<u>Dept</u>	Discription of Changes from Ver. 3.12 to 3.13 GUH LAB USER GUIDE	Page No. Ver. 3.12	Page No. Ver. 3.13
General	Changed: USE OF THE LABORATORY from page 12 to 13 Changed: PHLEBOTOMY SERVICE from page 18 to 19 Changed: TRANSPORT OF SPECIMENS TO THE LABORATORY from page 20 to 21 Changed: REPORTING RESULTS from page 23 to 24 Changed: BLOOD AND TISSUE ESTABLISHMENT from page 25 to 26 Changed: DIVISION OF ANATOMIC PATHOLOGY from page 43 to 42 Changed: MORTUARY SERVICES-AUTOPSY from page 53 to 52 Changed: IMMUNOLOGY DEPARTMENT (SUPRAREGIONAL SERVICE) from page 57 to 56 Changed: HAEMATOLOGY LABORATORY from page 69 to 70 Changed: MEDICAL MICROBIOLOGY DEPARTMENT from page 73 to 74	1	1
General	Changed: ALPHABETICAL TEST DIRECTORY from page 91 to 90 Updated header spacing	2	2
General	Updated footer to current version, updated dates of authorisation and review Removed: " Martina Doheny ", added: "Prof. Murray and Dr. Phelan	2	2
General	The Laboratory Medicine Service removed: "Laboratory Manager: Ms. Martina Doheny Email: martina.doheny@hse.ie" Added: "Associate Clinical Director for the Laboratories: Dr. Sine Phelan Email: Sine.Phelan@hse.ie"	3	3
General	General Information Gentact Information Updated laboratory manager Removed: "Ms. Martina Doheny", Added: "Not appointed"	10	10
General	General Information Gontact Information Specimen reception: Removed: "Ms. Bridie O'Donnell", added: "Ms. Karen Mullins Central Reception Manager"	10	10
General	3. Use of the Laboratory 3.1 Register of Users: Removed: "Liz Neville, liz.neville@hse.ie", Added: "Pearse Timothy, Pearse.Timothy@hse.ie"	12	13
вю	2. General Information 2.3 Contact Information Clinical Biochemistry: removed Ms-Michelle-Finnegan Michellee-finnegan@hse-ie, added Ms. Martina Doheny martina.doheny@hse-ie; removed: Mr. Liam Blake liamM.blake@hse-ie, Phone Ext: 2709, added: Dr. Janice Reeve janice.reeve@hse-ie Phone Ext: 8752, added: Ms. Karen Heverin Principal Clinical Biochemist karen.heverin@hse-ie Phone Ext: 8752 added: Dr. Verena Gounden Consultant Chemical Pathologist Verena.Gounden@hse-ie Phone ext. 8200	6	6
BIO	Clinical Biochemistry Department 4.4 Biochemistry Tests Replaced: "Summary of Blood Specimen volume requirements" with "Summary of Request Forms and Blood Specimen including Volume Requirements"	38	38
вю	8. Clinical Biochemistry Department 8.4 Biochemistry Tests Added: "A single request form may be utilised for General Biochemistry, Glucose and HbA1c requesting, ensuring the appropriate number of specimens are provided. Specialist tests performed in-house and special assays referred to external laboratories require individual request forms and separate specimens. Once collected, submit the entire specimen to the laboratory with the appropriate request form."	38	38
BIO	8. Clinical Biochemistry Department 8. 6 GP Specimens Updated: The target turnaround time for routine GP requests is 2 4 working days.	39	39
ВІО	8. Clinical Biochemistry Department 8.13 Near Patient Testing (NPT) added: "ketone monitoring, "and "Ketone meters are available in critical care and diabetic outpatient services."	40	40
вю	8. Clinical Biochemistry Department 8.13 Near Patient Testing (NPT) The development of an integrated laboratory-connected and managed NPT service for critical care analysers, glucose meters, added: ", ketone meters " added: "Feedback The clinical biochemistry department welcomes feedback from clinical users and patients, both positive and negative. All feedback is communicated to management and staff to allow us to shape our processes. Complaints are recorded in our quality management system and fully investigated, with feedback on root cause and actions required, where relevant, to the complainant. " added: "Patient Consent For most routine laboratory procedures, consent can be inferred when the patient willingly submits to the sample collecting procedure, for example, venepuncture. Any further patient consent requirements are outlined in the alphabetical test directory contained in section 16 of this document. Patient consent remains the responsibility of the requesting clinician and the laboratory cannot accept responsibility for referral laboratory rejection of requests due to patient consent being unavailable."	41	41
BIO	16. Alphabetical Test Directory added: Anti-Mullerian Hormone (AMH) Laboratory: Clinical Biochemistry Specimen: 7.0 mL blood in a plain gel tube Turnaround: Priority: 1 working day. Routine: 4 working days Ref. Range: On report form	99	99
ВІО	16. Alphabetical Test Directory Calcitonin Specimen: added "on ice"	112	113
BIO	Calcium-ionised Turnaround: replaced: "15 mins" with "15 minutes"	112	113
BIO	To historical Test Directory Cerebrospinal Fluid - Lactate Specimen: replaced: "300 μL." with "300 μL"	115	116
BIO	Dihydropyrimidine Dehydrogenase (DPD) Activity Specimen: removed: " , and a urine specimen "	127	128

	,		
BIO	16. Alphabetical Test Directory Glucose: removed: Fluoride Oxalate blood , added: Vacuette FC mix tube NaF/Citrate/EDTA	137	138
вю	16. Alphabetical Test Directory removed: "Interleukin 6 Laboratory: Clinical Biochemistry. Specimen: 7.0 mL blood in a plain gel tube Comment: Specimen must be received in the laboratory on the day of venepuncture. Turnaround: Urgent: 2 hours. Priority: 3hours. Routine: 4 working days Ref. Range: See report form"	147	149
BIO	16. Alphabetical Test Directory Lactate Turnaround: replaced: "15 mins" with "15 minutes"	149	151
BIO	16. Alphabetical Test Directory Metanephrines (Metanephrine/Normetanphrine/3-methoxytyramine - Plasma) Specimen: replaced: "30 mins " with "30 minutes"	154	156
BIO	16. Alphabetical Test Directory Porphyrin Screen Specimen: replaced: "24 hour" with "spot" urine collection Comment: added: "St. James's Hospital Porphyrin Request Form must be completed, available in GUH Useful Resources"	165	166
ВІО	16. Alphabetical Test Directory removed: "Procalcitonin Laboratory: Clinical Biochemistry. Specimen: 7.0 mL blood in a plain gel tube, received in the laboratory within 6 hours of venepuncture. Turnaround: Urgent: 2 hours. Priority: 3hours. Routine: 4 working days Ref. Range: See report form"	166	167
