Molecular Quest Healthcare Pvt Ltd

QUADRUPLE MARKER

NAME: MRS PREETI

PATIENT CODE: 012003120268 DOB: 07/07/90 (DDMMYY)

LMP: 20/10/19 EDD: 03/08/20 PHYSICIAN:

SPECIMEN

SPECIMEN CODE: MOLQ COLLECTION DATE: 12/03/20 RECEIVED: 12/03/20 REFERRING LAB #: MOLQ

REPORTED: 13/03/20

CLINICAL INFORMATION

GESTATIONAL AGE: 19 weeks 3 days from BPD of 37.8 mm on 26/02/20

MATERNAL AGE AT TERM: 30.1 years

MATERNAL WEIGHT: 72.0 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

| Assay | L RESULTS Results | MoM | DOWN C | VNDDOME | | | ODEN NE | UDAL TI | IDE DEEEC |
|---------------------------|--------------------|---------|--------|-----------------|-------------|-----------|----------------|-----------------|-----------------------|
| Assay | IVESUIIS | IVIOIVI | DOWN S | YNDROME | | | OPEN NE | URAL IL | JBE DEFEC |
| AFP | 47.2 ng/mL | 0.97 | | serum screen | age only | | | serum screen | population prevalence |
| uE3 | 1.50 ng/mL | 0.83 | Higher | 30,001 | | 1:10 | | 3010011 | prevalence |
| ßhCG | 21006.0 mIU/mL | 0.85 | Risk | | | | Higher Risk | | |
| InhA | 176 pg/ml | 0.85 | - | | | 1:100 | 1:104 — | | |
| | , 0 | | 1:250 | | | | (2.47 MoM) | | |
| Risk Assessment (at term) | | | | | | 1:1000 | | | |
| NTD: | | 1:26700 | | | | 1.1000 | | | |
| Down Syndrome | | 1:12300 | Lower | | | | Lower | | |
| Age alone | | 1:986 | Risk | | | - 1:10000 | Risk | | |
| Equivalent Age Risk <15.0 | | | | | | | | | |
| Trisomy 18 | | 1:63600 | | 1:12300 | 1:986 | | | 1:26700 | 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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