


# Schedule of Accreditation

issued by

## United Kingdom Accreditation Service

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

 <p><b>UKAS</b> PROFICIENCY TESTING</p> <p><b>7872</b></p> <p>Accredited to <b>ISO/IEC 17043:2010</b></p>	<p><b>Oxford University Hospitals NHS Foundation Trust, operating GenQA</b></p> <p><b>Issue No: 008 Issue date: 04 June 2021</b></p>	
	<p><b>Level 1, The Women's Centre</b> <b>John Radcliffe Hospital</b> <b>Oxford University Hospitals NHS</b> <b>Foundation Trust</b> <b>Oxford</b> <b>OX3 9DU</b></p>	<p><b>Contact: Dr Ros J Hastings</b> <b>Tel: +44 (0)7825 905868</b> <b>E-Mail: ros.hastings@genqa.org</b> <b>Website: www.genqa.org</b></p>
<p>Proficiency Testing provided from the locations specified below</p>		

### Locations covered by the organisation and their relevant activities

<u>Location details</u>	<u>Activity</u>	<u>Location code</u>
<p>GenQA Level1, The Women's Centre John Radcliffe Hospital Oxford University Hospitals NHS Trust Oxford OX3 9DU</p>	<p><b>Local contact</b> Dr Ros J Hastings Contact details above</p>	<p>Scheme operations Cytogenomics</p> <p>JRH</p>
<p>GenQA Room 39 G 03 Nuffield Orthopaedic Centre Windmill Road Oxford OX3 7LD</p>	<p><b>Local contact</b> Mrs Bettina Quellhorst-Pawley Email: bettinaqp@genqa.org</p>	<p>QMS operations</p> <p>NOC</p>
<p>GenQA Laboratory Medicine NHS Lothian NINE, Edinburgh BioQuarter Little France Road Edinburgh EH16 4UX</p>	<p><b>Local contact</b> Prof Sandi Deans Tel: 0131 242 6898 Email: Sandi.Deans@ed.ac.uk</p>	<p>Scheme operations and laboratory services</p> <p>RIE</p>



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#### DETAIL OF ACCREDITATION

Materials/Products	Scheme Name/Type of Test/Properties Measured	Scheme Protocols/Procedures/ Techniques Used	Location Code
	<u>Cytogenomics EQA Schemes</u>	All protocols available through <a href="http://www.genqa.org">www.genqa.org</a>	
Online images	<b>Constitutional EQAs:</b> <b>Prenatal Karyotyping</b> Cytogenetic analysis of images and interpretation of results		JRH
Online Images	<b>Recurrent Miscarriage Karyotyping</b> Cytogenetic analysis of images and interpretation of results		JRH
Online images	<b>Sex Chromosome Disorders karyotyping</b>		JRH
Online images	<b>Pregnancy Loss (G-banded)</b> Cytogenetic analysis of images and interpretation of results		JRH
Online images	<b>PGT Blastomere FISH</b> Analysis of FISH images and interpretation of results		JRH
Online images	<b>Haemato-Oncology EQAs: Acute Lymphoblastic Leukaemia</b> Cytogenetic analysis of images and interpretation of results		JRH
Online images and Fixed cell suspensions	<b>Chronic Lymphocytic Leukaemia</b> Cytogenetic analysis of images and interpretation of results. Technical ability to obtain a result using FISH, analysis and interpretation of results		JRH / RIE



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Online images	<u>Cytogenomics EQA Schemes</u> (cont'd)  <b>Myeloid Disorders</b> Cytogenetic analysis of images and interpretation of results	All protocols available through <a href="http://www.genqa.org">www.genqa.org</a>	JRH
DNA	<b>Postnatal Constitutional CNV detection</b> Technical ability to obtain a result on an array or NGS platform, analysis and interpretation of results		JRH
Fixed cell suspensions and DNA	<b>Rapid Prenatal Testing for common aneuploidies</b> Technical ability to obtain a result using FISH, analysis and interpretation of results. Technical ability to obtain a molecular result for rapid detection of aneuploidies of chromosomes 13, 18, 21, X and Y, analysis and interpretation of results		JRH, RIE
FFPE slides	<b>CNS Tumours</b> Technical ability to obtain a result using FISH and molecular methods, analysis and interpretation of results		RIE
DNA	<b>Acquired Array (CLL and MDS)</b> Technical ability to obtain a result on an array platform, analysis and interpretation of results		RIE
DNA	<b>Prenatal CNV Detection</b> Technical ability to obtain a result on an array platform, analysis and interpretation of results		RIE



