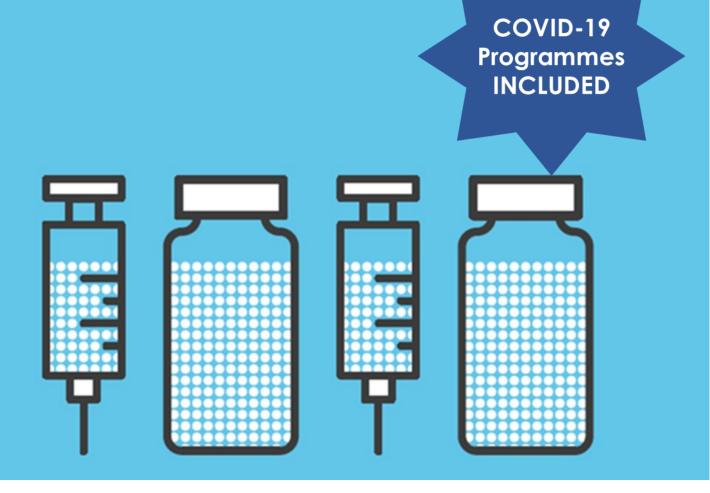
Services Available List 2021-2022

United Kingdom
National External
Quality Assessment Service





To find out more about our services visit our website: www.ukneqas.org.uk or our twitter/LinkedIn page: @UKNEQAS

Who are UK NEQAS?

Our Mission:

Improving global diagnostic testing for the benefit of patients through quality assessment and education

For over 50 years members of UK National External Quality Assessment Service (UK NEQAS) have been the world leading providers of Clinical Laboratory External Quality Assessment (EQA)/Proficiency Testing (PT) programmes. At present UK NEQAS offers over 500 different EQA/PT programmes covering over 5000 unique analytes, ensuring full coverage of all aspects of clinical laboratory testing worldwide. These programmes support the delivery of accurate and precise clinical laboratory tests on patients regardless of where they are performed. Additionally, findings from UK NEQAS programmes support the identification of best practice amongst service users and are often used to produce national and international guidelines. This promotes the implementation of best practice procedures worldwide and benefits patient care on a global scale.

The EQA/PT services offered by UK NEQAS are continually developing in response to changes in clinical laboratory requirements, and innovation is an area in which UK NEQAS has always led. This has been highlighted recently with the COVID-19 pandemic, where the UK NEQAS consortium has developed several EQA/PT programmes. UK NEQAS rapidly deployed EQA/PT programmes with some achieving accreditations to ISO 17043:2021 in just 12 weeks from conceptual development. We now have a number of programmes which cover the detection and monitoring of COVID-19 along with several educational competency programmes for staff undertaking swabbing and sample handling. These programmes are listed in this brochure on a specific COVID-19 page for ease of identification.

Our EQA/PT services have always included assessment of the 'end-to-end' process of patient investigation, including interpretation of results, either in stand-alone programmes or integrated into standard exercises. These traditional EQA/PT services are enhanced by our Pre and Post Analytical Quality Monitoring service (known as PrepQ), in which we have established a true end-to-end system, allowing laboratories to monitor pre and post analytical issues to support the requirements of ISO 15189.

UK NEQAS EQA/PT services are provided by expert centres in 11 cities around the UK, with each covering a specific area of clinical laboratory EQA/PT. These centres form the UK NEQAS Consortium, a not-for-profit company limited by guarantee and a UK Registered Charity (no: 1044013). To ensure the provision of high quality and independent EQA/PT services every UK NEQAS member centre has no commercial interests, is staffed by a dedicated team of EQA/PT specialists, operates on a strict not-for-profit basis, is accredited to ISO 17043, and operates programmes specifically designed to improve patient care.

At UK NEQAS we are committed to the provision of EQA/PT services that support the continual improvement of healthcare services by

- providing education
- facilitating clinical audit
- monitoring clinical effectiveness
- assisting in the management of risk
- · generating information used for research and development

By providing EQA/PT that fulfil these requirements, members of the UK NEQAS Consortium support laboratories in the accurate and precise delivery of their services ensuring patients receive the highest quality diagnostic testing wherever in the world they may be.

The worldwide partnerships between UK NEQAS and laboratories are ultimately for the benefit of patients and we are proud to support high quality healthcare through these relationships. We realise that there is always work required to develop EQA/PT systems, we are committed to undertake this process and always welcome feedback and suggestions on how we can improve our services.

Thank you for your continuing support of UK NEQAS.

UK NEQAS President

L. Whiley

Contents

Page 5 Cellular Pathology Technique Head & Neck Pathology Cellular Pathology Neuropathology Immunocytochemistry & In-Situ Hybridisation Page 6 - 7 Birmingham Quality Cardiac Markers **Clinical Chemistry** Trace Flements **Guildford Peptide Hormones Edinburgh Peptide Hormones** Page 8 - 10 **Blood Coagulation** Blood Transfusion Laboratory Practice (BTLP) Haematology Haematology Vitamin K Leucocyte Immunophenotyping Page 11-12 **Immunology** Immunology, Immunochemistry & Allergy Histocompatibility and Immunogenetics Page 13 Microbiology **Medical Microbiology** Parasitology Page 14 - 17 **Genomics &** Reproductive Science **Reproductive Science** Genomics **Covid-19 Programmes** Page 18 Point of Care Testing Page 19 **A-Z Index** Page 20 - 26 **Participating Countries** Page 27 **UK NEQAS Contacts** Page 28

Cellular Pathology

Cellular Pathology Technique
Tel: +44 (0) 191 816 1030
Email: cpt@ukneqas.org.uk
Legal Entity: LabXCell LTD
Services Available
Tissue Diagnostics
Specialist Techniques
Bone Marrow Trephine Biopsy (BMT)
Diagnostic Cytopathology
Direct Immunofluorescence (DIF)
Mohs' Procedure
Muscle Histochemistry
Neuropathology
Renal Biopsy Pathology
Transmission Electron Microscopy (TEM)
Frozen Section (Companion Scheme to Specialist Techniques and Neuropathology)
Mega Block (Companion Scheme to Specialist Techniques and Neuropathology)
Digital Interpretative Diagnostic Cytopathology
Digital Diagnostic Ultrastructural Pathology*
Digital Pathology Imaging*

Head and Neck Pathology				
Tel: +44 (0) 121 371 5723				
Email: neckpath@ukneqas.org.uk				
Legal Entity : British Society for Oral & Maxillofacial				
Services Available				
Head and Neck Pathology (Oral, ENT & Combined)				

Neuropathology		
Tel : +44 (0) 1865 234 904		
Email: neuropath@ukneqas.org.uk		
Legal Entity : Oxford University Hospitals NHS Foundation Trust		
Services Available		
Neuropathology Interpretative EQA Scheme		

Tel: +44 (0) 208 187 9174			
Email: info@ukneqasic			
Legal Entity: External Quality Assessment			
Services for Cancer Dic	Ü		
Services	Available		
	Alimentary Tract Pathology (GIST)		
	Breast Pathology (HER2 IHC)		
	Breast Pathology (Hormonal Receptors ER & PR)		
	Breast Pathology (Hormonal Receptors ER only)		
	Cytopathology		
	Gastric HER2 IHC		
	General Pathology		
	Lymphoid Pathology		
Immunocytochemistry	Mismatch Repair (MMR) Proteins		
	Neuropathology		
	Non-Small Cell Lung Cancer (NSCLC) ALK IHC		
	Non-Small Cell Lung Cancer (NSCLC) PD-L1 IHC*		
	Non-Small Cell Lung Cancer (NSCLC) ROS1 IHC*		
	Triple Negative Breast Cancer (TNBC) PD-L1 IHC*		
In-Situ Hybridisation	Breast HER2 ISH (Interpretive and Technical)		
	Non-Small Cell Lung Cancer (NSCLC) ALK ISH*		
	Non-Small Cell Lung Cancer (NSCLC) ROS1 FISH*		

