrophoresis
Thrombophilia Screen Please Contact Laboratory for further advise (RESTRICTED TESTING)	
Sample received for Heritable Thrombophilia testing must meet the guideline as detailed on MMUH website – see attached link. http://www.mater.ie/healthcare-professionals/gp-referrals/Guideline_for_heritable_thrombophilia_testing.pdf	
Requests not meeting the guideline will generate one of the following comments: “No clinical details are provided, /Incomplete clinical details are provided/The request falls outside MMUH guideline for testing.	
These samples will not be processed but will be held in storage in the Haematology laboratory until additional clinical information had been received. In the event of no communication the samples will discarded after eight weeks.	
Specimen:	6 x 3ml Coagulation tube (green cap) or 2 x 10ml coagulation tube (green cap) and 1 x 2.7 ml blood in EDTA tube (Pink cap FBC tube). Tests performed include PT, APTT, Fibrinogen, Lupus Anticoagulant, Protein C, Free Protein S, Antithrombin, Activated Protein C Resistance, Factor V Leiden. Patients should be 2-3 weeks post thrombotic event before testing, samples should be sent to the lab as soon as possible after phlebotomy. The following tests will not be performed if the patient is on VKA: PC, PS, LA. The following tests will not be performed if the patient is on heparin: LA, APCRV (and also FVL), Antithrombin.
I.M.S (Infectious Mononucleosis) Screen	
Specimen:	2.7ml blood in EDTA tube (Pink Cap FBC tube)
Turnaround time:	3 hours
Reference range:	Positive or negative result

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Immunology

ANCA antibodies (Immuno fluorescence test for ANCA Pattern)	
Specimen:	4.9 ml in BROWN Serum Gel tube for clotted blood
Turnaround time:	3 days routine samples, same day for phoned urgent samples if received before 12 pm.
Reference range:	Negative is normal.
Antibody Screen	
Specimen:	4.9 ml in BROWN SERUM GEL tube for clotted blood
Turnaround time:	7 days
Reference range:	Negative is normal The antibody screen includes parietal cell antibody (PCA), mitochondrial antibody (AMA), smooth muscle antibody (SMA), which are tested on rat tissue. If the PCA is positive, the sample will automatically be tested for thyroid peroxidase antibody. If the AMA is positive, the sample will automatically be tested for M2 (PBC) and LKM antibodies.
Additional Information:	Positive samples will not be repeated within 3 months
BJP Identification	
Specimen:	Urine samples analysed by electrophoresis and immunofixation for presence of BJP.
Turnaround time:	14 days
Reference range:	No BJP detected is normal.
CCP (cyclic citrullinated peptide) antibody	
Specimen:	4.9 ml in BROWN SERUM GEL tube for clotted blood
Turnaround time:	4 days
Reference range:	Less than 11 U/mL
Complement (C3 & C4)	
Specimen:	4.9 ml in BROWN SERUM GEL tube for clotted blood
Turnaround time:	2 days
Reference range:	Adult C3: 0.75 – 1.88 g/L. C4: 0.14 – 0.61 g/L
dsDNA antibody (Immunoassay)	

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Specimen:	4.9 ml in BROWN SERUM GEL tube for clotted blood
Turnaround time:	21 days
Reference range:	Less than 15 IU/mL If the result is greater than 10 IU/mL, DNA CL will automatically be requested (unless it was previously DNACL+)
ENA antibodies (Ro, La, RNP, Sm, Scl-70 & Jo-1)	
Specimen:	4.9 ml in BROWN SERUM GEL tube for clotted blood
Turnaround time:	21 days
Reference range:	If the antinuclear antibody was ANA +, it is automatically tested for ENA antibodies. The sample is initially screened for ENA antibodies, if this is positive, the sample is automatically tested for the individual antibodies (ENAC). ENA Screen ratio <0.7 is negative. Samples with levels >0.7 will be tested for specific ENA antibodies
Additional Information:	Positive samples will not be repeated within 3 months
Endomysial (IgA) antibody	
Specimen:	4.9 ml in BROWN SERUM GEL tube for clotted blood
Turnaround time:	14 days
Reference range:	Negative is normal.
Additional Information:	EMA is only used to confirm TTG positive samples. Previously positive samples are not repeated for EMA. The tTG antibody assay is used to monitor patients.
IgE	
Specimen:	4.9 ml in BROWN SERUM GEL tube for clotted blood
Turnaround time:	8 days
Reference range:	0 - 3 years old: less than 56 kU/L 3 - 7 years old: 56-110 kU/L 8 - 10 years old: 124-148 kU/L Greater than 10 years old: Less than 114 kU/L
Immunoglobulins IgG, IgA & IgM	
Specimen:	4.9 ml in BROWN SERUM GEL tube for clotted blood Requests for Immunoglobulins are also assayed for Protein Electrophoresis and vice versa.
Turnaround time:	3 days