BIO	16. Alphabetical Test Directory SHBG removed: "Comment: Female — only analysed where testosterone >1.2nmol/L."	172	173
DAP	Added: "Special counselling may be needed for examination results with serious implications for the patient. It is the responsibility of the test requester to ensure that examination results with serious implications for the patient are not communicated to the patient without the opportunity for acceptable counselling."	2	2
DAP	2. General Information 2.3 Contact Information Anatomic Pathology: Histopathology, Cytopathology and Molecular Pathology added: "Dr Aliaa Shalaby Consultant Pathologist Aliaa.Shalaby@mailn.hse.ie Phone Ext: 3445" Added: "Dr Kevin Culligan Consultant Pathologist Kevin.Culligan@hse.ie Phone Ext: 3853" removed: Terri Muldoon, Chief Medical Scientist, Terri.muldoon@hse.ie; added: "Loretta Lydon, Chief Medical Scientist, Loretta.Lydon@hse.ie"	8	8
DAP	General Information Suboratory Opening Hours Anatomic Pathology: Deadline for sample in Lab: changed from 16:30 to 16:00 Mon-Fri	11	11
DAP	2. General Information 2.7 Complaints: Added: "Feedback, including complaints is open to patients and laboratory users throughout the "Your Service Your Say" mechanism accessible on the HSE Saolta website. Complaints are processed in accordance with the HSE policy- Your Service Your Say – management of service user feedback for comments, compliments and complaints- publicly accessible on the HSE site."	11	12
DAP	3. Use of the Laboratory 3.2 Requests to the Laboratory Histopathology requirement: Added: "Where the clinician is submitting slides to the DAP for analysis that the number of slides being submitted should be recorded on the request form."	14	15
DAP	3. Use of the Laboratory 3. 2 Requests to the Laboratory Histopathology requirement: Added: "• For fixed specimens, ensure the specimen container selected is large enough to allow the specimen to be immersed in at least twice its own volume of buffered formalin." Formatted to bullet point below: "• The specimen site must be indicated and detailed on the request form and on the container. • In the case of multipart container submission on a case each part must be clearly identified as to the site and nature of the specimen. The detail on the request form and the specimen container must match. • The lid must be securely closed to prevent spillage • Radioactive specimens: The Request Form and specimen containers must have a radiation label. When a radioactive specimen is being sent information on the radiation dose should be given. The specimen should be delivered to the dedicated lab room for radioactive specimens. It should be placed behind the lead shield, and the lab staff informed of its presence there. • The Request Form and specimen containers must indicate if specimen is high risk (eg TB, COVID-19, HIV or Hepatitis). • The Colorectal Programme specimen request form must include the NCSS COR number. SHARPS containers must not be used as specimen containers. • Note: It is not possible or safe at the moment of receipt of the specimen(s) in the Division of Anatomic Pathology to check each pot for the presence of a specimen. Therefore acceptance of a test request by the DAP staff is not confirmation that the described specimen is present in the container, but rather that the form details and the container details, and where applicable the sign off book details, match and contain the information required. The absence of a described specimen may not be noted until the specimen container is opened in the sampling area of the lab. The absence of a described specimen is recorded as a non conformance. The sender is informed of the issue as soon as possible by the DAP staff." 3. Use of the Laboratory 3.4 Coll	15	16
	the sample is collected must be in place and followed. Any difficulty in obtaining the specimen should be noted on the request form. In the case of short or scanty specimens list tests requested in order of priority." 5. Transport of Specimens to the Laboratory		
DAP	Added: "5.6 BreastCheck Unit Specimens from the BreastCheck unit for the Division of Anatomic Pathology are delivered directly to DAP specimen reception by BreastCheck staff."	22	23

	9. Division of Anatomic Pathology		
DAP	9.1 Division of Anatomic Pathology 9.1 Division Profile: In: Histopathology provides Routine Histology and Advanced Diagnostic services, added: ", perinatal"	42	42
DAP	9. Division of Anatomic Pathology 9.1 Division Profile: Corrected: from Histolopathology to Histopathology	42	42
DAP	9. Division of Anatomic Pathology 9.1 Division of Anatomic Pathology 9.1 Division Profile: Changed: from ISO15189 2012 to ISO15189 2022 Added: "The Division of Anatomic Pathology ensure that patients well-being, safety, and rights primary considerations. The laboratory conforms to the HSE Code of Conduct and Behaviour in the provision of its service, including the rights of patients to care that is free from discrimination."	43	43
DAP	9. Division of Anatomic Pathology 9.2 General Information Added: "Special counselling may be needed for examination results with serious implications for the patient at the discretion of the clinical team. The DAP provides opportunities for patients and laboratory users to provide helpful information to aid the laboratory in the selection of the examination methods, and the interpretation of the examination results. Contact may be made directly with the Chief Medical Scientist (CMS), the Head of Department (HoD), and members of the Consultant staff. DAP staff Email and telephone contact information is given in section 2.3 of this document. Feedback may also be given via the "Your Service Your Say" mechanism accessible on the HSE website."	43	43
DAP	9. Division of Anatomic Pathology 9.3 Specimen Acceptance Added: "Note: Non adherence to the requirements for the specimen and the test request poses risks to the quality of the service the DAP is able to provide for the case concerned and for the patient. These risks include: rejection of the specimen, compromise to the specimen prior to receipt by the lab, compromise to the report, compromise to patient management, and or patient impact." Added: "Multiple samples on one form are acceptable and should be labelled A, B, C etc. where possible."	43	44
