Molecular Quest Healthcare Pvt Ltd

QUADRUPLE MARKER

NAME: MRS PRIYANKA PATIENT CODE: 012008080098 DOB: 16/12/87 (DDMMYY)

LMP: 27/03/20 EDD: 02/01/21 PHYSICIAN:

SPECIMEN

SPECIMEN CODE: MOLQ COLLECTION DATE: 08/08/20 RECEIVED: 08/08/20 REFERRING LAB #: MOLQ

REPORTED: 10/08/20

CLINICAL INFORMATION

GESTATIONAL AGE: 19 weeks 0 day from BPD of 43.4 mm on 08/08/20

MATERNAL AGE AT TERM: 33.0 years MATERNAL WEIGHT: 58.0 kg

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

GESTATION: Singleton

MATERNAL RACE: ASIAN

SCREENING STATUS: Initial sample

PARA / GRAVIDA: 0 / 1

| CLINICAL RESULTS | | | | | | | | | |
|---------------------------|---------------|----------|----------------|-----------------|-------------|-------------------------|---|-----------------|--------------------------|
| Assay | Results | MoM | DOWN SYNDROME | | | OPEN NEURAL TUBE DEFECT | | | |
| AFP | 62.8 ng/mL | 1.22 | | serum screen | age only | | | serum screen | population prevalence |
| uE3 | 2.05 ng/mL | 1.15 | Higher Risk | | , | 1:10 | | | |
| ßhCG | 14561.1 mIU/m | L 0.49 | | | | | Higher Risk 1 : 104 (2.47 MoM) | | |
| InhA | 153 pg/ml | 0.76 | | | | 1:100 | | | |
| Risk Assessment (at term) | | | 1:250 | | 4,4000 | | | | |
| NTD: | , | 1:12000 | | | | 1:1000 | | | |
| Down Syndrome <1: | | <1:50000 | Lower Risk | | | | Lower Risk | | |
| Age alone | | 1:628 | | | | — 1:10000 | | | |
| Equivalent Age Risk | | <15.0 | | | | | | | |
| Trisomy 18 | | 1:38000 | | <1:50000 | 1:628 | | | 1:12000 | 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> - |
|-------------|------------|
| | |