


Schedule of Accreditation

issued by

United Kingdom Accreditation Service

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

 <p>Accredited to ISO 15189:2012</p>	Invitro Genetics Ltd, trading as CooperGenomics	
	Issue No: 009 Issue date: 31 July 2018	
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Testing performed at the above address only		

DETAIL OF ACCREDITATION

Materials/Products tested	Type of test/Properties measured/Range of measurement	Standard specifications/ Equipment/Techniques used
HUMAN BODY TISSUE AND FLUIDS Biopsy samples from IVF generated embryos Biopsy samples from IVF generated embryos	<u>Molecular and Cytogenetics examination activities for the purposes of clinical diagnosis:</u> Copy number assessment for chromosome aneuploidy Detection of unbalanced structural chromosomal abnormalities via copy number analysis And Detection of chromosome aneuploidy via copy number analysis.	Documented In-house Methods and Manufacturer's instructions for analyser with specific reference to: Preimplantation Genetic Screening (PGT-A) using Illumina SurePlex whole genome amplification (WGA) and VeriSeq™ NGS; Illumina MiSeq. (SOP 59) Preimplantation Genetic Disorders (PGT-SR) by Whole genome amplification (WGA) of embryo DNA by the polymerase chain reaction (ABI 3130), followed by preparation (quantification using the Qubit) and the analysis of WGA DNA for variation of chromosome copy number by Next Generation Sequencing (NGS) using a MiSeq analyser: Illumina's VeriSeq NGS protocol to measure DNA /chromosome copy number.
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