DAP	9. Division of Anatomic Pathology 9.4 Histopathology Radiation Specimen Added: "The request form and specimen containers must each be labelled with a radiation label. The radiation dose information must be given." Removed: "Ensure that radiation information is included on the request form and specimen container/s."	46	47
DAP	9. Division of Anatomic Pathology 9.4 Histopathology Outside normal working hours Please notify the Histopathology Department (ext. 4589) in advance of changed: from 17:00 to 16:00 Post Vasectomy Analysis It is best that the semen sample is delivered within 1 hour of production to the laboratory, Monday to Friday 09:00 to 11.30 and changed: from 14.00 to 16:00-h to 14.00 to 15:00 h	47	47
DAP	9. Division of Anatomic Pathology 9.5 Cytopathology Added: "Note: Where slides are being submitted for DAP analysis- the number of slides being submitted should be recorded on the request form."	48	48
DAP	Division of Anatomic Pathology S. Cytopathology EBUS (Ultra sound guided Endobronchial Specimens)	48	49
DAP	Added: "The number of slides being submitted should be recorded on the request form." 9. Division of Anatomic Pathology 9.6 Molecular Pathology removed: "a mutation service for", added: "in situ hybridisation service for confirmation of" Breast and Gastric HER-2; added: "status, and a mutation service for" Non Small Cell Lung Cancer; added: "adenocarcinoma"; EGFR/ALK/ROS-1 added: "/KRAS"; added: "NSCLC adenocarcinoma negative by the in-house panel will be referred to Cancer Molecular Diagnostics in St James Hospital for NGS analysis	49	49
DAP	with the Lunf Adenocarcinoma Focus assay." 9. Division of Anatomic Pathology 9.8 Turnaround times: removed: Revised Interim TAT (X working Days) 3-14 3-14 7-16 5	50	51
DAP	16. Alphabetical Test Directory Skin Punch Biopsy for Direct Immunofluorescence (DIF) In Specimen added: "Send the skin punch biopsy for DIF fresh."	173	174
GBTE	7. Services and Products available at GBTE removed: "*Praxbind (Idarucizumab) - reversal agent for Dabigatran"	25	26
GBTE	7. Blood and Tissue Establishment 7.2 Services and Products available at GBTE added: "cfDNA testing of the mothers blood can also be completed in early pregnancy. This is performed in a referral site (generally the IBTS). It predicts the fetus blood group and women who have a predicted Rh Negative fetus then do not enter the RAADP program for prophylactic Anti-D. The infants' blood group is then confirmed at birth."	26	27
GBTE	7. Blood and Tissue Establishment 7.3 Sample / Request Form Labelling Policy added: "cfDNA/ On Request/ 2 x 6ml EDTA blood/ Referred to Irish Blood Transfusion Service"	27	28
GBTE	7. Blood and Tissue Establishment 7.3 Sample / Request Form Labelling Policy Changed from "Please contact GBTE for external request forms or any queries regarding specimen referral." to "Please contact GBTE for external request forms or any queries regarding specimen referral."	27	28
GBTE	7. Blood and Tissue Establishment 7.17 Clinical Advice and Service Removed: *Dr. Sørcha Ni Løingsigh added: *Dr Maria Eduarda Couto, * Dr Abdelrahman Moutaz	34	34

	2. General Information		
MM	2.3 Contact Information	6	6
	Clinical Immunology		
	replaced: "Mr. Mike Cullina michael.cullina@hse.ie" with: "Mr. Arthur McQuaid arthur.mcquaidmichael.cullina@hse.ie" 2. General Information		
	2.7 Complaints		
	Added: "Complaints/compliments may be received verbally, by letter, fax or email.		
	Alternatively the complainant may:		
IMM	- Complete the HSE feedback form titled 'your service your say'	11	12
	- Email: yoursay@hse.ie		
	- Contact HSE your service your say contact number: 1800 424 555		
	A complaint can also be made by contacting the Laboratory manager or the relevant laboratory Chief Medical scientist at the contacts		
	given. The relevant laboratory will follow up complaints promptly as per their laboratories procedures."		
	11. Immunology Department (Supraregional Service)		
IMM	11.1 Department Profile	56	56
	removed: "or the joint immunology / rheumatology clinic (for connective tissue diseases)."		
	11. Immunology Department (Supraregional Service) 11.2 Guidelines for Requesting Allergy Tests		
IMM	Anaphylaxis	57	57
1101101	removed: "In the refractory period after anaphylaxis, specific IgE to the causative allergen may be falsely negative. Testing should	37	37
	generally be deferred for 3-4 weeks."		
	11. Immunology Department (Supraregional Service)		
	11.2 Guidelines for Requesting Allergy Tests		
IMM	Asthma and Rhinitis	57	57
	removed: " (usually by skin testing) "		
	replaced: " kiwi" with "fresh"		
	11. Immunology Department (Supraregional Service)		
	11.3 Guidelines for Requesting Tests for Autoimmune Disease		
	Coeliac disease		
	replaced: "IgA anti-tissue transglutaminase antibodies (tTg) or IgA anti-endomysial antibodies are found in active disease, and can be		
	used to monitor compliance with treatment. IgA anti-tTG is used as the screening test (more sensitive) and positive results confirmed		
	by IgA anti-endomysial testing (more specific). As part of quality assurance the test method can detect samples with absent IgA that		
	may cause false negative results. In patients with selective IgA deficiency the IgG anti-tTG assay is performed. NICE Guidelines, 2016s state that 'Testing for Coeliac disease is only accurate if the person continues to follow a gluten-containing diet during the testing		
IMM	period. Some gluten should be eaten in more than one meal every day for a minimum of 6 weeks before testing."	58	58
	with: "IgA anti-tissue transglutaminase antibodies (tTg) or IgA anti-endomysial antibodies are found in active disease, and can be used		
	to monitor compliance with treatment. IgA anti-tTG is used as the initial screening test (more sensitive) and only positive results are		
	confirmed once by IgA anti-endomysial testing (more specific).		
