Molecular Quest Healthcare Pvt Ltd

QUADRUPLE MARKER

NAME: MRS NAVDEEP KAUR PATIENT CODE: 012003150005 DOB: 20/03/88 (DDMMYY)

LMP: 10/11/19 EDD: 13/08/20 PHYSICIAN:

SPECIMEN CODE, MOLO

SPECIMEN CODE: MOLQ COLLECTION DATE: 15/03/20

RECEIVED: 15/03/20 REFERRING LAB #: MOLQ

REPORTED: 15/03/20

CLINICAL INFORMATION

GESTATIONAL AGE: 18 weeks 3 days by U/S (18 wks 1 days on 13/03/20)

MATERNAL AGE AT TERM: 32.4 years

MATERNAL WEIGHT: 72.3 kg MATERNAL RACE: ASIAN

MATERNAL HISTORY: IDDM(N), SMOKER(U), RH(N), VPA(U), SSRI(U), CBZ(U), IVF(N)

GESTATION: Singleton

SCREENING STATUS: Initial sample

PARA / GRAVIDA: 0 / 1

| CLINICA | L RESULTS | | | | | | | | |
|-----------------------------------|----------------|------------------|---------------|-----------------|-------------|------------------------|----------------|-----------------|-----------------------|
| Assay | Results | MoM | DOWN SYNDROME | | | OPEN NEURAL TUBE DEFEC | | | |
| AFP | 45.2 ng/mL | 1.06 | | serum screen | age only | | | serum screen | population prevalence |
| uE3 | 1.45 ng/mL | 0.93 | Higher | | | 1:10 | | | |
| 3hCG | 10854.0 mIU/mL | 0.39 | Risk | | | | Higher Risk | | |
| InhA | 103 pg/ml | 0.52 | | | | 1:100 | 1:104 — | | |
| | | | 1:250 | | | | (2.50 MoM) | | |
| Risk Assessment (at term) | | | | | | 1:1000 | | | |
| NTD: | | 1:27200 | | | | 1.1000 | | | |
| Down Syndrome | | -1:50000 | Lower | | | | Lower | | |
| Age alone | | 1:709 | Risk | | | - 1:10000 | Risk | | |
| Equivalent Age Risk Trisomy 18 | | <15.0 1:11500 | | ~1:50000 | 1:709 | | | 1:27200 | 1:1000 |

Interpretation* (This is Screening Test Only Not a Diagnostics Confermatory Test)

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 NEURAL TUBE DEFECT 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.

| Reviewed by: | |
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| neviewed by | |