MICROBIOLOGICAL LABORATORY



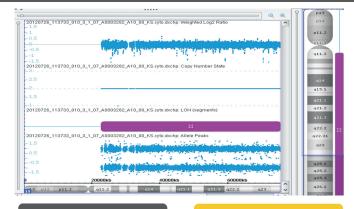


CYTOSCAN ASSAY

Results matter



Most of the miscarriages and congenital anomalies are caused by chromosomal abnormalities including aneuploidies, triploidy, uniparental disomy (UPD), etc. Traditional cytogenetic techniques such as karyotyping and FISH are limited by cell culture failure, maternal contamination, usually lack the appropriate sensitivity and increasing the turn-around time for the results. CytoScan Chromosome Microarray (CMA) analysis is a FDA approved highly controlled & validated assay protocol run on completely regulated diagnostic (Dx) system, thus avoiding plagiarism. The assay set-up applied with high density array chips, enable to perform high-resolution genome-wide DNA copy number analysis. Besides non-polymorphic markers, inclusion of SNP markers targeting whole genome provides genotyping information, enabling detection of LOH, which can also be used to detect UPDs. The combined high resolution DNA copy number data and the ability to detect gains, losses, and UPDs on a single array makes this analysis a great tool for next generation cytogenetics studies.



3% - 7%

ID Prevalence

Average age

1 in 33 Babies

Born with

Interesting Facts

4 yrs

Diagnosis by

FIRST

line Genetic Test

CytoScan Assay is the first FDA-cleared wholegenome diagnostic test to aid physicians in identifying the underlying genetic cause of developmental delay, intellectual disability, congenital anomalies, or dysmorphic features in children.

First of-its-kind diagnostic test 1

Analyze the patient's entire genome with one go 2

Exceptional performance 3 Highlights

Dual probe content with high-density SNPs 4

Streamlined data analysis 5

Test Name : CYTOSCAN

Test Code : 90312

Test Details Methodology: Microarray

Specimen :Blood/ Tissue (Fresh or Frozen)

Volume : 20 µl genomic DNA (>50 ng/µl CONC)

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