	As part of quality assurance the test method can detect samples with absent IgA that may cause false negative results. In patients with		
	selective IgA deficiency i.e. undetectable levels of IgA the IgG anti-tTG assay is performed. NICE Guidelines, 2016s state that 'Testing		
	for Coeliac disease is only accurate if the person continues to follow a gluten-containing diet during the testing period. Some gluten		
	should be eaten in more than one meal every day for a minimum of 6 weeks before testing'."		
	11. Immunology Department (Supraregional Service)		
IMM	11.3 Guidelines for Requesting Tests for Autoimmune Disease	58	58
	Anti-mitochondrial antibodies		
	replaced " cirrhosis" with "cholangitis" 11. Immunology Department (Supraregional Service)		
	11.4 Endocrine Disorders		
IMM	Thyroid	59	59
	Added: "Anti-TSH receptor antibodies are highly sensitive for the diagnosis of Grave's hyperthyroidism and related thyroid eye disease	33	55
	but can also be present in some individuals with Hashimoto's thyroiditis."		
	11. Immunology Department (Supraregional Service)		
	11.4 Endocrine Disorders		50
	11.4 Endocrine Disorders		
IMM	Diabetes Mellitus	59	59
IMM		59	59
IMM	Diabetes Mellitus Added: "For newly diagnosed type 1 diabetes it is recommended to request anti-GAD, anti-IA2 anti-ZnT8 antibodies."	59	29
IMM	Diabetes Mellitus Added: "For newly diagnosed type 1 diabetes it is recommended to request anti-GAD, anti-IA2 anti-ZnT8 antibodies." 11. Immunology Department (Supraregional Service)	59	29
IMM	Diabetes Mellitus Added: "For newly diagnosed type 1 diabetes it is recommended to request anti-GAD, anti-IA2 anti-ZnT8 antibodies." 11. Immunology Department (Supraregional Service) 11.6 Autoimmune Rheumatic and Renal Diseases	59	29
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IMM	Diabetes Mellitus Added: "For newly diagnosed type 1 diabetes it is recommended to request anti-GAD, anti-IA2 anti-ZnT8 antibodies." 11. Immunology Department (Supraregional Service) 11.6 Autoimmune Rheumatic and Renal Diseases Antinuclear antibody (ANA) added: ANA testing from General Practitioners for autoimmune rheumatic diseases is performed using the Connective Tissue Disease (CTD) screen. The CTD Screen is an automated method for the detection of anti-nuclear antibodies (ANA) in autoimmune rheumatic diseases such as SLE, mixed connective tissue disease, Sjogrens syndrome, Scleroderma and Myositis. The CTD Screen tests for anti-RNP, Sm, Ro, La, centromere B, Scl-70, Jo-1, Fibrillarin, RNA polymerase III, Ribosomal-P, PM-Scl, PCNA, Mi-2 and anti-dsDNA. Positive CTD screen results will have further testing for ANA (by indirect immunofluorescence), anti-ENA and anti-dsDNA where appropriate." 11. Immunology Department (Supraregional Service) 11.6 Autoimmune Rheumatic and Renal Diseases Cytoplasmic antibodies detected on ANA testing replaced: "Cirrhosis" with "Cholangitis" 11. Immunology Department (Supraregional Service)	60	60

ІММ	11. Immunology Department (Supraregional Service) Added: "11.10 Therapeutic Drug Monitoring Biologic therapies, including the anti-tumor necrosis factor (anti-TNF) agents (Infliximab, Adalimumab), the adhesion molecule inhibitors (Vedolizumab), and the p-40 interleukin-12/23 inhibitor Ustekinumab are effective treatments for patients with moderate to severe inflammatory bowel disease (IBD). Nevertheless, up to 1/3 of patients with Crohn's disease (CD) and ulcerative colitis (UC) show primary non-response (PNR) to biologic therapies and up to 50% of patients after an initial clinical response stop therapy either for secondary loss of response (SLR) or a serious adverse event. Drug trough levels and anti-drug antibodies enable the clinician, based on patient's clinical status, to make rational therapeutic decisions in different clinical situations: Reactive TDM: Guide therapy after a treatment failure and follow-up therapeutic adjustment (switch or optimization). Reactive TDM should be performed in patients with primary non-response or secondary loss of response to biologic therapy. Proactive TDM: Proactive TDM should be performed post induction for patients treated with anti-TNF therapy. Proactive TDM should be performed at least once during maintenance therapy for patients treated with anti-TNF therapy. Guides treatment de-escalation for patients in remission. When infliximab de-escalation (dose reduction) is considered in patients in remission, proactive TDM both prior to and after de- escalation should be performed. Reactive TDM has been proven more cost-effective than empiric anti-TNF therapy optimization. Decrease the risk of allergic reactions during infusion or other adverse effects The department of Immunology provides testing for trough levels and antibodies (where indicated) for the following biological drugs: Infliximab, Vedolizumab and Adalimumab. Ustekinumab analysis is referred externally for testing."	67	67