Immunocytochemistry & In-Situ Hybridisation

Clinical Chemistry

Birmingham Quality

Tel: +44 (0) 121 414 7300

Email: birminghamquality@uhb.nhs.uk

Legal Entity: University Hospitals Birmingham

NHS Foundation Trust

Services Available

Antibiotic Assays

Antifungal Assays

Clinical Chemistry

Colorectal Cancer Screening

Erythropoietin

Faecal Haemoglobin

Faecal Markers

Faecal Pancreatic Elastase

Fluids

Fructosamine

GFR Estimations (Creatinine, Cystatin C &

eGFR)

Glycated Haemoglobins

Haematinics

Holotranscobalamin

Immunosuppressants

Interpretative Comments in Clinical Chemistry

Intrinsic Factor Antibodies

Lipid Investigations

Newborn Screening

Paediatric Bilirubin

Quantitative Amino Acids

Serum Indices (HIL)

Specific Proteins

Steroid Hormones

Sweat Testing

Thyroglobulin

Thyroid Hormones

Toxicology (TOX) and Therapeutic Drug

Monitoring (TDM)

Urinary Catecholamines & Metabolites

Urine Chemistries

Urine Dipsticks

Vitamin Assays (Carotene, Vitamin A & E)

Vitamin D

Cardiac Markers

Tel: +44 (0) 141 440 2888

Email: info@uknegas-cm.org.uk

Legal Entity: NHS Greater Glasgow & Clyde

Services Available

Cardiac Markers Schemes (Laboratory)

Cardiac Markers Schemes (Point of Care)

B-Type Natriuretic Peptide (BNP)

Cardiac Troponin I

Cardiac Troponin T

СКМВ

Myoglobin

NT-proBNP

Trace Elements

Tel: +44 (0) 1483 571 122 ext. 3611

Email: rsc-tr.Guildford-EQA@nhs.net

Legal Entity: Frimley Health NHS Foundation

Trust

Services Available

Lead, Cadmium, Manganese, Arsenic, Magnesium, Mercury, Zinc, Selenium, Thallium, Chromium and Cobalt in Blood

Mercury, Cadmium, Copper, Iron, Zinc, Chromium, Cobalt, Arsenic, Lead, Manganese, Nickel, Thallium and Selenium in Urine

Zinc, Copper, Aluminium, Selenium, Chromium and Cobalt in Serum

Aluminium in Water/Dialysis Fluid (Educational)

Copper and Iron in Solid Matrices e.g. Liver (Educational)

Guildford Peptide Hormones

Tel: +44 (0) 1483 571 122 ext. 3611

Email: rsch.peptideeqa@nhs.net

Legal Entity: Frimley Health NHS Foundation

Trust

Services Available

C-Peptide

Gastrin

Insulin

Insulin-like Growth Factor 1

Insulin-like Growth Factor Binding Protein-3

Clinical Chemistry

Edinburgh Peptide Hormones			
Tel: +44 (0) 131 242 6885			
Email: ukneqas@ed.ac.u	k		
Legal Entity: NHS Lothian			
Services A	vailable		
	1st Trimester (Down's syndrome and Trisomies T13 and T18)		
Maternal Serum	2nd Trimester (Down's syndrome)		
Screening	2nd Trimester (Neural tube defects)		
	1st Trimester (Down's syndrome using dried blood spots)		
D	Placental Growth Factor (PLGF)		
Pre-eclampsia Markers*	SFIt-1		
	SFlt-1: PLGF ratio		
Peptide I	FSH, LH, AMH, Prolactin and Growth Hormone		
Peptide II	PTH, ACTH and Calcitonin		
Tumour Markers	AFP, CEA and hCG		
Pregnancy Testing	Urinary hCG (Qualitative and Quantitative)		
	Type III Procollagen Peptide (PIIINP)		
Liver Fibrosis Markers*	Hyaluronic acid		
	Tissue inhibitor of metalloproteinase I (TIMP-I)		
	Enhanced Liver Fibrosis (ELF) Score		
	FIB-4 Score		

Haematology

Blood Coagulation			
Tel : +44 (0) 114 267 3300			
Email: neqas@coageq	a.org.uk		
Legal Entity: Sheffield To	eaching Hospitals NHS		
Foundation Trust	A		
Services	Available Activated Partial		
	Thromboplastin Time (APTT)		
	D-Dimer		
	Fibrinogen Evaluation (Clauss method)		
	Heparin Assay (HA) - anti-Xa		
Screening tests	Heparin Dosage Assessment (HDA) by APTT		
	Prothrombin Time (PT)/INR (Venous/Capillary methods)		
	Prothrombin Time for Diagnosis (PT/ Diagnostic) (PTD)		
	Thrombin Time (TT)		
	Factor II, V, VII VIII, IX, X, XI, XII Assays		
	Direct Oral Anticoagulant (DOAC) Assay*		
	Von Willebrand Factor Antigen: Assay		
Assays	VWF:RCo (activity) Assay		
Assays	VWF:CB (collagen-binding) Assay		
	Antithrombin Antigen and Activity Assays		
	Protein C Antigen and Activity Assay		
	Protein S Total and Free Antigen Assay		

Blood Coagu	lation Continued
Assays	Protein S Activity Assay
	Activated Protein C Resistance Assay
	Plasminogen Assay
	DOAC assays
	FXIII assays
	Homocysteine assays
	Quantitative Factor VIII Inhibitor
	ADAMTS13 assays
	Lupus anticoagulant screening/assay
Other programmes	FVIII treatment monitoring
Other programmes	FIX treatment monitoring
	Emicizumab assays
	HIT screening/assays
	Molecular Genetics of Thrombophilia Testing (Factor V Leiden)
	Genetics of Heritable Bleeding and Thrombotic disorders
	Activated Clotting Time (ACT)
	POCT D-dimer
POCT	CoaguChek XS, CoaguChek XS Plus, CoaguChek XS Pro & Pro II (PT/INR)
	Thromboelastometry and Thromboelastography

Haematology

Blood Transfusion Laboratory Practice (BTLP)

Tel: +44 (0) 1923 217 933

Email: btlp@uknegas.org.uk

Legal Entity: West Hertfordshire Hospitals NHS

Trust

11031				
Services Available				
	ABO & D Grouping			
	Antibody Screening			
	Antibody			
Pre-Transfusion Testing (PTT)	Identification			
(٢11)	Crossmatching			
	Rh and K			
	Phenotyping			
Fetomaternal	FMH Screening			
Haemorrhage (FMH)	FMH Quantification			
456 = 11	ABO Titration			
ABO Titration (ABOT)	A ₁ typing			
	Detection of cells			
Direct Antiglobulin	coated with IgG			
Testing (DAT)	Detection of cells			
	coated with C3d			
Point of Care D typing (POCTD)	RhD typing			
	Red Cell Genotyping (RCG)*			
	,			
Pilot schemes	Antenatal Antibody Titration (ANT)*			
	Extended Red Cell			
	Phenotyping (ERP)*			
Other services	Training, Assessment			
(supplementary to EQA)	and Competency Tool (TACT)			
(Joppielliellial)	IOOI (IACI)			

Vitamin K

Tel: +44 (0) 207 188 6815

Email: keqas@ukneqas.org.uk

Legal Entity: Viapath Analytics

Services Available

Vitamin K1 at endogenous concentrations in human serum

MK-4 at endogenous concentrations in human serum

MK-7 at endogenous concentrations in human serum

Vitamin K1, 2, 3-epoxide at endogenous concentrations in human serum

PIVKA-II (undercarboxylated prothrombin) at endogenous concentrations in human serum

