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|  |               |            |  | Date of Report<br>PRISCA | 14-09-2023<br>5.2.0.13 |
|--|---------------|------------|--|--------------------------|------------------------|
| Patient Data   |               |            |  |                          | 0.210120               |
| Name   | MRS MAHIMA    |            |  |                          | 012309130047           |
| Birthday   |               | 22-11-1997 | Sample ID  |                          | 11805158               |
| Age at Sample date   |               | 25.8       | Sample Date  |                          | 13-09-2023             |
| Gestational age 13+2   |               |            | 2  |                          |                        |
| Correction factors   |               |            |  |                          |                        |
| Fetuses  | 1 IVF         |            | unknown  | Previous trisomy 21      | unknown                |
| Weight in kg   | 40.7 Diabetes |            | NO   | Pregnancies              | unknown                |
| Smoker   | NO Origin     |            | Asian  |                          |                        |
| Biochemical Data   |               |            | Ultrasound Data  |                          |                        |
| Parameter  | Value         | Corr Mom   | Gestational age  |                          | 13+2                   |
| PAPP-A   | 5.48 mIU/n    | nl 0.49    | Method   |                          | CRL (<>Robinson)       |
| fb-hCG   | 34.6 ng/ml    | 1          | Scan date  |                          | 13-09-2023             |
| Risks at sampling date   |               |            | Crown rump length in mm 71.1   |                          |                        |
| Age Risk   |               | 1:948      | Nuchal translu   | cency MoM                | 0.62                   |
| Biochemical T21 risk   |               | 1:1060     | Nasal bone   |                          | PRESENT                |
| Combined trisomy 21 risk   |               | 1:6356     | Sonographer  |                          | DR SAHIL LOOMBA        |
| Trisomy 13/18 + NT   |               | <1:10000   | Qualifications   | in measuring NT          | MBBS                   |
| Risk   |               |            | Down's Syndrome Risk (Trisomy 21 Screening)  |                          |                        |
| Risk   1:10   1:250   00   1:250   00   1:1000   1:1000   1:1000   1:1000   1:1000   1:1000   1:1000   1:10000   1:10000   1:10000   1:10000   1:10000   1:10000   1:10000   1:10000   1:10000   1:10000 |               |            | The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.<br>After the result of the Trisomy 21 test (with NT) it is expected that among 6356 women with the same data, there is one woman with a trisomy 21 pregnancy and 6355 women with not affected pregnancies.<br>The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that the risk calculations are statistical aapproaches and have no diagnostic value!<br>The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998).<br>The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values |                          |                        |
| Risk Above Cut Off   |               |            |  | e Risk                   | Risk below Age risk    |