IMM	11. Immunology Department (Supraregional Service) Added: "11.11 Interferon Gamma Release Assay (IGRA/Quantiferon) Quantiferon TB Gold is an indirect test for latent Mycobacterium Tuberculosis infection (LTBI) and M. Tuberculosis complex infection. Latent Tuberculosis (LTBI) is an asymptomatic condition that may progress to active Tuberculosis in some individuals. The primary goal for the diagnosis of LTBI is to initiate medical treatment to prevent progression to active disease. Testing for LTBI is indicated when the risk of developing disease from latent infection (if present) is increased e.g Recent close contact of TB, immunosuppression, HIV infection, before commencing immunosuppression with biologic drugs that increase the risk of TB reactivation (e.g. anti-TNF), and occupational health screening for healthcare workers. The Interferon-Gamma Release Assay (IGRA/Quantiferon) measures the level of the cytokine, interferon-gamma (IFN-gamma) released by patient lymphocytes in a cell-mediated immune response to mycobacterial proteins. These proteins include ESAT-6, CFP-10 and TB7.7, and are absent from all BCG strains and most non-tuberculous mycobacteria. Although the assay quantitatively detects the IFN-gamma, the interpretation of the result for a single patient is strictly qualitative. The IGRA/Quantiferon assay requires specialised blood collection tubes. These tubes (set of 4) are available for collection from the Immunology laboratory. Correct handling of the blood collection tubes is essential. A negative Interferon-Gamma Release Assay (IGRA) result does not preclude the possibility of M. tuberculosis infection. False negatives can be due to incorrect handling of the blood collection tubes, the stage of the infection (e.g. sample taken prior to development of cellular immune response), or co-morbid conditions which affect immune function. All positive results should be followed by further medical evaluation. If the result is indeterminate for TB antigen responsiveness, this may be related to a wide vari	68	68
IMM	11. Immunology Department (Supraregional Service) 11.10 Guidelines relating to Genetic Referrals: Changed number to "11.12"	67	69
IMM	11. Immunology Department (Supraregional Service) 11.12 Guidelines relating to Genetic Referrals	67	69
IMM	Changed website from: "http://www.olchc.ie" to "www.childrenshealthireland.ie" 11. Immunology Department (Supraregional Service) 11.12 Guidelines relating to Genetic Referrals replaced: "Haemochromatosis genetic testing by the Molecular Genetics Lab, Northern Molecular Genetics Service, Biomedicine East, Central Parkway. Newcastle Upon Tyne, NE1 3BZ, UK: refer to http://www.newcastle-hospitals.org.uk/services/northern- genetics_services_molecular genetics.aspx Newcastle report is issued to the Clinician by Immunology. Paper report issued only-results not available on Healthlinks." with "Haemochromatosis genetic testing by Eurofins-Biomnis Dublin; Eurofins-Biomnis Haemochromatosis genetic report is issued to the Clinician by Immunology GUH. Paper report issued only - results not available on Healthlinks."	67	69
IMM	16. Alphabetical Test Directory Adalimumab (trough levels and antibodies) Laboratory: removed: ":- referred to Immunology Dept, Northern General hospital, Sheffield" Turnaround: replaced: "6-weeks" with "10 working days" Report: replaced: "Drug levels (mg/L): Suboptimal, therapeutic and supratherapeutic drug levels Antibodies: Negative = <10AU/ml" with: "Drug levels (mg/L): Suboptimal (<3ug/ml), therapeutic (3-7ug/ml) and supratherapeutic drug (>7ug/ml) levels Antibodies: Negative = <10ngAU/ml"	90	90
IMM	16. Alphabetical Test Directory Anti Cardiolipin Antibodies: added: "(IgG, IgM cardiolipin & Beta 2 glycoprotein)" Turnaround: replaced " 7 working days " with "5 working days"	96	96
IMM	16. Alphabetical Test Directory Added: "Anti-CV2/ CRMP5 Laboratory: Immunology: – referred to Immunology Department, Churchill Hospital, Oxford OX3 7LJ Specimen: 5.0 mL blood in plain gel tube. CSF analysis also available. Turnaround: 6 weeks"	96	96
IMM	16. Alphabetical Test Directory replaced: "Anti-GABA (anti-glutamate receptor antibodies)" with "Anti-GABA a /GABA b (anti-glutamate receptor antibodies)"	97	97
	16. Alphabetical Test Directory		
IMM	Anti-IA2 Antibodies replaced: "Positive/ Negative" with "Positive: >10 IU/ml; Negative 0-10 IU/ml"	98	98

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	16. Alphabetical Test Directory Added: "Anti-Parietal Cell Antibodies		
	Laboratory: Immunology		
IMM	Specimen: 5.0 mL blood in plain gel tube	101	101
	Turnaround: 5 Working days		
	Report: 0 – 10 EliA U/ml" 16. Alphabetical Test Directory		
	Added: "Anti-Thyroid Receptor Antibodies		
	Laboratory: Immunology		
IMM	Specimen: 5.0 mL blood in plain gel tube	103	103
	Turnaround: 10 working days Ref. Range: Negative: <2.9 IU/I		
	Equivocal: 2.9-3.3 IU/I		
	Positive: >3.3 IU/I"		
	16. Alphabetical Test Directory		
	Added: "Anti-ZNT8 Antibodies Laboratory: Immunology: – referred to Immunology Dept, Northern General hospital, Sheffield		
IMM	Specimen: 5.0 mL blood in plain gel tube	104	104
	Turnaround: 6 weeks		
	Report: Positive: >15U/ml; Negative <15U/ml"		
	16. Alphabetical Test Directory replaced: "AutoImmune Inflammatory Myopathy panel includes anti-: Mi-2 alpha, Mi-2 beta, TIF1 gamma, MDA5, NXP2, SAE1, Ku, PM-		
IMM	Sci100 and PM-Sci75, OJ, EJ, Jo-1, PL-7, PL-12, SRP and Ro-52" with "AutoImmune Inflammatory Myopathy panel includes anti-: Mi-2	105	106
	alpha, Mi-2 beta, TIF1 gamma, MDA5, NXP2, SAE1, Ku, PM-Scl100 and PM-Scl75, OJ, EJ, Jo-1, PL-7, PL-12, SRP, Ro-52, HMGCR and		
	CN1a."		
	16. Alphabetical Test Directory Cerebrospinal Fluid - Neurodegenerative biomarkers (CSF Tau/Phospho Tau/ Beta amyloid)		
	replaced: "Immunology St James Hospital Dublin" with "Clinical Chemistry, Tallaght University Hospital"		
	Specimen: repaced: "-2.5mls" with "minimum 2mL"		
IMM	Comment: replaced: "CSF by LP; received in Sarstedt 2ml screw cap tubes (contact lab for supply of tubes). Sample must reach the lab	115	116
	within 2hours of collection, Mon Friday Specific request form to be completed — obtained from Immunology lab" with "CSF by LP; received in Blue top Sarstedt CSF Collection tube (contact lab for supply of tubes)."		