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Reference range:	Adults IgG: 7.00 –16.00 g/L IgA: 0.70 – 4.00 g/L IgM: 0.4 – 2.30 g/L The results are not released until compared with Protein Electrophoresis results. If there is a band present, sample is immunofixed.
Nuclear Antibody (ANA)	
Specimen:	4.9 ml in BROWN SERUM GEL tube for clotted blood
Turnaround time:	7 days
Additional information:	Negative is normal ANA is tested on Hep2 cells to detect and identify anti nuclear antibody (centromere, homogeneous, nucleolar and Speckled patterns). If the ANA is greater than a weak positive, it will automatically be tested for dsDNA and ENA (extractable nuclear antigen, which includes anti Ro, La, RNP, Sm, Jo-1 & Scl-70) antibodies. Positive samples will not be repeated within 3 months
Paraprotein identification/Protein Electrophoresis	
Specimen:	4.9 ml in BROWN SERUM GEL tube for clotted blood Requests for Protein Electrophoresis (SPEP) are also automatically assayed for serum Immunoglobulins. If a possible Paraprotein band is detected then, the sample is automatically reflexed for immunofixation, unless a paraprotein previously typed and identified.
Turnaround time:	3 days
Reference range:	No paraprotein detected is normal. Report comments added on an individual case basis
Parietal Cell Antibody	
Specimen:	See antibody screen
Rheumatoid Factor	
Specimen:	4.9 ml in BROWN SERUM GEL tube for clotted blood Fluids cannot be tested.
Turnaround time:	2 days

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Reference range:	Less than 20 IU/mL (nephelometry)
Smooth Muscle Antibody	
Specimen:	See antibody screen
Thyroid Antibodies	Please refer to Clinical Chemistry and Diagnostic Endocrinology
Tissue Transglutaminase antibody	
Specimen:	4.9 ml in BROWN SERUM GEL tube for clotted blood
Turnaround time:	2 days
Reference range:	Less than 3 U/mL
Additional information:	The IgA level is automatically checked, as IgA deficiency causes false negative results for tTG IgA antibody. If the patient is IgA deficient; the tTG IgG antibody assay is completed instead of the tTG IgA antibody. The endomysial IgA antibody assay is used to confirm positive tTG IgA samples, if they were not previously positive.
Cardiolipin IgG & IgM antibody	
Specimen	4.9 ml in BROWN SERUM GEL tube for clotted blood
Turnaround Time:	8 days
Reference Range:	Less than 10 U/ml


Other Possible Referral Immunology Tests include

Acetyl choline esterase antibody John Radcliffe Hospital Headington Oxford UK	
Specimen:	4.9ml in Brown Serum Gel tube for clotted sample
Turnaround Time:	14 days
Referral Laboratory	
Tryptase Beaumont Hospital	
Specimen Type:	4.9ml serum sample x 3 (timed at 0,4-6 hrs and >24 hours post event)
35 days	