н	ae	m	a	t٥	In	a	v
	u		•	_	•	9	7/

Tel: +44 (0) 1923 217 878

Email: haem@ukneqas.org.uk

Legal Entity: West Hertfordshire Hospitals NHS

Trust				
Services	Available			
Automated Counting & Related	Automated Differential Leucocyte Count			
	ESR			
	Full Blood Count (FBC)			
	Haemoglobin only			
	Plasma Viscosity			
	Infectious			
	Mononucleosis*			
	Reticulocyte Count			
	Abnormal			
	Haemoglobins			
	DNA Diagnostics for Haemoglobinopathies			
Haemoglobinopathy	Liquid Newborn Haemoglobinopathy Screening			
	Newborn Sickle Screening			
	Sickle Screening only			
	Blood Films for			
	Morphology			
	Blood Parasite			
	Screening and Identification			
	(Haematology)			
AA a wala ala wa Balada d	Cytochemistry: Haemosiderin staining			
Morphology Related	Digital Blood Film			
	Morphology for CPD			
	Malaria Rapid			
	Diagnostic Test			
	(Haematology)			
	Manual WBC			
	Differential Count			
Red Cell Enzyme	G6PD Screen and Assay			

^{*}Pilot/Non Accredited

For accreditation status please refer to www.ukas.com

Haematology

Leucocyte Immunophenotyping		Leucocyte Immu	nophenotyping Continued
Tel: +44 (0) 114 267 3600			BCR-ABL1 Minor
Email: admin@ukneqasli.co.uk			Quantification*
Legal Entity: Sheffield Teaching Hospitals NHS Foundation Trust			BRAF p.Val600Glu (V600E) Mutation Status
Services Available			for Hairy Cell Leukaemia
	CD34+ Stem Cell Enumeration		FLT3 Mutation Status
	Immune Monitoring		lgH/TCR Clonality Status
	Leukaemia Immunophenotyping (Part 1)		JAK2 p.Val617Phe (V617F) Mutation Status
	Leukaemia Diagnostic Interpretation (Part 2)		KIT p.Asp816Val (D816V) Mutation Status for Mast Cell Disease
	Low Level Leucocyte Enumeration		NPM1 Mutation Status
	Minimal Residual Disease for ALL by Flow Cytometry	Molecular	Paediatric Acute Leukaemia Translocations
	Minimal Residual Disease for AML by Flow Cytometry*		Post-Stem Cell Transplant Chimerism Monitoring
Flow Cytometry	Minimal Residual Disease for CLL by Flow Cytometry*		Acute Myeloid Leukaemia and Myelodysplastic Syndrome Gene Panels*
	Minimal Residual Disease for Plasma Cell Myeloma by Flow Cytometry*		Chronic Lymphocytic Leukaemia Gene Panels*
	Haematological Malignancy Bone Marrow Aspirate Assessment*		Lymphoplasmacytic Lymphoma/ Waldenström Macroglobulinemia*
	Cerebrospinal Fluid (CSF) Immunophenotyping*		Myeloproliferative Neoplasm Gene Panels*
	Paroxysmal Nocturnal Haemoglobinuria		
Molecular	BCR-ABL1 and AML Translocation Identification		
	BCR-ABL1 Kinase Domain Variant (Mutation) Status		
	BCR-ABL1 Major Quantification	*Pilot/Non Accredited For accreditation status p	lease refer to www.ukas.com

Immunology

Immunology, Immunochemistry & Allergy			
Tel : +44 (0) 114 271 5715			
Email: immunology@ukr	neqas.org.uk		
Legal Entity: Sheffield Te	eaching Hospitals NHS		
Foundation Trust Services	Available		
Jei vices /	Allergen Component		
	Testing		
	Allergen Specific IgE		
Allergy and	Antibody to Fungal & Related Antigens		
Immunodeficiency	lgG Subclasses		
	Specific Microbial Antibodies		
	Total IgE		
	Tryptase		
	Acetylcholine Receptor Antibodies		
	ANCA/GBM		
	Antibodies		
	Antibodies to Nuclear and Related Antigens		
	Bullous Dermatosis Antibodies		
	Coeliac Disease Antibodies		
	Diabetic Markers		
Autoimmunity	Ganglioside Antibodies		
,,	General Autoimmune Serology		
	Interferon Gamma Release Assays (Mycobacterium tuberculosis) IGRA TB		
	Myelin Associated Glycoprotein IgM Antibodies (MAG)		
	Myositis Associated Antibodies		
	Paraneoplastic Antibodies		

IIA continued			
Autoimmunity	Phospholipase A2 Receptor Antibodies (PLA2R)		
	Phospholipid Antibodies		
	SARS-CoV-2 Antibody Detection		
	Alkaline Phosphatase (ALP) Isoenzymes		
	Alpha 1 Antitrypsin and Phenotype Identification		
	C-Reactive Protein (CRP)		
	C1 Esterase Inhibitor and Functional Complement Assays		
	CSF Haem Pigments		
Immunochemistry	CSF IgG Oligoclonal Bands		
	CSF Proteins and Biochemistry		
	CSF β2 Transferrin/Beta Trace Protein		
	Interleukin-6 (IL-6)*		
	Point of Care CRP		
	Procalcitonin		
	Ultrasensitive C-Reactive Protein		
	β2 Microglobulin		
	Cryoproteins (image based)		
Oncology	Monoclonal Protein Identification		
	Prostate Specific Antigen (PSA)		
	Ultrasensitive PSA		
	Tumour Markers (CA Series)		

Immunology

Histocompatibility and Immunogenetics

Tel: +44 (0) 1443 622 185

Email: handi@ukneqas.org.uk

Legal Entity: Velindre University NHS Trust

Services Available

Crossmatching by Flow Cytometry

Cytotoxic Crossmatching

DNA HLA Typing at 1st Field Resolution

DNA HLA Typing to 2nd or 3rd Field Resolution

Educational Cross Match

Educational Interpretive Clinical Scenarios

Educational Scheme

HFE Typing

HLA Antibody Detection

HLA Antibody Specificity Analysis

HLA Genotyping for Coeliac and other HLA

associated diseases

HLA Phenotyping

HLA-B*57:01 Typing for Drug Hypersensitivity

HLA-B27 Testing

HPA Antibody Detection/Specification

HPA Genotyping

Interpretative: HFE Genotype and Hereditary

Haemochromatosis

Interpretive HLA Genotype

KIR Genotyping

Medical Microbiology

Microbiology			
Tel: +44 (0) 208 905 9890			
Email: organiser@ukneq	asmicro.org.uk		
Legal Entity: Public Heal	th England		
Services A	Available		
	AAFB Microscopy		
	Antimicrobial		
	Susceptibility		
	Clostridioides Difficile		
	(prev. Clostridium		
	Difficile)		
	Community Medicine		
Bacteriology	Faecal Pathogens		
	General Bacteriology		
	Genital pathogens		
	MRSA Screening		
	Mycobacteria Culture		
	Syphilis Serology		
	Urinary Antigens		
	C. trachomatis and		
	N. gonorrhoeae		
	CMV DNA		
	Quantification		
	EBV DNA		
	Quantification		
	HBV DNA		
	Quantification		
	Hepatitis C RNA Detection		
	HIV 1 RNA Quantification		
	Molecular Detection		
	Mycobacteria		
Molecular	Molecular detection		
	of HEV RNA		
	Molecular Detection		
	of HPV		
	Molecular Detection		
	of Respiratory Viruses		
	Molecular Detection		
	of SARS-CoV-2		
	Molecular Detection		
	of Viruses in CSF		
	Viral Gastroenteritis		
	(also suitable for		
	antigen testing)		

