Molecular Quest Healthcare Pvt Ltd

QUADRUPLE MARKER

NAME: MRS SHAESTA KHAN PATIENT CODE: 012010070069 DOB: 12/10/91 (DDMMYY)

LMP: 14/05/20 EDD: 15/02/21 PHYSICIAN:

SPECIMEN

SPECIMEN CODE: MOLQ COLLECTION DATE: 07/10/20 RECEIVED: 07/10/20 REFERRING LAB #: MOLQ

REPORTED: 08/10/20

CLINICAL INFORMATION

GESTATIONAL AGE: 21 weeks 2 days from BPD of 41.0 mm on 16/09/20

MATERNAL AGE AT TERM: 29.3 years MATERNAL WEIGHT: 52.0 kg

MATERNAL WEIGHT: 52.0 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 | 94.1 ng/mL | 1.13 | | serum screen | age only | | | serum screen | population prevalence |
| uE3 | 2.66 ng/mL | 1.03 | Higher | | | 1:10 | _ | | |
| ßhCG | 24169.5 mIU/mL | 1.22 | Risk | | | | Higher Risk | | |
| InhA | 244 pg/ml | 1.13 | _ | | | 1:100 | 1:104 — | | |
| | , 0 | | 1:250 | | | | (2.33 MoM) | | |
| Risk As | sessment (at tern | n) | | | | 1:1000 | | | |
| NTD: | | 1:5120 | | | | - 1.1000 | | | |
| Down Syndrome | | 1:15800 | Lower | | | 4.40000 | Lower | | |
| Age alone | | 1:1060 | Risk | | | - 1:10000 | Risk | | |
| Equivalent Age Risk | | <15.0 | L | | | | | | |
| Trisomy 18 | | 1:204000 | | 1:15800 | 1:1060 | | | 1:5120 | 1:1000 |

Interpretation* (This is a Screening Test only Not a Confirmatory Diagnostic 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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