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Microbiology


Routine Bacteriology

Clostridium difficile toxin (Faeces)	
Specimen	Approximately 5-6 grams faeces are sufficient for Clostridium difficile toxin testing. This should be collected into a Sterile leak proof container (MSU). Transport to laboratory as soon as possible. Turnaround time Same day of receipt Monday-Friday
Storage	Specimens are stored at 4oC until they are processed.
Additional information	Clostridium difficile toxin testing will only be performed on diarrhoeal samples. Repeat testing will not be performed for 28 days on patients who are Clostridium difficile toxin positive. Sample must be received in the Cherry Orchard laboratory for processing by 9am, and must be delivered directly to the NOHC Microbiology Laboratory by 08.00 for same day results
CSF (Cerebrospinal Fluid)	
Specimen	CSF is normally collected sequentially into 3 or more separate disposable sterile screw-capped containers, which should be numbered sequentially 1, 2, 3 etc. A minimum volume of 1 ml in each container is required. Please prioritise tests and ensure there is sufficient sample if a large number of tests are ordered.
Turnaround time	Processed on receipt. Microscopy report: <2hours Final negative report: 48hours Final culture positive report: 48-72hours
Special precautions	All positive results are telephoned to a member of attending team by Microbiology medical staff.
Additional information	Samples of CSF must be brought to the laboratory and handed directly to a member of staff. Oligoclonal bands: When oligoclonal bands are requested, a clotted blood sample (white cap or brown cap) must also be taken. These are referred to a laboratory in Sheffield. CSF samples not accompanied by serum will not be referred.

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Sputum	
Specimen	a) Sputum-expectorated b) endotracheal tube specimen (induced sputum) A minimum volume of 1 ml sample in a sterile screwcap container e.g. MSU container Turnaround Times: Routine final report: 2-4 days. Transplant patients: 5-7 days Special Precautions: Salivary or mucosal specimens are unsuitable for culture and will not be processed. Exceptions are samples from ITU/HDU/CTH DU and HLTW Where the patient has difficulty producing a specimen, postural drainage and physiotherapy may be required Samples should reach the laboratory within 4 hours of collection to avoid overgrowth with Gram-negative bacilli. Also <i>H. influenzae</i> and <i>S. pneumoniae</i> may not survive beyond this time Results from specimens not received in the laboratory on the same day, as collection should be interpreted with care. Clinical information e.g. cystic fibrosis is essential Storage: Specimens are stored at 4oC until they are processed. Additional information: Early morning fresh expectorated sputum is recommended for <i>Mycobacterium tuberculosis</i> . See section- Mycobacteriology. If required, a separate sample should be taken for cytology and sent to the Histology laboratory.


Mycobacteriology

Tuberculosis	
Specimen	Early morning freshly expectorated sputum is recommended for <i>Mycobacterium tuberculosis</i> <ul style="list-style-type: none"> • Sputum (3 consecutive mornings): 5ml per sample • EMU- not appropriate for the diagnosis of respiratory TB. Please contact a Clinical Microbiologist (4634) before ordering this test. Early morning urine on 3 consecutive days; minimum of 50ml of each collection. Note: appropriate containers are available from biochemistry

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- Aspirate brushing
- Bronchoalveolar Lavage (BAL) Bronchial washing - 5 ml
- CSF: minimum volume 1ml
- Body fluids, aspirates, pus: minimum 1ml
- Gastric Lavage: collected early morning before breakfast. Used for children where a sputum cannot be obtained, preferably 5ml per sample
- Skin or Tissue biopsies. Note: do not add formalin
- Blood/bone marrow: (specific culture bottles available from Microbiology laboratory) Add 3mls aseptically into the culture bottle and return promptly to the microbiology laboratory in the transport container provided.
- Post mortem specimens

Samples must be taken into a sterile screwcap container e.g. MSU container and sealed in a plastic biohazard bag

Turnaround time

Auramine Stains:

Routine: Same day if received by 8am Monday to Friday

Urgent: Direct stains can be carried out on samples received before 22.00 using special requisition form available in laboratory.

Culture:

7 weeks (most cultures become positive within the first four weeks of incubation, however if the strain is slow-growing or if there are scanty numbers of organisms present in the specimen prolonged incubation may be required) Note: Positive microscopy and positive cultures are phoned immediately to the requesting source


Sensitivity Testing:

The first isolate from each positive patient is identified and the culture is sent to a reference laboratory for susceptibility testing. Results are normally available within three weeks of the date on which the culture is dispatched.

