

Schedule of Accreditation

issued by

United Kingdom Accreditation Service

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

 7872 Accredited to ISO/IEC 17043:2023	Oxford University Hospitals NHS Foundation Trust, operating GenQA	
	Issue No: 017 Issue date: 20 March 2025	
	Level 1, The Women's Centre John Radcliffe Hospital Oxford University Hospitals NHS Foundation Trust Oxford OX3 9DU	Contact: Melody Tabiner E-Mail: melody.tabiner@genqa.org bettinaq@genqa.org Website: www.genqa.org
Proficiency Testing provided from the locations specified below		

Locations covered by the organisation and their relevant activities

<u>Location details</u>		<u>Activity</u>
GenQA Level1, The Women's Centre John Radcliffe Hospital Oxford University Hospitals NHS Trust Oxford OX3 9DU	Local contact Melody Tabiner Email: melody.tabiner@genqa.org Bettina Quellhorst Email: bettinaq@genqa.org	QMS operations
GenQA Laboratory Medicine NHS Lothian NINE, Edinburgh BioQuarter Little France Road Edinburgh EH16 4UX	Local contact Prof Sandi Deans Tel: 0131 242 6898 Email: Sandi.Deans@genqa.org	Scheme operations and laboratory services



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DETAIL OF ACCREDITATION

Materials/Products	Scheme Name/Type of Test/Properties Measured	Scheme Protocols/Procedures/ Techniques Used
		All protocols available through www.genqa.org
FFPE Tissue	Sample Handling DNA extraction from formalin fixed paraffin embedded (FFPE) tissue samples	
Fresh Tissue	DNA extraction from fresh tissue	
Blood	DNA extraction from venous blood samples	
DNA	DNA Quantification	
DNA	Technical Next Generation Sequencing (NGS) - Germline	
	Genomic and Inherited Disorders	
Case scenario or DNA	Ataxia and spastic paraplegias	
Case scenario or DNA	Cardiac disorders	
Case scenario or DNA	Charcot Marie Tooth disease and related sensory and motor neuropathies	
Case scenario or DNA	Cystic fibrosis and CFTR-related disorders	
Case scenario or DNA	Epilepsy disorders	
Case scenario or DNA	Eye disorders	
Case scenario or DNA	Familial colorectal cancer and polyposis	
Case scenario or DNA	Familial endocrine tumour predisposition disorders	
Case scenario or DNA	Familial hypercholesterolaemia	
Case scenario or DNA	Fragile X syndrome and <i>FMR1</i> -related disorders	
Case scenario or DNA	Gastroenterology and hepatology disorders	
Case scenario or DNA	Hereditary breast and ovarian cancer	
Case scenario or DNA	Huntington disease and DRPLA	
Case scenario or DNA	Inborn errors of metabolism	
Case scenario or DNA	Mitochondrial disorders	



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Materials/Products	Scheme Name/Type of Test/Properties Measured	Scheme Protocols/Procedures/ Techniques Used
<p>Case scenario or DNA Case scenario or DNA Case scenario or DNA Case scenario or DNA</p> <p>Case scenario or DNA</p> <p>Case scenario or DNA Case scenario or DNA Case scenario or DNA Case scenario or DNA Case scenario</p> <p>Images and DNA Case scenario Case scenario or DNA</p> <p>Case scenario or DNA Case scenario or DNA Case scenario Case scenario or DNA Fixed Cell or DNA DNA</p> <p>Images Case scenario or DNA</p>	<p>Genomic and Inherited Disorders</p> <p>Muscular dystrophies Neurodegenerative disorders Neurofibromatosis and rasopathies Parathyroid and calcium regulation disorders Primary immunodeficiency disorders Renal disorders Respiratory disorders Skeletal dysplasia's X-inactivation Classification and interpretation of germline SNVs and indels Chromosome instability syndromes Developmental Delay Differences in sex development (DSD) Hypotonic infant Imprinting disorders Infertility Variant validation Microdeletion syndromes Postnatal Constitutional CNV detection Postnatal Karyotyping Trio Sequencing Postnatal</p>	<p>All protocols available through www.genqa.org</p>



