Molecular Quest Healthcare Pvt Ltd

QUADRUPLE MARKER

NAME: MRS SHALINI SRIVASTAVA PATIENT CODE: 012006100060 DOB: 01/12/93 (DDMMYY)

LMP: 08/02/20 EDD: 29/10/20 PHYSICIAN:

SPECIMEN CODE MOLO

SPECIMEN CODE: MOLQ COLLECTION DATE: 10/06/20 RECEIVED: 10/06/20 REFERRING LAB #: MOLQ

REPORTED: 11/06/20

CLINICAL INFORMATION

GESTATIONAL AGE: 19 weeks 6 days from BPD of 46.0 mm on 10/06/20

MATERNAL AGE AT TERM: 26.9 years MATERNAL WEIGHT: 53.0 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 DEFECT | | | |
| AFP | 72.0 ng/mL | 1.19 | | serum screen | age only | | | serum screen | population prevalence |
| uE3 | 2.72 ng/mL | 1.32 | Higher | | | 1:10 | _ | | |
| ßhCG | 24239.9 mIU/mL | 0.83 | Risk | | | | Higher Risk | | |
| InhA | 100 pg/ml | 0.47 | | | | 1:100 | 1 : 104 — (2.47 MoM) | | |
| Risk As | sessment (at term |) | 1:250 | | | 4.4000 | (2.11 1110111) | | |
| | | 1:13300 | | | | _ 1:1000 | | | |
| Down Syndrome | | 1:50000 | Lower | | | 4:40000 | Lower | | |
| Age a | lone | 1:1260 | Risk | | | - 1:10000 | Risk | | |
| Equivalent Age Risk | | <15.0 | L | | | | | | |
| Trisomy 18 | | 1:187000 | | <1:50000 | 1:1260 | | | 1:13300 | 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 | <i>r</i> - |
|-------------|------------|
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