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Storage:	<p><u>Molecular Tests:</u> Direct detection of Mycobacterium tuberculosis complex in a clinical sample: available on request, 7day turnaround time</p> <p><u>Identification of Mycobacterial cultures to species level:</u> Routinely carried out on the first isolate from each patient, results within 7 working days</p> <p><u>Molecular detection of resistance to rifampicin and isoniazid:</u> Routinely carried out on all first isolates of <i>Mycobacterium tuberculosis</i> complex, results available within 7 working days.</p> <p>Specimens are stored at 4oC until they are processed.</p>
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Parasitology


Faeces: Ova, Cysts and Parasites – only by request of Consultant Microbiologist	
Specimen:	Approximately 5 – 6 grams collected into a sterile leakproof container (MSU)
Turnaround Time:	3-5 days
Special Precautions	Three examinations spaced 2-3 days apart are recommended for optimal recovery, as shedding of ova and cysts tends to be intermittent. Full clinical details are Essential eg foreign travel, immunocompromised patients. Samples received without appropriate clinical information will not be processed.
Stoage:	Specimens are stored at 4oC until they are processed

Quantiferon Testing – By prior arrangement only	
Specimen:	4 blood samples collected into 4 Quantiferon TB Gold collection Tubes These must be ordered in advance and collected from the Microbiology Laboratory. Date and time of specimen collection must be recorded on each tube. Inoculated tubes must not be refrigerated.
Turnaround time:	7-10 days
Special precautions/ Additional Information:	

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Specimens are accepted Monday to Thursday up to 4pm only – specimens will not be accepted on Fridays or after hours. Specimen tubes must be filled to the black line on each tube – DO NOT OVERFILL Tubes must be thoroughly mixed and delivered promptly to the Microbiology Laboratory – do NOT refrigerate

Fluids and Aspirates: Total cell count and Differential Cell counts	
Specimen:	A minimum volume of 1ml is required for all joint/synovial fluids. Fluids are collected aseptically according to local protocols into a sterile leak-proof container. Fluids may also be sent in Blood Culture bottles.
Turnaround time:	Total and Differential cell count: same day. Final report 7 days except on Bank Holidays +24 Hours
Special precautions:	Fluids for total and differential cell counts must be collected into an additional EDTA PINKISH Tube. Specimens for cell count not received in EDTA will not be processed.
Storage:	Specimens are stored at 4oC until they are processed.


Therapeutic Drug Monitoring

Antibiotic Assays	Amikacin, Gentamycin,
Specimen:	4.9 ml in ORANGE LITHIUM HEPARIN GEL tube.
<u>When to monitor</u>	<ul style="list-style-type: none"> In serious sepsis within the first 72 hours of commencing therapy to ensure adequate peak blood levels and to ensure that accumulation of the dose does not occur. Thereafter at three day, intervals if renal function is normal Daily in patients with impaired renal function in order to ensure that accumulation of the drug is not taking place

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Therapeutic ranges

- Daily in patients on renal dialysis, on prolonged treatment (>10 days) and in patients with changes in renal function during aminoglycoside therapy.
- If an expected therapeutic response does not occur
- In patients with gross obesity, in whom levels tend to be high when dosed according to total body weight

ANTIBIOTIC	TROUGH	
Amikacin	<5.0mg/L	
Gentamycin (once daily dose)	<1.0 mg/L	

Additional Information:

Peak levels are no longer indicated


NATIONAL CENTRE FOR HEREDITARY COAGULATION DISORDERS - ST JAMES' HOSPITAL

Factor V Leiden (FVL)	
Specimen:	1 x 3ml Purple EDTA and 2 x 3ml Light Blue Tri-Sodium Citrate. Samples must be received by Coag Lab, NCHCD by 4pm Mon-Fri. Requests for Factor V Leiden must be accompanied by either sample for APCR analysis or an APCR result from an external source.
Turnaround time:	1 month
Extrinsic Factor Assay (Factor II, Factor V, Factor VII, Factor X, Factor 2, Factor 5, Factor 7, Factor 10)	
Specimen:	6 x 3 ml LIGHT BLUE tri—sodium Citrate
Turnaround time:	7 days
Reference range:	Adult (>18 years) Factor II = 0.72 – 1.31 IU/mL Factor V = 0.63 – 1.33 IU/mL Factor VII = 0.51 – 1.54 IU/mL