Microbic	ology continued
Mycology	Antifungal Susceptibility
	7 timorigai sosceptibility
	Cryptococcal antigen
	detection
	Fungal Biomarkers
	Mycology
	Anti-HBs detection
	Blood Borne Viruses
	Blood Donor Screen
	Diagnostic Serology:
	hepatitis screen
	Hepatitis B Serology
	Hepatitis C Serology
	Hepatitis E Serology
Virology	HIV Point of Care
Virology	HIV Serology
	Immunity Screen
	Measles and Mumps
	IgG Serology
	Parvovirus B19 and
	Rubella Serology
	Respiratory Rapid: RSV
	Rubella IgG Serology
	Virus identification

Parasitology
Tel: +44 (0) 203 908 1371
Email: parasite@ukneqas.org.uk
Legal Entity: Public Health England
Services Available
Blood Parasitology
Faecal Parasitology
Malaria Rapid
Molecular Diagnosis of Malaria
Parasite Serology
Toxoplasma Serology
Molecular detection of Faecal Parasites
Teaching for Blood and Faecal Parasites

Genon	nics ana kep	ro	anctive 2cie	nce
Reproducti	ve Science		Genomics	continued
Tel: +44 (0) 161 276 6437 Email: repscience@ukned	qas.org.uk			Acquired array (CLL and MDS)
Legal Entity: Manchester dation Trust	University NHS Foun-			Acute Lymphoblastic Leukaemia (ALL)
Services A	Available			Chronic Lymphocytic
		Chronic Lym		Chronic Lymphocytic Leukaemia (CLL)
Embryology: Embryo Mor notation	phology, Time Lapse An-			IGHV mutation status*
Genomics	(GenQA)		Haematological Neoplasms	Chronic Lymphocytic Leukaemia (CLL) TP53 mutation analysis*
, ,				Haematological Technical FISH
Email: info@genqa.org Legal Entity: Oxford Unive	ersity Hospitals NHS Foun-			Haematological Translocations*
				Lymphoma
Services A	Available Cardiovascular			Lymphoma Technical NGS*
	Disorders	-		Myeloid Disorders
	Dysmorphology			Myeloma
Services A	Monogenic Disorders			Tissue-i
	Oncogenetics			BRCA and HRR variant classification
	Genetic Counselling*	NHS Foun- Motility, Con- ime Lapse An- Base Ba		COVID-19 Care
	Chromosome Breakage syndromes			Home Swabbing COVID-19 Laboratory
	Disorders of Sexual Development*	Individual Competency		, Testing
	Induced Pluripotent stem (IPS) cells		Assessment	COVID-19 Direct LAMP assay
	Infertility*	-		COVID-19 LamPORE assay
	Microdeletion syndromes*			COVID-19 High throughput Direct RT-
Constitutional - Postnato	Postnatal constitutional Copy			LAMP assay
	Number Variant (CNV) detection			Aminoglycoside- Induced deafness*
	Recurrent miscarriage karyotyping		Molecular Genetic	Ataxia & Hereditary Spastic Paraplegia (HSP)
	Severe Developmental		Disorders	Cardiac Disorders
Services Av Andrology: Semen Analysis centration & Morphology) Embryology: Embryo Morphology: Embryo Morphology notation Genomics (Tel: Oxford Office: +44 (0) 1865 Edinburgh Office: +44 (0) 1 Email: info@genqa.org Legal Entity: Oxford Universidation Trust/ NHS Lothian Services Av	Delay*			Charcot Marie Tooth

Sex chromosome

disorders karyotyping

disease (CMT) and

related sensory and

motor neuropathies

^{*} Pilot/Non Accredited

Genomi	cs continued
O C I I O I I I	Cystic Fibrosis (CF) and
	CFTR-related disorders
	Fuella and Discount and
	Epilepsy Disorders
	Eye Disorders
	Familial Colorectal Cancer and Polyposis
	, ·
	Familial Hypercholesterolaemia
	(FH)
	Familial endocrine
	tumour predisposition
	disorders
	F 'l V l
	Fragile X syndrome and FMR1-related disorders
	TWIRT-TOTATE A GISOTACTS
	Gastrohepatology
	disorders*
	Hereditary Breast and
	Ovarian Cancer
	(HBOC) disorders
	Huntington disease
Molecular Genetic	(HD)
Disorders	Hypotonic Infant
	Imprinting disorders
	Inborn Errors of
	Metabolism (IEM)
	Linkage analysis
	Mitochondrial and
	POLG-related
	disorders Muscular Dystrophics
	Muscular Dystrophies
	Neurodegenerative Disorders
	Neurofibromatosis type
	1 and Rasopathies
	Osteogenesis
	Imperfecta (OI)
	Prediction of 5-
	Fluorouracil Toxicity
	(DPYD)*
	Primary
	Immunodeficiency
	disorders (PID)

Genomics	continued	
Molecular Genetic Disorders	Renal Disorders	
	Respiratory disorders	
	Skeletal Dysplasias	
	X-Inactivation	
Molecular Newborn Screening	Cystic Fibrosis (CF) Molecular Newborn screening (bloodspots)	
	MCADD Molecular Newborn Screening (bloodspots)	
	SCID Molecular Newborn Screening (bloodspots)*	
	BRCA testing for ovarian cancer - somatic	
	BRCA testing for ovarian, breast, prostate and pancreatic cancer - germline	
	Breast cancer (PIK3CA testing)*	
	Breast cancer - Tumour Expression profiling*	
Molecular Pathology	Central Nervous System (CNS) Tumours	
	Circulating free (cf) DNA testing in lung cancer	
	Gastrointestinal stromal tumours (GIST)	
	Microsatellite instability testing (MSI)	
	Molecular Tissue Identification	
	NTRK fusions*	
	Neuroblastoma*	

Genomics continued			
	Prostate cancer (HRR		
	testing)*		
	Renal Tumours*		
	Sarcoma		
	Thyroid Cancer*		
	Colorectal cancer - Extended MMR		
	Colorectal cancer - MMR		
Molecular Pathology	Colorectal cancer - core		
	Lung cancer - EGFR only		
	Lung cancer- comprehensive		
	Lung cancer - core		
	Lung cancer - fusions		
	Melanoma		
	Preimplantation genetic testing (PGT) for aneuploidy		
Preimplantation Genetic Testing (PGT)	Preimplantation genetic testing (PGT) for chromosomal rearrangements		
	Preimplantation genetic testing (PGT) of Blastomere (FISH)		
	Preimplantation genetic testing (PGT) of Polar Body		
	Preimplantation Genetic Testing (PGT) for monogenic disorders		
Prenatal Testing	Maternal cell contamination (MCC) & sexing		
	Non-invasive prenatal testing (NIPT) for aneuploidies		

Genomics continued			
Prenatal Testing	Non-invasive prenatal testing (NIPT) for common microdeletions*		
	Non-invasive prenatal testing (NIPT) for sexing		
	Pregnancy loss (G-banding)		
	Pregnancy loss (Molecular methods)		
	Prenatal constitutional Copy Number Variant (CNV) detection		
	Prenatal karyotyping		
	Rapid Prenatal Testing for common aneuploidies		
	DNA Quantification*		
Sample Handling	DNA extraction from formalin-fixed paraffin -embedded (FFPE) tissue		
	DNA extraction from fresh frozen (FF) tissue*		
	DNA extraction from saliva*		
	DNA extraction from venous blood		
Technical (Next Generation Sequencing)	Next Generation Sequencing (NGS) germline		
	Next Generation Sequencing (NGS) somatic WES/WGS*		
	Next Generation Sequencing (NGS) somatic panel		

Genomics continued

Exome Sequencing Data Interpretation*

ISCN (International System for Human Cytogenomic Nomenclature) (Classification Only)*