	Turnaround: replaced: "-4-6 weeks" with "2-3 weeks"		
	Report: replaced: "St James" with "TUH"		
	16. Alphabetical Test Directory		
IMM	removed: Complement: CH100 (Total Haemolytic Complement) Functional Activity CH100 (Total) and CH100A (Alternate Pathway)	120	121
	Comment: replaced: "collectionon the same day it was taken" with "within 6 hrs of collection"		
	16. Alphabetical Test Directory		
	added: "Connective Tissue Disease Screen (CTD)		
	Laboratory: Immunology Specimen: 5.0 mL blood in plain gel tube		
IMM	Turnaround: 5 working days	120	122
	Ref. Range: Negative: <1.0		
	Positive: >1.0. Positive CTD screen results will have further testing for ANA (by indirect immunofluorescence), anti-ENA and anti-		
	dsDNA." 16. Alphabetical Test Directory		
	removed: "C V2/ CRMP5		
IMM	Laboratory: Immunology: – referred to Immunology Department, Churchill Hospital, Oxford OX3 7LJ	124	125
	Specimen: 5.0 mL blood in plain gel tube. CSF analysis also available.	124	123
	Turnaround: 6 weeks Report: Positive/Negative"		
	16. Alphabetical Test Directory		
IMM	Cystic Fibrosis – Genetic Test	124	126
	Comment: replaced: "www.olchc.ie" with "https://www.childrenshealthireland.ie"		
IMM	16. Alphabetical Test Directory Cytogenetics: Chromosome Analysis /Karyotyping Adults (age >5 years)	124	126
	Replaced: "(age >5 years)" with "(age >18 years)"		120
	16. Alphabetical Test Directory		
IMM	Cytogenetics: Chromosome Analysis /Karyotyping Paediatrics (age <5 years)	125	126
	Replaced: "(age <5 years)" with "(age <18 years)" Comment: replaced "www.olchc.ie" with "https://www.childrenshealthireland.ie"		
	16. Alphabetical Test Directory		
IMM	Cytogenetics: Microarray / aCGH	125	126
	Comment: replaced "www.olchc.ie" with "https://www.childrenshealthireland.ie" 16. Alphabetical Test Directory		
IMM	Cytotoxic Antibodies (solid organ transplantation)	126	127
	Ref. range: removed "form" added: "issued by Beaumont."	120	
	16. Alphabetical Test Directory		
IMM	Fragile X Chromosome Comment: replaced "ways alche is" with "ways children healthireland is"	133	135
	Comment: replaced "www.olchc.ie" with "www.childrenshealthireland.ie" 16. Alphabetical Test Directory		
	Haemochromatosis: replaced: "C282Y and H63D Genetic Mutations" with "C282Y, H63D and S65C Genetic Mutations"		
IMM	Laboratory: replaced: ":- referred to Molecular Genetics Lab. Northern Molecular Genetics Service, Biomedicine East, Central Parkway-	137	139
	Newcastle Upon Tyne, NE1 3BZ, UK" with "- referred to Eurofins Biomnis, Dublin." Turnaround: up to 8.3 weeks		
	Turnaround: up to 8 2 weeks		
	16. Alphabetical Test Directory		
	HLA B27 Typing Specimen: added: "5.0 mL EDTA blood (to be kept at room temperature only)"		
IMM	Comment: replaced: "Restricted test-restricted to the following disciplines Rheumatology, Ophthalmology & Orthopaedics. Please	143	145
	phone laboratory if there are exceptional reasons why this test is essential" with "Eurofins Biomnis Consent form for HLA testing to be		
	submitted with samples for (available at www.eurofins.ie/biomnis/test-information/test-request-forms)"		
	16. Alphabetical Test Directory		
15.45.4	HLA Typing	143	4.45
IMM	Specimen: replaced: "7.0 mL EDTA blood" with "5.0 mL EDTA blood (to be kept at room temperature only)" Comment: replaced: "Restricted test" with "Eurofins Biomnis Consent form for HLA testing to be submitted with samples for (available	143	145
	at www.eurofins.ie/biomnis/test-information/test-request-forms)."		

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removed: Richoechic CoFactor (RICOG) (WW F- RICOF) Laboratory Heamstology Specimen- 2 x 2.7 mL blood in a 0.109m Sodium Citrate tube. (1.0 mL Paediatric-tubes are available). Comment: Prior authorization by Consultant Haematologist or SPR. Arrange with coagulation laboratory before taking specimen. Must lill-bottles to mark. Turnaround-5 weeks Ref. Asange. Refer to report 16. Alphabetical Test Directory Von Willebrands removed: Faster Antigen (wWF-Ag), added "Screens" Specimen: 2 x 2.7 mL blood in 0.109m Sodium Citrate tubes added: "C. 1.0 mL Paediatric tubes are available)." 2. General Information: Medical Microbiology: Added: "Dr. Roisin Mulqueen Consultant Microbiologist Roisin.mulqueen3@hse.ie" 13. Medical Microbiology Department 13. Medical Microbiology Department 13.4 Out of Hours Service 13.4 Out of Hours Service 13.4 Out of Hours Service 23. Sonsultation Service 23. Sonsultation Service 23. Sonsultation Service 23. Medical Microbiology Department 13.4 Out of Hours Service 23. Sonsultation Service 23. Sonsultation Service 23. Medical Microbiology Department 13.5 Guidelines for Requesting Out Of Hours service 23. Medical Microbiology Department 13.5 Guidelines for Requesting Microbiology Tests Unine Samples replaced: "Culture is performed on all Unines. Unine microscopy is only performed routinely on children -18 years of age and pregnant women, however microscopy may be requested in sertian circumstances following discussion with a Consultant Microbiologist. Unine specimens that are received in plain universal containers that are older than 48 hours or unine specimens that are received in bioric acid containers and are more than 96 hours old are unsuitable for culture and will be rejected. Unine specimens that are received in anything but a yellow topped vacuette container as as shown in the image below are unsuitable for culture and will be rejected. Unine specimens that are received in anything but a yellow topped vacuette container as shown in the image below are unsuitable for culture and will b				