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Factor X = 0.64 – 1.50 IU/mL	
Intrinsic Factor Assay (Factor VII, Factor IX, Factor XI, Factor XII, Factor 8, Factor 9, Factor 11, Factor 12)	
Specimen:	6 x 3 ml LIGHT BLUE tri—sodium Citrate
Turnaround time:	7 days
Reference range:	Adult (>18 years)
	Factor VIII = 0.60 – 1.36 IU/mL
	Factor IX = 0.80 – 1.47 IU/mL
	Factor XI = 0.72 – 1.52 IU/mL
	Factor XII = 0.52 – 1.64 IU/mL
VWD Screen (Von Willebrand Screen)	
Specimen:	4 x 3 ml LIGHT BLUE tri—sodium Citrate
Turnaround time:	3 weeks (including multimers 6 weeks)
Reference range:	Adult (>18 years)
	Factor VIII = 0.60 - 1.36 IU/mL
	VW Antigen = 0.49 - 1.73 IU/mL
	VW Ristocetin Co-Factor = 0.55 - 1.56 IU/mL
	VW Collagen Binding = 0.50 - 1.50 IU/mL
	VW Multimers = Qualitative result

NATIONAL VIRUS REFERENCE LABORATORY


Bone Bank Serology	
Specimen:	5.5 ml in WHITE SERUM tube. Stand upright once drawn
Turnaround time:	2 weeks

NATIONAL VIRUS REFERENCE LABORATORY – SEROLOGY

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Needle Stick Serology	
Specimen:	5.5 ml in WHITE SERUM tube. Stand upright once drawn
Turnaround time:	Routine: 3 working days, Urgent: 1 hour
Post Vaccination Titre (Hep B Titre)	
Specimen:	5.5 ml in WHITE SERUM tube. Stand upright once drawn
Turnaround time:	24 hours
Reference range:	>10 mIU/mL

NATIONAL VIRUS REFERENCE LABORATORY – MOLECULAR VIROLOGY

Norovirus (Molecular Qualitative)	
Specimen:	Stool sample in a sterile universal container
Turnaround time:	5 working days
Influenza (PCR Qualitative)	
Specimen:	BAL, NPA, nose and throat swab
Turnaround time:	Same day if received before 10am (in season) 5 working days (out of season)


PUBLIC HEALTH LABORATORY SERVICES – CHERRY ORCHARD

Faeces (Culture and Sensitivity)	
Specimen:	Stool sample in a sterile universal container
Turnaround time:	Negative cultures 48 hours
VTEC detection by PCR	

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Specimen:	Stool sample in a sterile universal container
Turnaround time:	1 working day

IRISH BLOOD TRANSFUSION SERVICES

Antibody Investigations	
Specimen:	2 x 7.5ml EDTA RED transfusion tube.
Turnaround time:	5-10 working days
HLAB27	
Specimen:	7.5 ml EDTA tube. (FULL)
Turnaround time:	10 working days
D Variant Typing	
Specimen:	7.5ml EDTA RED transfusion tube.
Turnaround time:	10 working days

CHARING CROSS HOSPITAL


Fluid samples for Chromium/Cobalt (ASR)	
Specimen. :	Fluid sample must be kept in fridge until delivery. Samples delivered at ambient temperature through Eurofins
Address :	Ground floor Medical Oncology Block, Charing Cross Hospital, Fulham Palace Road, London, W68RF, UK
Turnaround time:	4 weeks
Reference range:	N/A

Berkshire and Surrey Hospital

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Tissue Samples for Chromium/Cobalt (ASR)

