

MOLQ LABORATORY (A UNIT OF MOLECULAR QUEST HEALTHCARE PVT. LTD.)

Triple Screen

PATIENT INFORMATION NAME MRS, KANUPRIYA PATIENT CODE: 011709210122 DOB: 22/09/85 (DDMMYY)

LMP: 11/05/17 EDD: 17/02/18 PHYSICIAN:

SPECIMEN RECEIVED: 21/09/17

SPECIMEN CODE: MOLQ LAB REFERRING LAB #: MOLO LAB COLLECTION DATE: 21/09/17 REPORTED: 22/09/17

CLINICAL INFORMATION

GESTATIONAL AGE: 18 weeks 5 days from BPD of 18.0 mm on 03/08/17 MATERNAL AGE AT TERM: 32.4 years MATERNAL WEIGHT: 60.0 kg

MATERNAL IDDM: Not specified (Non-diabetic assumed)

GESTATION: Singleton

MATERNAL RACE: INDIAN

SCREENING STATUS: Initial sample

PARA / GRAVIDA: 0 / 1

| CLINICA | L RESULTS | | | | | | | | |
|---------------------------|----------------|---------|---------------|-----------------|-------------|----------|-------------------|-----------------|-----------------------|
| Assay | Results | MoM | DOWN SYNDROME | | | | OPEN SPINA BIFIDA | | |
| AFP | 40.5 ng/mL | 0.76 | | serum screen | age only | | | serum screen | population prevalence |
| uE3 | 1.29 ng/mL | 0.72 | Higher | | | 1:10 | | | |
| hCG | 24731.5 mIU/ml | 0.88 | Risk | | | | Higher Risk | | |
| | | | | | | 1:100 | - Nisk | | |
| | | | 1:250 | | | | | | |
| Risk Assessment (at term) | | | | | | _ 1:1000 | 1 : 721 | | |
| OSB: 1:60100 | | 1:60100 | | | | - 1.1000 | (2.00 MoM) | | |
| Down Syndrome | | 1:1110 | Lower Risk | | | | Lower | | |
| Age alone | | 1:694 | | | | 1:10000 | Risk | | |
| Equivalent Age Risk | | 28.7 | | | | | | | |
| Trisomy 18 | | 1:30700 | | 1:1110 | 1:694 | | | 1:60100 | 1:1000 |

Interpretation* (based on partial information supplied)

DOWN SYNDROME Screen Negative

The risk of Down syndrome is LESS than the screening cut-off. No follow-up

is indicated regarding this result.

OPEN SPINA BIFIDA **Screen Negative**

The maternal serum AFP result is NOT elevated for a pregnancy of this gestational age. The risk of an open neural tube defect is less than the

screening cut-off.

TRISOMY 18 Screen Negative

> These serum marker levels are not consistent with the pattern seen in Trisomy 18 pregnancies. Maternal serum screening will detect approximately

60% of Trisomy 18 pregnancies.