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Urine specimens that are received in plain universal containers that are older than 48 hours or urine specimens that are received in boric acid containers and are more than 96 hours old are unsuitable for culture and will be rejected. Urine samples submitted for microscopy and culture must be submitted in a urine sample tube, a Vacuum urine tube. Urine is initially collected in a primary urine beaker, then transferred via integrated transfer device to the Yellow Vacuette® urine tube, which is submitted to the laboratory. Do not submit the transfer beaker to the Laboratory as it will be rejected." with: "Urine microscopy is performed on all Urines. Urine culture is only routinely performed on samples from children <16, maternity patients, clinical details specifying patient is neutropenic and patients with a microscopy result with a white cell count of >20cmm, however culture may be requested in certain circumstances following discussion with a Consultant Microbiologist. Urine specimens that are received in anything but a yellow topped vacuette container as shown in the image below are unsuitable for culture and will be rejected. Urines must be decanted from the beaker into the tube before being sent to the laboratory. Beakers send to the laboratory that have not been decanted into the urine vacuette will not be processed and will be disposed of immediately. Urine is initially collected in a primary urine beaker, then transferred via integrated transfer device to the Yellow Vacuette® urine tube,	Micro			
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Micro	13. Medical Microbiology Department 13.5 Guidelines for Requesting Microbiology Tests GUH National Reference Laboratory National Salmonella, Shigella and Listeria Reference Laboratory replaced: "https://saolta.ie/documents/national-salmonella-shigella-listeria-reference-laboratory-users-guide" with: "https://saolta.ie/documents/galway-reference-laboratory-service-incorporating-national-salmonella-shigella-listeria"	81	82
Micro	13. Medical Microbiology Department 13.5 Guidelines for Requesting Microbiology Tests National Carbapenemase Producing Enterobacterales Reference Laboratory (CPERL) rempaced: "https://saolta.ie/documents/national-carbapenemase-producing-enterobacterales-cpe-reference-laboratory-users-guide" with: "https://saolta.ie/documents/galway-reference-laboratory-service-incorporating-national-salmonella-shigella-listeria"	82	83
Micro	15. Out Of Hours (Emergency Service) removed referral to: Blood Culture 2	86 & 88	87 & 89
Micro	15. Out Of Hours (Emergency Service) Requiring Consultation removed: "2. Submit vial before 23:00" replaced: 3 with 2, 4 with 3, 5 with 4, 6 with 5 in 4. removed: "mid"	89	89
Micro	16. Alphabetical Test Directory Amikacin: added: "Referred to external laboratory."	93	93
Micro	16. Alphabetical Test Directory Bacterial PCR (For sterile fluids and Tissues) S.aureus PCR (Mec A and CoA), Group A Streptococcus DNA, N. meningitidis DNA, Haemophilus influenzae DNA and Streptococcus pneumoniae DNA. Laboratory: added: "- referred to Great Ormonde Street Hospital" Turnaround: repaced: "1 week (Verbal report available on positive samples)" with "2 weeks (Verbal report available on detected targets)"	106	106
Micro	16. Alphabetical Test Directory Blood Culture Comment: removed: "Delivery by Porter if glass bottles."	108	109
Micro	16. Alphabetical Test Directory Bone Marrow Culture Comment: removed: "Delivery by Porter if glass bottle."	109	110
Micro	16. Alphabetical Test Directory Bordetella Species (Whooping cough / Pertussis)- culture replaced: "Laboratory: Medical Microbiology Specimen: Perinasal swab (available from Medical Microbiology) Comment: Contact Laboratory prior to sending to ensure fresh media is available. If delay refrigerate @ 2-80C. Turnaround: 10 days Report: "Bordetella-pertussis" Not-isolated or "Bordetella-pertussis" isolated with "See Whooping Cough"	109	110
Micro	16. Alphabetical Test Directory Cerebrospinal Fluid (Molecular analysis for Pathogens) Laboratory: removed: ":- referred to Irish Meningococcal and Streptococcal Reference Laboratory /National Virus Reference Laboratory when unavailable on site." Turnaround: replaced: "1-2 weeks (Verbal report available on positive samples within 2-5 working days)" with: "1-2 working days (Verbal report available on detected targets)" Repoert: added: "Targets"	114	115
Micro	16. Alphabetical Test Directory Cerebrospinal Fluid – Viral PCR removed: "(HSV and VZV)" Turnaround: replaced: "1-2 weeks (Verbal report available on positive samples)"	114	115
Micro	with: "1-2 working days (Verbal report available on detected targets)" 16. Alphabetical Test Directory Faeces – Molecular analysis, Microscopy, Culture and Antigen Detection report: removed: "Any clinically significant isolate-all samples with pathogen DNA detected (Except Campylobacter spp)" added: "When Salmonella DNA or Shigella/EIEC DNA is detected. Referral to Cherry Orchard when VTEC DNA is detected."	131	133
Micro	16. Alphabetical Test Directory Herpes simplex virus - PCR Laboratory: Removed: "Medical-Microbiology", added: "Virology – referred to National Virus Reference Laboratory, Dublin" Specimen: removed: "0.5 mL CSF in plain leak-proof sterile container or" swab in viral transport medium from genital site.	142	143
Micro	16. Alphabetical Test Directory Legionella culture Comment: Added: "Routinely on ICU specimens. On request following approval by a Consultant Microbiologist on non-ICU specimens."	150	151
