

Client: Example Client ABC123 123 Test Drive Salt Lake City, UT 84108 UNITED STATES

Physician: Doctor, Example

## **Patient: Patient, Example**

**DOB** 8/22/1991 Gender: Female

**Patient Identifiers:** 01234567890ABCD, 012345

**Visit Number (FIN):** 01234567890ABCD **Collection Date:** 00/00/0000 00:00

## **Chorionic Villus, FISH**

ARUP test code 0040203

Chorionic Villus, FISH

See Note

(Ref Interval: Normal)

Test Performed: Chorionic Villus, FISH (FISHCVS) Specimen Type: Direct (uncultured) villi Indication for Testing: Trisomy 21 mosaicism in mother (2%), pregnancy related conditions, family history of other congenital malformation, deformations and chromosomal abnormalities

RESULT

Normal FISH Result (Female)

INTERPRETATION

There was no evidence for aneuploidy of chromosomes 13, 18, 21, X and Y in 50 interphase cells scored.

This panel will not detect approximately one third of prenatal chromosome abnormalities. Aneuploidy of other chromosomes, structural abnormalities, and mosaicism have not been ruled out by this analysis. Additional testing is recommended for the final interpretation of this result; pending results will be reported separately.

FISH analysis performed on CVS presumes that the fetal chromosome complement is accurately reflected in the extra-embryonic tissue. Rarely, testing of placenta/villi will yield results that differ from those obtained from testing the fetus or newborn. In addition, contamination of the sample with cells of maternal origin may result in the analysis of the maternal rather than fetal cells.

This analysis was performed with chromosome enumeration probes for 13, 18, 21, X and Y using the AneuVysion probe kit (Abbott Molecular).

Cytogenomic Nomenclature (ISCN): nuc ish(DXZ1x2,DYZ3x0,D18Z1x2),(RB1,D21S259/D21S341/D21S342)x2

This result has been reviewed and approved by

A portion of this analysis was performed at the following location(s):

H=High, L=Low, \*=Abnormal, C=Critical

4848



INTERPRETIVE INFORMATION: Fluorescence in Situ Hybridization, CVS

This test was developed and its performance characteristics determined by ARUP Laboratories. It has not been cleared or approved by the US Food and Drug Administration. This test was performed in a CLIA certified laboratory and is intended for clinical purposes.

Counseling and informed consent are recommended for genetic testing. Consent forms are available online.

EER Chorionic Villus, FISH

See Note

Authorized individuals can access the ARUP Enhanced Report using the following link:

VERIFIED/REPORTED DATES				
Procedure	Accession	Collected	Received	Verified/Reported
Chorionic Villus, FISH	23-019-105063	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
EER Chorionic Villus, FISH	23-019-105063	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00

END OF CHART

H=High, L=Low, \*=Abnormal, C=Critical

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