

**To : Aarohan Fetal Medicine Centre-Indore**

C/o Dr.Preeti Parekh Tomar.

239 A Anoop Nagar

Madhya Pradesh

Indore- 452001

Mobile : 9285559556

**Report Of : Mrs. AKANKSHA PURI**

Pt.Contact :



Sample ID : 2110028226

Patient ID : 10021248

Received on : 01/04/2021 13:30

Registered on : 01/04/2021 19:05

Reported on : 17/04/2021 17:02

**Referred By : DR.PREETI PAREKH TOMAR**

**Cytogenetics Chromosome Analysis Report**

Patient Name : Mrs. AKANKSHA PURI

Gender : Female

Age : 24 Yrs

City : INDORE

Physician Name : DR.PREETI PAREKH TOMAR

Specimen Type : Amniotic Fluid

Test Requested : QF PCR + Prenatal Karyotype

Referral Reason : **TRISOMY 21 SCREEN POSITIVE (1:45)**

46,Normal

46,Normal

**Metaphase Counted** : 20  
**Metaphase karyotyped** : 5  
**Autosome** : Normal  
**Karyotype** : 46,Normal  
**ISCN** : 2016

**Banding technique** : GTG  
**Metaphase Analyzed** : 5  
**Sex Chromosome** : Normal  
**Estimated Band Resolution** : 375-450

**Result** : The cultured amniotic fluid cells showed a normal karyotype.

PRENATAL SEX OF THE FETUS CANNOT BE REVEALED DUE TO CENTRAL GOVERNMENT 2003 ACT ON PRENATAL DIAGNOSIS.

**Interpretation** : Normal Karyotype

**Recommendations** : NA

**Note** :- We have exercised our best efforts to accurately analyze the chromosome karyotypes of this specimen. However, the level of resolution in this G-banded analysis cannot detect cryptic / submicroscopic deletions. The above analysis is based on the sample received in the laboratory. In addition, maternal cell contamination (Only for Prenatal) or low level mosaicism may not be detected.

**Ajinkya Jadhav**  
Incharge Cytogenetics

**Dr. Yamini Jadhav**  
Consultant Cytogenetics

|End of Report |