Micro	16. Alphabetical Test Directory Meningococcal PCR Laboratory: added: " - Referred to the Irish Meningococcal and Sepsis Reference Laboratory" Specimen: removed: "G reater than 200 ul CSF in a sterile plain tube or " Comment: removed: "S ample to be handed to Medical Microbiology staff member " Report: replaced: "Meningococcal DNA" with "Target"	154	155
Micro	16. Alphabetical Test Directory MSU – Midstream Urine Specimen: Replaced: "Specimen in Boric Acid Universal container. Use plain sterile Universal container for Paediatric specimen or urine volumes < 20 mL." with "Specimen in urine vacuette tube."	157	158
Micro	Micros Ced micros With Specimen in unine vacuette tube. 16. Alphabetical Test Directory Mycobacteria Testing Comment: added: "Culture is only performed on all tissue and fluid samples or where clinical details query MOTT "	157	159
Micro	16. Alphabetical Test Directory Mycobacteria PCR - Xpert assay Comment: replaced: "Xperts are performed on all initial specimens with AAFB seen on microscopy or by prior arrangement with Microbiology Medical Staff: " with: "Xperts are performed on all samples requesting TB. Culture is only performed on all tissue and fluid samples or where clinical details query MOTT."	158	159
Micro	16. Alphabetical Test Directory Pneumococcal PCR Laboratory: added: "– referred to the Irish Meningococcal and Sepsis Reference Laboratory" Specimen: removed: "Greater than 200 ul CSF in a sterile plain tube or "	164	165

	16. Alphabetical Test Directory		
Micro	Tuberculosis Testing Comment: added: "Culture is only performed on all tissue and fluid samples or where clinical details query MOTT"	180	181
	16. Alphabetical Test Directory		
Micro	Whooping Cough	185	186
IVIICIO	Laboratory: added: " – referred to Our Lady's Children's Hospital Crumlin [OLCHC]"	163	100
Mort	Removed: "Comment: Contact Laboratory prior to ensure fresh media is available:"		
WIGHT	2. General Information		
PHLEB	2.3 Contact Information	10	10
PHLER	Phlebotomy Department	10	10
	Added: "Senior Phebotomist, Bleep: 835		
	4. Phlebotomy Service UHG OPD:		
PHLEB	replaced: "Mon – Thurs 08:00 – 18:00 Friday 08:00 – 14:00" with "Mon – Thurs 09:00 – 18:00 Friday 09:00 – 14:00"	18	19
	Merlin Park University Hospital	10	13
	added: "Friday 09:00 – 14:00"		
	2. General Information		
Viro	2.5 Laboratory Opening Hours	11	11
	Virology Changed routine hours from: "08:00 – 17:30 h Mon-Fri" to "08:00 – 17:00 h Mon-Fri"		
	13. Medical Microbiology Department (Division of Clinical Microbiology)		
Viro	13.3 Consultation Service	73	75
	Added: "Dr Roisin Mulqueen"		
	13. Medical Microbiology Department (Division of Clinical Microbiology)		
	13.4 Out of Hours Service SARS COV-2 PCR testing Out Of Hours service		
	Removed: "Monday - Friday 20.00 - 08.00 the following day		
Viro	Saturday 16.00 – 08.00 the following day	74	76
	Sunday 08.00 - 08.00 the following day"		
	Added: "Monday – Friday 17.00 – 18.30		
	Saturday 08.00 – 13.00		
	Sunday 08.00 – 13.00" 14. Virology Department (Division of Medical Microbiology)		
Viro	14.2 Availability of Clinical Advice and Interpretation	83	84
	Added: ", Dr. Roisin Mulqueen (Ext4146)"		
	14. Virology Department (Division of Medical Microbiology)		
Viro	14.5 Virology Tests	85	86
	Influenza and RSV Detection Removed: "A" from "AA nasal/nasopharyngeal" and added: "swab "		
	14. Virology Department (Division of Medical Microbiology)		
Viro	14.8 Telephoning for Virology Results	85	86
	Added "require."		
\ fi==	16. Alphabetical Test Directory	100	110
Viro	Bordetella pertussis antibodies Corrected: from "Collindale Avenue" to "Colindale Avenue"	109	110
	16. Alphabetical Test Directory		
	Erythrovirus B19 IgM + IgG antibodies		
Viro	Added: "(Parvovirus)"	130	131
	Removed: "-referred to the National Virus Reference Laboratory, Dublin"		
	Updated: from " 3 weeks " to "5 days" 16. Alphabetical Test Directory		
Viro	Hepatitis A Virus Total Antibody	140	142
	Replaced: "Hepatitis A Virus Total Antibody" with "IgG"		
	16. Alphabetical Test Directory		
Viro	Hepatitis B DNA / Viral Load	140	142
	removed: ":-referred to the National Virus Reference Laboratory, Dublin" turnaround: replaced: "1—3 weeks" with "10 days"		
	16. Alphabetical Test Directory		
	Removed: "Hepatitis C Antigen		
	Laboratory: Virology		
Viro	Specimen: 7.0 mL blood in a plain gel tube Comment: Only available in very specific cases and following approval by a Consultant Microbiologist	141	143
	Turnaround: 3-5 working days		
	Report: Not Detected/Detected"		
	16. Alphabetical Test Directory		
Viro	Hepatitis C PCR / Viral Load / Genotype	141	143
	In Laboratory added: "Hep C Genotype is performed in the NVRL." 16. Alphabetical Test Directory		
	Human Immunodeficiency (HIV) PCR / Viral Load / Genotype		
Viro	Turnaround: replaced: "1—3 weeks" with "10 days"	144	145
	Report: replaced: "Detected/Not detected" with: "Not Detected/copies/ml with comment where relevant."		
Vi	16. Alphabetical Test Directory	457	450
Viro	Mumps IgG antibody Added: "Report: Detected/Not Detected/Equivocal"	157	158
	16. Alphabetical Test Directory		
Viro	SARS COV – 2 (PCR)	171	172
VIIO	Report: added: "Detected weak", removed: "Indeterminate", added: "Whole Genome Sequencing (WGS) is performed upon request of	1/1	1/2
	SARS-CoV-2 positive samples."		
Viro	16. Alphabetical Test Directory Varicella-zoster Virus IgG antibodies	183	184
VIIO	Report: added: "/Indeterminate"	103	104
	16. Alphabetical Test Directory		
	Added:		
	"Zika		
Viro	Laboratory: Virology. Referred to the National Virus Reference Lab.	186	187
	Specimen: 7.0 ml blood in a plain gel tube Comment: Only available in very specific cases and following approval by a Consultant Microbiologist		
	Turnaround: 3 weeks"		