

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

Client: [REDACTED]  
Physician: [REDACTED]

ARUP Test Code: 2002297  
Collection Date: 12/14/2016  
Received in lab: 12/15/2016  
Completion Date: 12/19/2016

**Interpretation**

Specimen Received  
Specimen Type: Amniotic Fluid  
Reason for Referral: Advanced Maternal Age; Abnormal NIPT T21  
Test Performed: FISH, Prenatal

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ABNORMAL FISH RESULT: Trisomy 21

**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 FDA-approved AneuVysion probe kit (Abbott Molecular). 50 interphase cells were scored for each probe.

This analysis showed three hybridization signals for chromosome 21, consistent with trisomy 21.

Sex chromosomes: XY (male)

ISCN:  
nuc ish(DXZ1x1,DYZ3x1,D18Z1x2),  
(RB1x2,D21S259/D21S341/D21S342x3)

**Recommendations:**  
The American College of Medical Genetics recommends that (1) clinical decision-making should be based on information from two of three of the following: positive FISH results, confirmatory chromosome analysis, or consistent clinical information, (2) for genetic counseling of families that have a fetus identified as positive by FISH, chromosome analysis to determine the mutational mechanism accounting for the FISH result is performed, and (3) genetic counseling for all abnormal results.

This result has been reviewed and approved by [REDACTED],  
Ph.D., FACMG  
Electronic Signature

Test developed and characteristics determined by ARUP Laboratories. See Compliance Statement A: [aruplab.com/CS](http://aruplab.com/CS)



Patient: [REDACTED]  
ARUP Accession: 16-349-111954