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DNA	<u>Cytogenomics EQA Schemes</u> (cont'd)	All protocols available through <a href="http://www.genqa.org">www.genqa.org</a>	RIE
Fixed Cell Suspensions / online images	<b>Pregnancy Loss (Molecular Methods)</b> (Array/MLPA/PCR/NGS) Technical ability to obtain a result using molecular methods, analysis and interpretation of results		
Online Images and Fixed cell suspension FFPE slides	<b>Myeloma</b>  Technical ability to obtain a result using FISH, analysis and interpretation of results Cytogenetic analysis of images and interpretation of results		
	<b>Lymphoma</b>  Technical ability to obtain a result using FISH, analysis and interpretation of results Cytogenetic analysis of images and interpretation of results		JRH, RIE



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DNA	<p><u>Molecular Genetics EQA Schemes:</u></p> <p><b>Molecular Genetic disorders</b>            Achondroplasia            Angelman syndrome            Alzheimer disease            Arrhythmia            Beckwith Wiedemann syndrome            BEST disease            Brugada syndrome            C9orf72 related            Frontotemporal Dementia and/or Amyotrophic Lateral Sclerosis            CADASIL            Campomelic dysplasia            Catecholaminergic polymorphic ventricular tachycardia (CPVT)            Charcot Marie Tooth disease (X-linked)            Congenital bilateral absence of the vas deferens            Connexin 26            Cystic fibrosis            Dravet syndrome            Duchenne and Becker muscular dystrophies            Emery Dreifuss muscular dystrophy            Fabrys disease            Familial adenomatous polyposis            Hereditary breast and ovarian cancer            Familial hypercholeresterolaemia            Familial medullary thyroid carcinoma            FGFR2 related skeletal dysplasias            FGFR3 related skeletal dysplasias</p>	<p>All protocols available through <a href="http://www.genqa.org">www.genqa.org</a></p>	RIE



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DNA	<p><u>Molecular Genetics EQA Schemes (cont'd)</u></p> <p><b>Molecular Genetic Disorders (cont'd)</b>            Fragile X syndrome            Fragile X associated tremor ataxia            Friedreich ataxia            Gaucher disease            Hereditary and motor sensory neuropathy            Hereditary neuropathy with liability to pressure palsies            Hereditary spastic paraplegia            Huntington disease            Hypertrophic Cardiomyopathy            Li-Fraumeni syndrome            Limb Girdle muscular dystrophies            Long QT syndrome            Lynch syndrome / Hereditary non-polyposis colon cancer            Marfan syndrome            Maternal cell contamination and sexing            MUYTH-associated polyposis            Medium chain acyl CoA dehydrogenase deficiency            Mitochondrial diseases            Molecular aneuploidy testing            Multiple endocrine neoplasia types 2A &amp; 2B            Myotonic dystrophy type 1            Neurofibromatosis type 1            Neurofibromatosis type 2            Paraganglioma (SDHD related)            Parkinson disease            Pathogenicity of sequence variants            POLG mutation testing            Prader-Willi syndrome            Retinitis pigmentosa            Retinopathies            Rett syndrome</p>	<p>All protocols available through <a href="http://www.genqa.org">www.genqa.org</a></p>	RIE



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DNA	<u>Molecular Genetics EQA Schemes (cont'd)</u>  <b>Core molecular diseases (cont'd)</b>  Schwannomatosis SDHD related paraganglioma SOX-9 associated skeletal dysplasias Spinal bulbar muscular atrophy Spinal muscular atrophy Spinocerebellar ataxias Tay Sachs disease Tuberous sclerosis Von Hippel-Lindau disease X-inactivation Variant Validation	All protocols available through <a href="http://www.genqa.org">www.genqa.org</a>	RIE
DNA	BRCA testing in ovarian cancer - germline		RIE
FFPE	BRCA testing in ovarian cancer - somatic		RIE
DNA	PGT- SR Preimplantation Genetic Testing of Blastomere/Trophectoderm for chromosomal rearrangements by NGS and/or arrays		RIE
DNA	PGT-A Preimplantation Genetic Testing of Trophectoderm and/or Blastomere for aneuploidies by NGS and/or arrays		RIE
DNA	Preimplantation Genetic Testing of Polar Bodies by NGS and/or array		RIE



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Dried blood spots	<u>Molecular Genetics EQA Schemes (cont'd):</u> <b>Dried blood spot testing</b> Cystic fibrosis (genotyping only) Medium chain acyl Co-A dehydrogenase deficiency (genotyping only)	All protocols available through <a href="http://www.genqa.org">www.genqa.org</a>	RIE
DNA and lymphocytes	<b>Pre-implantation genetic testing</b> Monogenic disorders		RIE
Formalin fixed paraffin embedded (FFPE) tumour section or DNA	<b>Molecular pathology</b> Microsatellite instability testing Molecular analysis in colorectal cancer Molecular analysis in gastrointestinal stromal tumours Molecular analysis in lung cancer Molecular analysis in melanoma		RIE
FFPE FFPE FFPE	Mismatch Repair Sarcoma Molecular Tissue Identification		
Blood	<b>DNA extraction schemes</b> DNA extraction from blood samples		RIE
FFPE tissue	DNA extraction from FFPE tissue samples		
END			