Pathogenicity of germline postnatal copy number variants (CNV) (Classification Only)*

Variant Classification

Pathogenicity of germline sequence variants

(Classification & Interpretation)

Pathogenicity of germline sequence variants (Classification only)

Pathogenicity of prenatal copy number variants (Classification only)*

Pathogenicity of somatic sequence variants (Classification only)*

Variant Validation

Covid-19 Programmes

Purpose	Services Available	UK NEQAS Provider
Diagnostic Programmes	Molecular Detection of Respiratory Viruses	UK NEQAS Microbiology
	Molecular Detection of SARS-CoV-2	UK NEQAS Microbiology
	SARS-CoV-2 Antibody Detection	UK NEQAS Immunology, Immunochemistry & Allergy
	SARS-CoV-2 Antigen*	UK NEQAS Microbiology
	Activated Partial Thromboplastin Time (APTT)	UK NEQAS Blood Coagulation
	B-Type Natriuretic Peptide (BNP)	UK NEQAS Cardiac Markers
	C-Reactive Protein	UK NEQAS Immunology, Immunochemistry & Allergy
	Point of Care CRP*	UK NEQAS Immunology, Immunochemistry & Allergy
	D-Dimer Assay (within Screening Tests)	UK NEQAS Blood Coagulation
	Ferritin (within Haematinics EQA Programme)	UK NEQAS Birmingham Quality
	Full Blood Count (FBC)	UK NEQAS Haematology
Treatment & Monitoring Programmes	Heparin Induced Thrombophilia (HIT) Screening (within Screening Tests)	UK NEQAS Blood Coagulation
	Interleukin-6 (IL-6)*	UK NEQAS Immunology, Immunochemistry & Allergy
	Monitoring Anticoagulation with Low Molecular Weight Heparin (UFH) (within Screening Tests)	UK NEQAS Blood Coagulation
	NT-pro BNP	UK NEQAS Cardiac Markers
	Plasma Viscosity	UK NEQAS Haematology
	Procalcitonin	UK NEQAS Immunology, Immunochemistry & Allergy
	Prothrombin Time for Diagnosis (PT/Diagnostic) (PTD)	UK NEQAS Blood Coagulation
	Routine Biochemistry	UK NEQAS Birmingham Quality
	Cardiac Troponin I	UK NEQAS Cardiac Markers
	Cardiac Troponin T	UK NEQAS Cardiac Markers
	Covid-19 Care Home Swabbing	UK NEQAS Genomics
	Covid-19 Laboratory Testing	UK NEQAS Genomics
Educational Modules	Covid-19 LAMP Assay	UK NEQAS Genomics
Educational Modules	Covid-19 LamPORE Assay	UK NEQAS Genomics
	Covid-19 High Throughput Direct RT -LAMP Assay	UK NEQAS Genomics

Point of Care Testing

Services Available	UK NEQAS Provider
Activated Clotting Time (ACT)	UK NEQAS Blood Coagulation
B-Type Natriuretic Peptide (BNP)	UK NEQAS Cardiac Markers
Cardiac Troponin I	UK NEQAS Cardiac Markers
Cardiac Troponin T	UK NEQAS Cardiac Markers
CKMB	UK NEQAS Cardiac Markers
Clinical Chemistry	UK NEQAS Birmingham Quality
CoaguChek XS, CoaguChek XS Plus, CoaguChek XS Pro & Pro II (PT/INR)	UK NEQAS Blood Coagulation
C-Reactive Protein (CRP)*	UK NEQAS Immunology, Immunochemistry & Allergy
D-Dimer	UK NEQAS Blood Coagulation
D-Typing	UK NEQAS BTLP
Fungal Biomarkers	UK NEQAS Microbiology
Glycated Haemoglobins	UK NEQAS Birmingham Quality
Haemoglobin only	UK NEQAS Haematology
HIV	UK NEQAS Microbiology
Malaria Rapid	UK NEQAS Parasitology
Molecular Detection of Respiratory Viruses	UK NEQAS Microbiology
Myoglobin	UK NEQAS Cardiac Markers
NT-pro BNP	UK NEQAS Cardiac Markers
Pregnancy Testing	UK NEQAS Edinburgh
Respiratory Rapid: RSV	UK NEQAS Microbiology
Thromboelastometry and Thromboelastography	UK NEQAS Blood Coagulation
Urinary Antigens	UK NEQAS Microbiology
Urine Dipsticks	UK NEQAS Birmingham Quality
Viral Gastroenteritis (also suitable for antigen testing)	UK NEQAS Microbiology

#		Antibody to Fungal & Related	11
1st Trimester (Down's syndrome and	7	Antigens Antifungal Assays	6
Trisomies T13 and T18)	/	Antifungal Susceptibility	13
1st Trimester (Down's syndrome using dried blood spots)	7	Anti-HBs detection	13
2nd Trimester (Down's syndrome)	7	Antimicrobial Susceptibility	13
2nd Trimester (Neural tube defects)	7	Antithrombin Antigen and Activity	8
β2 Microglobulin	, 11	Assays	
		Assays (Blood Coagulation)	8
Α		Ataxia & Hereditary Spastic Paraplegia (HSP)	14
A ₁ typing	9	Autoimmunity	11
AAFB Microscopy	13	Automated Counting & Related	
Abnormal Haemoglobins	9	(Haematology)	9
ABO & D Grouping	9	Automated Differential Leucocyte	9
ABO Titration ABO Titration (ABOT)	9	Count	
Acetylcholine Receptor Antibodies	11	В	
	14	Bacteriology	13
Acquired array (CLL and MDS) Activated Clotting Time (ACT)	8, 19	BCR-ABL1 and AML Translocation Identification	10
Activated Partial Thromboplastin Time (APTT)	8, 18	BCR-ABL1 Kinase Domain Variant (Mutation) Status	10
Activated Protein C Resistance Assay	8	BCR-ABL1 Major Quantification	10
Acute Lymphoblastic Leukaemia	14	BCR-ABL1 Minor Quantification*	10
(ALL) Acute Myeloid Leukaemia and		Blood Borne Viruses	13
Myelodysplastic Syndrome Gene	10	Blood Donor Screen	13
Panels*		Blood Films for Morphology	9
ADAMTS13 assays	8	Blood Parasite Screening and Identification (Haematology)	9
AFP, CEA and hCG	7	Blood Parasitology	13
Alimentary Tract Pathology (GIST)	5	Bone Marrow Trephine Biopsy (BMT)	5
Alkaline Phosphatase (ALP) Isoenzymes	11	BRAF p.Val600Glu (V600E) Mutation Status for Hairy Cell Leukaemia	10
Allergen Component Testing	11	BRCA and HRR variant classification	14
Allergen Specific IgE	11	BRCA testing for ovarian cancer -	15
Allergy and Immunodeficiency	11	somatic	10
Alpha 1 Antitrypsin and Phenotype Identification	11	BRCA testing for ovarian, breast, prostate and pancreatic cancer - germline	15
Aluminium in Water/Dialysis Fluid (Educational)	6	Breast cancer - Tumour Expression	15
Aminoglycoside-Induced deafness*	14	profiling* Breast cancer (PIK3CA testing)*	15
ANCA/GBM Antibodies	11	Breast HER2 ISH (Interpretive and	
Andrology: Semen Analysis (Sperm		Technical)	5
Motility, Concentration & Morphology)	14	Breast Pathology (HER2 IHC)	5
Antenatal Antibody Titration (ANT)	9	Breast Pathology (Hormonal Receptors ER & PR)	5
Antibiotic Assays	6	Breast Pathology (Hormonal	Г
Antibodies to Nuclear and Related	11	Receptors ER only)	5
Antigens		B-Type Natriuretic Peptide (BNP)	6, 18, 19
Antibody Identification	9	Bullous Dermatosis Antibodies	11
Antibody Screening	9		

