Molecular Quest Healthcare Pvt Ltd

QUADRUPLE MARKER

NAME: MRS KIRAN

PATIENT CODE: 012005270034 DOB: 11/02/92 (DDMMYY)

LMP: 15/01/20 EDD: 23/10/20 PHYSICIAN:

SPECIMEN

SPECIMEN CODE: MOLQ COLLECTION DATE: 27/05/20 RECEIVED: 27/05/20 REFERRING LAB #: MOLO

REPORTED: 28/05/20

CLINICAL INFORMATION

GESTATIONAL AGE: 18 weeks 5 days from BPD of 41.3 mm on 25/05/20

MATERNAL AGE AT TERM: 28.7 years

MATERNAL WEIGHT: 66.0 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 | 37.3 ng/mL | 0.81 | | serum screen | age only | | | serum screen | population prevalence |
| uE3 | 2.12 ng/mL | 1.27 | Higher | | | 1:10 | _ | | |
| ßhCG | 17221.0 mIU/m | L 0.61 | Risk | | | | Higher Risk | | |
| InhA | 239 pg/ml | 1.20 | | | | 1:100 | 1:104 | | |
| | 10 | | 1:250 | | | | (2.50 MoM) | | |
| Risk Assessment (at term) | | | | 1:1000 | | | | | |
| NTD: | | 1:55000 | | | | - 1.1000 | | | |
| Down Syndrome | | 1:11300 | Lower | | | | Lower | | |
| Age alone | | 1:1120 | Risk | | | - 1:10000 | Risk | | |
| Equivalent Age Risk <15.0 | | | | | | | | | |
| Trisomy 18 | | 1:65900 | | 1:11300 | 1:1120 | | | 1:55000 | 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>i</i> - |
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