


Schedule of Accreditation

issued by

United Kingdom Accreditation Service

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

 <p>UKAS MEDICAL 21163</p> <p>Accredited to ISO 15189:2012</p>	<h3>Juno Genetics Limited</h3> <p>Issue No: 004 Issue date: 01 August 2023</p>	
	<p>Juno Genetics Limited Hayakawa Building Oxford Science Park Edmund Halley Road Oxford OX4 4GB</p>	<p>Contact: Dagan Wells Tel: +44 (0)203 743 19944 E-Mail: Dagan.Wells@junogenetics.com Website: www.junogenetics.com</p>
<p>Testing performed at the above address only</p>		

DETAIL OF ACCREDITATION

Materials/Products tested	Type of test/Properties measured/Range of measurement	Standard specifications/ Equipment/Techniques used
HUMAN TISSUES AND FLUIDS	<u>Molecular Genetics</u>	
Peripheral Blood, Saliva samples (in Oragene Media) and Products of conception	Extraction procedures: Automated extraction	Qiagen QIAamp kit and Qiacube
Peripheral Blood, Saliva samples (in Oragene Media) and Products of conception	Manual extraction	Qiagen QIAamp kit and Qiacube
Plasma	Extraction of cf DNA	Hamilton Liquid handler and VeriSeq workflow. Quantification using Spectramax Spectrophotometer



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<p>Trophectoderm cells</p> <p>DNA extracted from Trophectoderm biopsy, Peripheral Blood or Saliva (also Genomic DNA received from an external source (mother/ father/ reference))</p> <p>DNA extracted from embryo biopsy material, blood, saliva (also genomic DNA received from samples from the mother/father)</p>	<p>Preimplantation Genetic Testing for Monogenic Disorders (PGT-M)</p> <p>Direct Mutation Testing (for direct mutation site confirmation testing in PGT-M) using in house SOPs and which includes the below and analysis for all using Genemapper</p> <p>Fragment analysis Minisequencing Expansion analysis Fragile X</p> <p>Preimplantation Genetic Testing for Monogenic Disorders (PGT-M): testing for inherited single gene disorders using linkage based and direct mutation site testing</p>	<p>Using: Repli-g SC or GenomiPi kit for DNA amplification by Multiple Displacement Amplification (MDA)</p> <p>Infinium SNP array/Karyomapping kit for genotyping the SNPs.Using Illumina NextSeq 550 or Illumina NextSeq 550 DX as a scanner to "read" the genotyped SNP. Analysis using BlueFuse multi software v5.5 (Illumina) EXAM_PGT-M_Primer Design_234 EXAM_PGT_M_SOP_197</p> <p>In house and kit-based methodology and analysis using ABI 3500DX /ABI 3500</p> <p>Illumina Next Seq 550 and Illumina NextSeq 550DX with the Applied Biosystems ViiA7 qPCR instrument EXAM_PGT_M_SOP_197</p>



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<p>HUMAN TISSUES AND FLUIDS (cont'd)</p> <p>DNA extracted from Trophectoderm biopsy</p> <p>DNA extracted from Peripheral Blood and Products of Conception</p>	<p><u>Molecular Genetics</u> (cont'd)</p> <p>Preimplantation Testing for Aneuploidies (PGT-A) – Chromosome copy number assessment:</p> <p>Preimplantation Genetic Testing for Structural Rearrangements (PGT-SR)</p> <p>Products of Conception (POC) Testing – Chromosome Copy Number Assessment Including Maternal Cell Contamination Determination</p>	<p>Hot Master-taq PCR and ABI 3500XL / ABI 3500 Hot Master-taq clean-up of PCR products using ExoSAP-IT, and Primer extension using SNaPshot and ABI 3500XL/ABI 3500</p> <p>FailSafe™ one step PCR and ABI 3500XL/ ABI 3500</p> <p>AmplideX® PCR/CE FMR1 AmplideX PCR kit</p> <p>PGT-SEQ kit using Juno Genetics USA Library Preparation, Sequencing using Illumina NextSeq 550 and Illumina NextSeq 550 DX, Analysis using NexCCsv2 analysis software EXAM_PGT-Seq SOP_149</p> <p>PGT-Seq Kit using Juno Genetics USA library Preparation, Sequencing using Illumina NextSeq 550 and Illumina NextSeq 550 DX, Analysis using NexCCsv2 analysis software EXAM_PGT-Seq SOP_149 EXAM_POCSOP_187</p>



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HUMAN TISSUES AND FLUIDS (cont'd) Plasma	<u>Molecular Genetics</u> (cont'd) Non-invasive prenatal testing (NIPT)	Non-Invasive Prenatal Testing (NIPT) for aneuploidy and sexing using VeriSeq automated workflow using VeriSeq Solution v 2.– Automated Cell free DNA extraction from plasma and Library, Preparation using the Hamilton Star Liquid Handler and Paralleled sequencing on a NextSeq 550 DX. Analysis with VeriSeq VeriSeq NIPT Assay Software v2 (for 5 and 24 chromosomes) SOP EXAM_Neo and Neo24 test_SOP_269
END		