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	Molecular Newborn Screening	
Dried blood spots	Cystic fibrosis molecular newborn screening	
Dried blood spots	Medium chain acyl Co-A dehydrogenase deficiency molecular newborn screening	
Dried blood spots	Severe Combined Immune Deficiency molecular newborn screening (SCID)	
	Molecular Pathology	
Formalin fixed paraffin embedded (FFPE) tumour section or DNA	BRCA testing for ovarian and prostate cancer – somatic	
DNA	BRCA testing for ovarian, breast, pancreatic and prostate cancer – Germline	
Formalin fixed paraffin embedded (FFPE) tumour section or DNA or plasma	Breast cancer	
Artificial plasma	cfDNA testing for tumour biomarkers	
Formalin fixed paraffin embedded (FFPE) tumour section or DNA	CNS Tumours	
Formalin fixed paraffin embedded (FFPE) tumour section or DNA	Microsatellite instability testing	
Formalin fixed paraffin embedded (FFPE) tumour section or DNA	Molecular analysis in melanoma	
Formalin fixed paraffin embedded (FFPE) tumour section or DNA	Molecular analysis in colorectal cancer	
Formalin fixed paraffin embedded (FFPE) tumour section or DNA	Molecular analysis in gastrointestinal stromal tumours	
Formalin fixed paraffin embedded (FFPE) tumour section or DNA	Molecular analysis in lung cancer	



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Materials/Products	Scheme Name/Type of Test/Properties Measured	Scheme Protocols/Procedures/ Techniques Used
<p>Formalin fixed paraffin embedded (FFPE) tumour section or DNA</p> <p>Formalin fixed paraffin embedded (FFPE) tumour section or DNA</p> <p>Formalin fixed paraffin embedded (FFPE) tumour section or DNA</p> <p>Formalin fixed paraffin embedded (FFPE) tumour section or DNA</p> <p>Formalin fixed paraffin embedded (FFPE) tumour section or DNA</p> <p>Case scenario or DNA</p> <p>Case scenario or DNA</p> <p>DNA and cells</p> <p>Images</p> <p>DNA</p> <p>DNA</p>	<p>Molecular Pathology</p> <p>Molecular Tissue Identification</p> <p>NTRK fusions</p> <p>Renal tumours</p> <p>Sarcoma</p> <p>Thyroid cancer</p> <p>Pharmacogenomics</p> <p>Aminoglycoside ototoxicity</p> <p>Dihydropyrimidine dehydrogenase (DPYD)</p> <p>Preimplantation Genetic Testing</p> <p>Preimplantation genetic testing for Monogenic disorders (PGT-M)</p> <p>Preimplantation genetic testing of Blastomere (FISH)</p> <p>Preimplantation genetic testing for structural rearrangements (PGT-SR)</p> <p>Preimplantation genetic testing for aneuploidies (PGT-A)</p>	<p>All protocols available through www.genqa.org</p>



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	Haematological Neoplasms	
Images or DNA	Acute Lymphoblastic Leukaemia (ALL)	
Images and fixed cell suspensions or DNA	Chronic Lymphocytic Leukaemia (CLL)	
DNA and lyophilized cells	Chronic Lymphocytic Leukaemia - IGHV	
DNA and lyophilized cells	Chronic Lymphocytic Leukaemia-TP53	
Fixed cell suspensions	Haematological Technical FISH	
Images and fixed cell suspensions	Lymphoma	
FFPE		
Images or DNA	Myeloid disorders	
Images, fixed cell suspensions or DNA	Myeloma	
	Reproductive Genomics	
Case scenario or DNA	Maternal cell contamination and fetal sexing	
Case scenario or DNA	Trio Sequencing - prenatal	
Plasma/artificial plasma	Non Invasive Prenatal Testing (NIPT) for common aneuploidies	
Plasma/artificial plasma	Non Invasive Prenatal Testing (NIPT) for common microdeletions	
Plasma/artificial plasma	Non Invasive Prenatal Testing (NIPT) for fetal sexing	
Images and/or DNA	Pregnancy Loss (G-banded and molecular methods)	
DNA	Prenatal Constitutional CNV detection	
Images	Prenatal Karyotyping	
Fixed Cell Suspensions or DNA	Rapid prenatal testing for common aneuploidies	



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Materials/Products	Scheme Name/Type of Test/Properties Measured	Scheme Protocols/Procedures/ Techniques Used
Case scenario Case scenario Case scenario Case scenario	<p>Clinical Genetics</p> Clinical Genetics - Cardiovascular Genetics Clinical Genetics - Dysmorphology Clinical Genetics - Monogenic disorders Clinical Genetics - Oncogenetics	All protocols available through www.genqa.org
END		