

Solid Tissue Cytogenetic Analysis

Chromosome analysis of solid tissue samples using an appropriate investigation method (see **Analysis** below) depending upon the referral reason and type of sample sent.

Referrals / Indications

- Abnormal foetus
- Perinatal, neonatal and other infant deaths with suspected chromosomal cause
- Miscarriage (POCs) three or more consecutive pregnancy losses
- Suspected mosaicism

Analysis / Services

- Single Nucleotide
 Polymorphism (SNP array)
- Aneuploidy testing by QF-PCR
- Fluorescence in situ Hybridisation (FISH)
- Chromosome analysis by karyotyping

Sample Requirements

Products of conception, foetal skin or other solid tissue samples – send in sterile transport medium supplied by laboratory or normal saline (sterile)

Please note, products of conception or placental samples must be accompanied by a 5ml maternal blood sample in an EDTA tube

Please label samples with three identifiers and date of collection

All samples must be accompanied by a completed request form

Consent for testing and sample storage is assumed when the request is received – it is the responsibility of the referring clinician to ensure that appropriate consent has been obtained

Prices* & Turnaround Times (TAT)

*Available on request

Analysis	TAT
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SNP array testing	42cd
Aneuploidy testing (QF-PCR)	42cd
FISH analysis	42cd
Chromosome analysis	42cd



Links Orphanet

http://www.orpha.net/

OMIM

http://www.omim.org/ GeneTests

http://www.genetests.org/

Genetic Test Registry

http://www.ncbi.nlm.nih.gov/gtr/ Decipher

http://decipher.sanger.ac.uk/

Unique

www.rarechromo.org

Contact Details

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