С		C-Peptide	6
C. trachomatis and N. gonorrhoeae	13	C-Reactive Protein (CRP)	11, 18, 19
C1 Esterase Inhibitor and Functional	Crossmatchina		9
Complement Assays	11	Crossmatching by Flow Cytometry	
Cardiac Disorders	14	Cryoproteins (image based)	
Cardiac Markers Schemes (Point of Care)	6	Cryptococcal antigen detection	13
Cardiac Markers Schemes (Laboratory)	6	CSF Haem Pigments	11
Cardiac Troponin I	6, 18, 19	CSF IgG Oligoclonal Bands	11
Cardiac Troponin T	6, 18, 19	CSF Proteins and Biochemistry	11
Cardiovascular Disorders	14	CSF β2 Transferrin/Beta Trace Protein	11
CD34+ Stem Cell Enumeration	10	Cystic Fibrosis (CF) and CFTR-related	15
Central Nervous System (CNS) Tumours	15	disorders	
Cerebrospinal Fluid (CSF) Immunophenotyping*	10	Cystic Fibrosis (CF) Molecular Newborn screening (bloodspots)	15
		Cytochemistry: Haemosiderin staining	9
Charcot Marie Tooth disease (CMT) and related sensory and motor neuropathies	14	Cytopathology	5
,	1.4	Cytotoxic Crossmatching	12
Chromosome Breakage syndromes Chronic Lymphocytic Leukaemia (CLL)	14 14	D	
Chronic Lymphocytic Leukaemia (CLL)		D-Dimer	8, 18, 19
IGHV mutation status*	14	Detection of cells coated with C3d	9
Chronic Lymphocytic Leukaemia (CLL) TP53	3 14	Detection of cells coated with IgG	9
mutation analysis*	14	Diabetic Markers	11
Chronic Lymphocytic Leukaemia Gene	10	Diagnostic Cytopathology	5
Panels*		Diagnostic Programmes (Covid-19)	18
Circulating free (cf) DNA testing in lung	15	Diagnostic Serology: hepatitis screen	13
cancer CKMB	6, 19	Digital Blood Film Morphology for CPD	9
Clinical Chemistry	6, 19	Digital Diagnostic Ultrastructural	5
Clinical Genetics	14	Pathology*	
Clostridioides Difficile (prev. Clostridium		Digital Interpretative Diagnostic Cytopathology	5
Difficile)	13	Digital Pathology Imaging*	5
CMV DNA Quantification	13	Direct Antiglobulin Testing (DAT)	9
CoaguChek XS, CoaguChek XS Plus,	8, 19	Direct Immunofluorescence (DIF)	5
CoaguChek XS Pro & Pro II (PT/INR)		Direct Oral Anticoagulant (DOAC) Assay*	8
Coeliac Disease Antibodies	11		
Colorectal cancer - core	16	Disorders of Sexual Development*	14
Colorectal cancer - Extended MMR	16	DNA Diagnostics for Haemoglobinopathies	9
Colorectal cancer - MMR	16	DNA extraction from formalin-fixed	- <i>.</i>
Colorectal Cancer Screening	6	paraffin-embedded (FFPE) tissue	16
Community Medicine	13	DNA extraction from fresh frozen (FF)	16
Constitutional Postnatal	14	tissue*	10
Copper and Iron in Solid Matrices e.g. Liver	6	DNA extraction from saliva*	16
(Educational)	14, 18	DNA extraction from venous blood	16
COVID-19 Care Home Swabbing COVID-19 Direct LAMP assay	14, 18	DNA HLA Typing at 1st Field	12
COVID-17 Direct LAWI assay COVID-19 High throughput Direct RT-LAMP	14, 10	Resolution	
assay	14, 18	DNA HLA Typing to 2nd or 3rd Field Resolution	12
COVID-19 Laboratory Testing	14, 18	DNA Quantification*	16
COVID-19 LamPORE assay	14, 18	DOAC assays	8
		Dysmorphology	14

-		Full Blood Count (FBC)	9, 18
EBV DNA Quantification			13, 19
	ıJ	Fungal Biomarkers	13, 19
Educational Cross Match (Histocompatibility & Immunogenetics)	12	FVIII treatment monitoring FXIII assays	
Educational Interpretive Clinical Scenarios	12	G	
(Histocompatibility & Immunogenetics)		G6PD Screen and Assay	9
Educational Scheme (Histocompatibility &	12	Ganglioside Antibodies	11
Immunogenetics) Embryology: Embryo Morphology, Time Lapse		Gastric HER2 IHC	5
Annotation	14	Gastrin	6
Emicizumab assays	8	Gastrohepatology disorders*	15
Enhanced Liver Fibrosis (ELF) Score	7	Gastrointestinal stromal tumours	15
Erythropoietin	6	(GIST)	
ESR	9	General Autoimmune Serology	11
Exome Sequencing Data Interpretation*	17	General Bacteriology	13
Extended Red Cell Phenotyping (ERP)	9	General Pathology	5
Epilepsy Disorders	15	Genetic Counselling*	14
Eye Disorders	15	Genetics of Heritable Bleeding and Thrombotic disorders	8
Educational Modules (Covid-19)	18	Genital pathogens	13
F		GFR Estimations (Creatinine, Cystatin C & eGFR)	6
Factor II, V, VII VIII, IX, X, XI, XII Assays	8	Glycated Haemoglobins	6, 19
Faecal Haemoglobin Faecal Markers	6		
Faecal Markers Faecal Pancreatic Elastase	6	H	/ 10
Faecal Parasitology	13	Haematinics	6, 18
Faecal Pathogens	13	Haematological Malignancy Bone Marrow Aspirate Assessment*	10
•	15	Haematological Neoplasms	14
Familial Colorectal Cancer and Polyposis	13	Haematological Technical FISH	14
Familial endocrine tumour predisposition disorders	15	Haematological Translocations*	14
Familial Hypercholesterolaemia (FH)	15	Haemoglobin only	9, 19
Ferritin	18	Haemoglobinopathy	9
Fetomaternal Haemorrhage (FMH)	9	HBV DNA Quantification	13
FIB-4 Score	7	Head and Neck Pathology (Oral, ENT & Combined)	5
Fibrinogen Evaluation (Clauss method)	8	Heparin Assay (HA) - anti-Xa	8
FIX treatment monitoring	8	Heparin Dosage Assessment (HDA)	
Flow Cytometry	10	by APTT	8
FLT3 Mutation Status Fluids	10 6	Heparin Induced Thrombophilia (HIT) Screening	18
FMH Quantification	9	Hepatitis B Serology	13
FMH Screening	9	Hepatitis C RNA Detection	13
Fragile X syndrome and FMR1-related	1.5	Hepatitis C Serology	13
disorders	15	Hepatitis E Serology	13
Frozen Section (Companion Scheme to Specialist Techniques and Neuropathology)	5	Hereditary Breast and Ovarian Cancer (HBOC) disorders	15
Fructosamine	6	HFE Typing	12
FSH, LH, AMH, Prolactin and Growth		HIT screening/assays	8
Hormone	7	HIV 1 RNA Quantification	13
		HIV Point of Care	13, 19

