

Patient: [REDACTED]
DOB: [REDACTED] Age: 36 Gender: F
Patient Identifiers: [REDACTED],
[REDACTED]
Visit Number (FIN): [REDACTED]

Client: [REDACTED]
Physician: Doctor Unknown

ARUP Test Code: 0040203
Collection Date: 12/26/2018
Received in lab: 12/27/2018
Completion Date: 12/28/2018

Interpretation

Specimen Received
Specimen Type: Chorionic Villi
Reason for Referral: FISHCVS
Test Performed: FISHCVS

NORMAL FISH RESULT

DIAGNOSTIC IMPRESSION:

Prenatal interphase fluorescence in situ hybridization (FISH) analysis was performed with chromosome enumeration probes for 13, 18, 21, X, and Y using the Aneuvysion probe kit (Abbott Molecular). 50 interphase cells were scored for each probe.

This analysis showed no evidence for a numerical abnormality for chromosomes 13, 18, 21, X, or Y.

Sex chromosomes: XX (female)

FISH analysis performed on CVS presumes that the fetal chromosome complement is accurately reflected in the extra-embryonic tissue. There are rare examples in which the karyotype of the CVS is not consistent with that of the fetus. In addition, contamination of the sample with cells of maternal origin may result in the analysis of maternal rather than fetal chromosomes.

ISCN:
nuc ish(DXZ1x2,DYZ3x0,D18Z1x2),
(RB1x2,D21S259/D21S341/D21S342x2)

PLEASE NOTE: Interphase FISH will not detect approximately one third of prenatal chromosome abnormalities, which include mosaicism for the above chromosomes, structural abnormalities, and other numerical chromosome abnormalities. Therefore, routine cytogenetic analysis or genomic microarray is recommended for the final interpretation. If either test is being performed, it will be reported separately.

This result has been reviewed and approved by [REDACTED], MD, FACMG

Test developed and characteristics determined by ARUP Laboratories. See Compliance Statement C: aruplab.com/CS



Patient: [REDACTED]
ARUP Accession: 18-360-401378
[REDACTED]