Specimen.:	Tissue samples kept in fridge until delivery. Samples delivered on dry ice through Eurofins
Address:	Berkshire and Surrey pathology services, SAS trace element laboratory, 15 Frederick Sanger road, Surrey Research Park, GU27YD, UK
Turnaround time:	4 weeks
Reference range:	N/A

Southampton General Hospital


Blood Samples for ASR and Titanium

Specimen.:	7.5ml METAL FREE LITHIUM HEPARIN ORANGE tube. Use metal free needle. Samples kept frozen and delivered at ambient temperature through Eurofins
Address:	SAS trace element unit, University Hospital Southampton, NHS Foundation trust, Southampton General Hospital, Tremona road, Southampton, S0166YD, UK.
Turnaround time:	4 weeks
Reference range:	Titanium 0 - 40 nmol/L Chromium 0 – 20 nmol/L Cobalt 0 - 17 nmol/L

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QUALITY POLICY

The Pathology Department of the National Orthopaedic Hospital Cappagh is committed to providing the highest quality diagnostic and consultative services for all its users.

In order to achieve this objective, it is the policy of the Department to implement and maintain an effective Quality Management System, which will include:

- The setting of quality objectives and plans at its Annual Management Review
- Integration of the Department's procedures, processes and resources.
- Commitment to the health, safety and welfare of patients, staff, students and visitors.
- Maintenance of professional practice including the safeguarding of patient information.
- Familiarising personnel with the content of this Quality Manual, Quality Policy and all procedures relevant to their work to ensure user satisfaction.


The Quality Management system will comply with the standards set by ISO 15189, AML-BB, S.I. No. 360 of 2005 European Communities (Quality and Safety of Human Blood and Blood Components) Regulations 2005 and S.I. No. 158 of 2006 European Communities (Quality and Safety of Human Tissues and Cells), EU Directive 2002/98/EC (setting standards of quality and safety for the collection, testing, processing, storage and distribution of human blood and blood components) and EU Directive 2004/23/EC (setting standards of quality and safety for the donation, procurement, testing, processing, preservation, storage and distribution of human tissues and cells) for the services and tests defined in the Quality Manual and is committed to:

- Staff recruitment, training and development at all levels to provide an effective and efficient service to its users.
- Providing and managing resources to ensure that all examinations are processed to produce the highest quality results possible.
- Reporting results in ways, which are timely, confidential, accurate and are supported by clinical advice and interpretation when required,
- Ensuring that laboratory examinations are fit for intended use

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National Orthopaedic Hospital Cappagh, Finglas, Dublin 11.		PATHOLOGY LABORATORY USER MANUAL

- Implementation of Internal Quality Control, External Quality Assessment, Audit and Assessment of User Satisfaction to continuously improve the quality of the service
- Compliance with relevant environmental legislation.
- Adherence to appropriate technical and professional standards.

Management and staff are committed to creating a quality culture within the Department by continuously improving our services based on the results of performance through data review, internal quality audits, equipment maintenance, Quality Control programmes and the assessment of users needs.

PROBLEMS, COMPLAINTS & SERVICE IMPROVEMENTS

If any problems are encountered with the service or any matter for complaint arises, please contact:

- Laboratory Manager 01- 814 0386
- Quality Systems Manager 01- 814 0456

In addition, suggestions for improvement of the service are always welcome.

CLINICAL INTERPRETATION

During Routine hours please contact:

Biochemistry:	Dr. Graham Lee	01 8034983
Haematology:	Prof. Peter O’Gorman	01 803 2438
	Dr. Khalil Alnajjar	01 850 0977
NOHC Microbiology:	Dr. Deirdre Brady	01 8545067
SSC Microbiology:	Dr. Edmond Smyth	086 252 4473
NOHC Histology:	Prof. Conor O’Keane	01 8032265
Blood Transfusion:	Prof. Peter O’Gorman	01 803 2438
	Dr. Khalil Alnajjar	01 850 0977

Out of hours:

Please contact the on call Medical Scientist, who will contact the required consultant (RF-PMG-33)

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