ontinued		
	13	
	12	
HLA Antibody Specificity Analysis	12	
HLA Genotyping for Coeliac and other HLA	12	
associated diseases	12	
,, ,	12	
,, , , , , , , , , , , , , , , , , , , ,	12	
3	12	
	6	
lomocysteine assays	8	
IPA Antibody Detection/Specification	12	
IPA Genotyping	12	(P
luntington disease (HD)	15	Leuko
Hyaluronic acid	7	Linkage
Hypotonic Infant	15	Lipid Inves
		Liquid Newb Screening
	1.1	Liver Fibrosis
gG Subclasses	11	Low Level Let
gH/TCR Clonality Status	10	Lung cancer -
mmune Monitoring	10	Lung cancer -
mmunity Screen	13 11	Lung cancer -
mmunochemistry	5	Lung cancer -
mmunocytochemistry		Lupus anticoa
mmunosuppressants ndividual Competency Assessment	6	Lymphoid Path
(Genomics)	14	Lymphoma
nduced Pluripotent stem (IPS) cells	14	Lymphoma Tec
nfectious Mononucleosis*	9	Lymphoplasma
nfertility*	14	Waldenström M
n-Situ Hybridisation	5	
nsulin	6	M
nsulin-like Growth Factor 1	6	Malaria Rapid
nsulin-like Growth Factor Binding Protein-3	6	Malaria Rapid Di (Haematology)
nterferon Gamma Release Assays	11	Manual WBC Dif
(Mycobacterium tuberculosis) IGRA TB		Maternal cell cor
nterleukin-6 (IL-6)*	11, 18	sexing
nterpretative Comments in Clinical Chemistry	6	Maternal Serum S
nterpretative: HFE Genotype and Hereditary Haemochromatosis	12	MCADD Molecule (bloodspots)
nterpretive HLA Genotype	12	Measles and Mur
ntrinsic Factor Antibodies	6	Mega Block (Coi
SCN (International System for Human		Specialist Techn
Cytogenomic Nomenclature) (Classification	17	Neuropathology
Only)*	1.5	Melanoma
mprinting disorders	15	Mercury, Cadn
nborn Errors of Metabolism (IEM)	15	Chromium, Co Manganese, N Selenium in Uri

M continued		Myeloproliferative Neoplasm Gene Panels*	10
Microdeletion syndromes*	14	Myoglobin	6, 19
Microsatellite instability testing (MSI)	15		11
Minimal Residual Disease for ALL by Flow	Myositis Associated Antibodies 10		11
Cytometry		N	
Minimal Residual Disease for AML by Flow	10	Neuroblastoma*	15
Cytometry* Minimal Residual Disease for CLL by Flow		Neurodegenerative Disorders	15
Cytometry*	10	Neurofibromatosis type 1 and Rasopathies	15
Minimal Residual Disease for Plasma Cell Myeloma by Flow Cytometry*	10	Neuropathology (Cellular Pathology Technique)	5
Mismatch Repair (MMR) Proteins	5	Neuropathology (Immunocytochemistry)	5
Mitochondrial and POLG-related		Neuropathology Interpretative EQA Scheme	
disorders	15	Newborn Screening	6
Mk-4 at endogenous concentrations in	9	Newborn Sickle Screening	9
human serum	,	Next Generation Sequencing (NGS)	
MK-7 at endogenous concentrations in	9	germline	16
human serum Mohs' Procedure	E	Next Generation Sequencing (NGS) somatic	16
	5	panel	10
, , , , , , , , , , , , , , , , , , , ,	10	Next Generation Sequencing (NGS) somatic	16
Molecular (Microbiology)	13	WES/WGS*	10
Molecular Detection Mycobacteria	13	Non-invasive prenatal testing (NIPT) for	16
Molecular detection of Faecal Parasites	13	aneuploidies	
Molecular detection of HEV RNA	13	Non-invasive prenatal testing (NIPT) for common microdeletions*	16
Molecular Detection of HPV	13	Non-invasive prenatal testing (NIPT) for	
, ,	13, 18, 19	sexing	16
Molecular Detection of SARS-CoV-2	13, 18	Non-Small Cell Lung Cancer (NSCLC) ALK	_
Molecular Detection of Viruses in CSF	13	IHC	5
Molecular Diagnosis of Malaria	13	Non-Small Cell Lung Cancer (NSCLC) ALK	5
Molecular Genetic Disorders	14, 15	ISH* Non-Small Cell Lung Cancer (NSCLC) PD-L1	
Molecular Genetic Disorders	15	IHC*	5
(Factor V Leiden)	8	Non-Small Cell Lung Cancer (NSCLC) ROS1 FISH*	5
Molecular Newborn Screening	15	Non-Small Cell Lung Cancer (NSCLC) ROS1	5
Molecular Pathology Molecular Tissue Identification	15, 16 15	IHC*	3
Monitoring Anticoagulation with Low	15	NPM1 Mutation Status	10
Molecular Weight Heparin (UFH)	18	NT-proBNP	6, 18, 19
Monoclonal Protein Identification	11	NTRK fusions*	15
Monogenic Disorders	14		
Morphology Related	9	0	2.4
MRSA Screening	13	Oncogenetics	14
	5	Oncology	11
Muscular Dystrophies	15	Osteogenesis Imperfecta (OI)	15
Mycobacteria Culture	13	Other Programmes (Blood Coagulation)	8
Mycology	13	Other Services Supplementary to EQA (BTLP)	9
Myelin Associated Glycoprotein IgM Antibodies (MAG)	11		
Myeloid Disorders	14		

P		Preimplantation genetic testing (PGT) of		
Paediatric Acute Leukaemia	Preimplantation genetic testing (PGT) of Blastomere (FISH)		16	
Translocations	10	Preimplantation genetic testing (PGT) of		
Paediatric Bilirubin	6	Polar Body		
Paraneoplastic Antibodies	11	Prenatal constitutional Copy Number		
Parasite Serology	13	Variant (CNV) detection		
Paroxysmal Nocturnal Haemoglobinuria	10	Prenatal karyotyping	16	
Parvovirus B19 and Rubella Serology	13	Prenatal Testing	16	
Pathogenicity of germline postnatal copy	. –	Pre-Transfusion Testing (PTT)	9	
number variants (CNV) (Classification	17	Primary Immunodeficiency disorders (PID)	15	
Only)* Pathogenicity of germline sequence		Procalcitonin	11, 18	
variants (Classification & Interpretation)	17	Prostate cancer (HRR testing)*	16	
Pathogenicity of germline sequence	1.7	Prostate Specific Antigen (PSA)	11	
variants (Classification only)	17	Protein C Antigen and Activity Assay	8	
Pathogenicity of prenatal copy number	17	Protein S Activity Assay	8	
variants (Classification only)*	17	Protein S Total and Free Antigen Assay	8	
Pathogenicity of somatic sequence variants (Classification only)*	17	Prothrombin Time (PT)/INR (Venous/ Capillary methods)	8	
Peptide I	7	Prothrombin Time for Diagnosis (PT/	8, 18	
Peptide II	7	Diagnostic) (PTD)		
Phospholipase A2 Receptor Antibodies (PLA2R)	11	PTH, ACTH and Calcitonin	7	
Phospholipid Antibodies	11	Q		
Pilot Schemes (BTLP)	9	Quantitative Amino Acids	6	
PIVKA-II (undercarboxylated prothrombin)	•	Quantitative Factor VIII Inhibitor	8	
at endogenous concentrations in human serum	9			
Placental Growth Factor (PLGF)	7	R		
Plasma Viscosity	, 9, 18	Rapid Prenatal Testing for common	16	
Plasminogen Assay	8	aneuploidies	10	
POCT (Blood Coagulation)	8	Recurrent miscarriage karyotyping	14	
POCT D-dimer	8, 19	Red Cell Enzyme		
Point of Care CRP	11, 18, 19	18. 19 Red Cell Genotyping (RCG)		
Point of Care D typing (POCTD)	9, 19	Renal Biopsy Pathology		
Postnatal constitutional Copy Number		Renal Disorders		
Variant (CNV) detection	14	Renal Tumours*		
Post-Stem Cell Transplant Chimerism	10	Respiratory disorders		
Monitoring	10	Respiratory Rapid: RSV	13, 19	
Prediction of 5-Fluorouracil Toxicity	15	Reticulocyte Count	9	
(DPYD)* Pre-eclampsia Markers*	7	Rh and K Phenotyping	9	
Pregnancy loss (G-banding)	16	RhD typing	9	
Pregnancy loss (Molecular methods)	16	Rubella IgG Serology	13	
Pregnancy Testing	7, 19	S		
Preimplantation Genetic Testing (PGT)	16	Sample Handling (Genomics)	16	
Preimplantation genetic testing (PGT) for		Sarcoma	16	
aneuploidy	16	SARS-CoV-2 Antibody Detection	11, 18	
Preimplantation genetic testing (PGT) for	1 /	SARS-CoV-2 Antigen*	18	
chromosomal rearrangements	16	SCID Molecular Newborn Screening		
Preimplantation Genetic Testing (PGT) for monogenic disorders	16	(bloodspots)*	15	

\$ continued	
Screening Tests (Blood Coagulation)	8, 18
Serum Indices (HIL)	6
Severe Developmental Delay*	14
Sex chromosome disorders karyotyping	14
SFIt-1	7
SFIt-1: PLGF ratio	7
Sickle Screening only	9
Skeletal Dysplasias	15
Specialist Techniques (Cellular Pathology Technique)	5
Specific Microbial Antibodies	11
Specific Proteins	6
Steroid Hormones	6
Sweat Testing	6
Syphilis Serology	13
T	
Teaching for Blood and Faecal	10
Parasites	13
Technical (Next Generation Sequencing)	16
Thrombin Time (TT)	8
Thromboelastometry and	
Thromboelastography	8, 19
Thyroglobulin	6
Thyroid Cancer*	16
Thyroid Hormones	6
Tissue Diagnostics (Cellular Pathology Technique)	5
Tissue inhibitor of metalloproteinase I (TIMP-I)	7
Tissue-i	14
Total IgE	11
Toxicology (TOX) and Therapeutic Drug Monitoring (TDM)	6
Toxoplasma Serology	13
Training, Assessment and Competency Tool (TACT)	9
Transmission Electron Microscopy (TEM)	5
Treatment & Monitoring Programmes (Covid-19)	18
Triple Negative Breast Cancer (TNBC) PD-L1 IHC*	5
Tryptase	11
Tumour Markers	7
Tumour Markers (CA Series)	11
Type III Procalloagen Peptide (PIIINP)	7

U	
Ultrasensitive C-Reactive Protein	11
Ultrasensitive PSA	11
Urinary Antigens	13, 19
Urinary Catecholamines & Metabolites	6
Urinary hCG (Qualitative and Quantitative)	7
Urine Chemistries	6
Urine Dipsticks	6, 19
V	
Variant Classification (Genomics)	17
Variant Validation (Genomics)	17
Viral Gastroenteritis (also suitable for antigen testing)	13, 19
Virology	13
Virus identification	13
Vitamin Assays (Carotene, Vitamin A & E)	6
Vitamin D	6
Vitamin K1 at endogenous concentrations in human serum	9
Vitamin K1, 2, 3-epoxide at endogenous concentrations in human serum	9
Von Willebrand Factor Antigen: Assay	8
VWF:CB (collagen-binding) Assay	8
VWF:RCo (activity) Assay	8
W, X, Y & Z	
X-Inactivation	15
Zinc, Copper, Aluminium, Selenium, Chromium and Cobalt in Serum	6

Current Participating Countries

Afganistan Falkland Islands Lesotho Reunion Liechtenstein Albania Romania Fiji **Finland** Lithuania Angola Russia Argentina France Luxembourg Rwanda Armenia French Polynesia Saudi Arabia Macau Aruba French Guiana Madagascar Senegal **Australia** Gambia Malawi Serbia **Austria** Sierra Leone Georgia Malaysia Azerbaijan Germany Martinique Singapore Bangladesh Ghana Mali Sint Maarten Gibraltar Malta Slovak Republic **Belarus** Belgium Greece Mauritius Slovenia Benin Greenland Mexico South Africa Bolivia Guadeloupe Monaco Spain Bosnia and Herzegovina Guatemala Mongolia Sri Lanka Botswana Guinea Sudan Montenegro Haiti Swaziland Brazil Morocco Bulgaria **Honduras** Mozambique Sweden Burkina Faso Hong Kong Namibia Switzerland Cambodia Hungary Nepal Taiwan Cameroon Iceland **Netherlands** Tajikistan India New Caledonia Tanzania Canada Indonesia New Zealand Central Africa Republic **TCHAD** Chile Iran Nicaragua Thailand China Iraq Niger Togo Colombia Ireland Nigeria Tunisia Congo, Democratic Republic Israel Turkey Norway **Turkmenistan** Costa Rica Italy Oman Cote d'Ivoire Jamaica Pakistan Uganda Croatia Japan UK Panama Curacao Jordan Papua New Guinea Ukraine Kazakhstan Paraguay **United Arab Emirates** Cyprus Czech Republic Kenya Peru Uruguay Denmark Kingdom of Bahrain Philippines USA Dominican Republic Kosovo Poland **Uzbekistan** Kuwait Venezuela Ecuador Portugal Egypt Kyrgyzstan Qatar Vietnam El Salvador Lao PDR Republic of Korea Yemen Estonia Latvia Republic of Macedonia Zambia Ethiopia Lebanon Republic of Moldova Zimbabwe

UK NEQAS Contact Information

Centre	Telephone	Email
Birmingham Quality	+44 (0) 121 414 7300	birminghamquality@uhb.nhs.uk
Blood Coagulation	+44 (0) 114 267 3300	neqas@coageqa.org.uk
Blood Transfusion Laboratory Practice (BTLP)	+44 (0) 1923 217 933	btlp@ukneqas.org.uk
Cardiac Markers	+44 (0) 141 440 2888	info@ukneqas-cm.org.uk
Cellular Pathology Technique	+44 (0) 191 816 1030	cpt@ukneqas.org.uk
Edinburgh Peptide Hormones	+44 (0) 131 242 6885	ukneqas@ed.ac.uk
Genomics	Oxford: +44 (0) 1865 857 644 Edinburgh: +44 (0) 131 242 6898	info@genqa.org
Guildford Peptide Hormones	+44 (0) 1483 571 122 ext. 3611	rsch.peptideeqa@nhs.net
Haematology	+44 (0) 1923 217 878	haem@ukneqas.org.uk
Head and Neck Pathology	+44 (0) 121 371 5723	neckpath@ukneqas.org.uk
Histocompatibility and Immunogenetics	+44 (0) 1443 622 185	handi@ukneqas.org.uk
Immunocytochemistry & In-Situ Hybridisation	+44 (0) 208 187 9174	info@ukneqasiccish.org
Immunology, Immunochemistry & Allergy	+44 (0) 114 271 5715	immunology@ukneqas.org.uk
Leucocyte Immunophenotyping	+44 (0) 114 267 3600	admin@ukneqasli.co.uk
Microbiology	+44 (0) 208 905 9890	organiser@ukneqasmicro.org.uk
Neuropathology	+44 (0) 1865 234 904	neuropath@ukneqas.org.uk
Parasitology	+44 (0) 203 908 1371	parasite@ukneqas.org.uk
Reproductive Science	+44 (0) 161 276 6437	repscience@ukneqas.org.uk
Trace Elements	+44 (0) 1483 571 122 ext. 3611	rsc-tr.Guildford-EQA@nhs.net
Vitamin K	+44 (0) 207 188 6815	keqas@ukneqas.org.uk

UK NEQAS International Quality Expertise

UK NEQAS Central Office PO BOX 401 Sheffield S5 7YZ

Tel: +44 (0) 114 261 1689

centraloffice@ukneqas.org www.ukneqas.org.uk

