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| Sample Collection |  |  |  |
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|  |                  |            |  | Date of Report<br>PRISCA | 03-04-2021<br>5.0.2.37 |
|--|------------------|------------|--|--------------------------|------------------------|
| Patient Data   |                  |            |  |                          |                        |
| Name   |                  | MRS SALMA  | Patient ID   |                          | 012104010214           |
| Birthday   |                  | 07-03-1992 | Sample ID  |                          | 10926675               |
| Age at delivery  |                  | 29.1       | Sample Date  |                          | 01/04/2021             |
| Gestational age  |                  | 13+1       |  |                          |                        |
| Correction factors   |                  |            |  | -                        |                        |
| Fetuses  | 1 IVF            |            | unknown  | Previous trisomy 21      | unknown                |
| Weight in kg   | 64.5 Diabetes    |            | unknown  | Pregnancies              | unknown                |
| Smoker   | Unknown Origin   |            | Asian  |                          |                        |
| Biochemical Data   |                  |            | Ultrasound Da  | ata                      |                        |
| Parameter  | Value            | Corr Mom   | Gestational age  | 2                        | 12+6                   |
| PAPP-A   | 2.42 mIU/ml      | 0.47       | Method   |                          | CRL(<>Robinson         |
| fb-hCG   | 18.3 ng/ml       | 0.43       | Scan date  |                          | 31-03-2021             |
| Risks at sampling date   | ;                |            | Crown rump le  | ength in mm              | 66.3                   |
| Age Risk   |                  | 1:734      | Nuchal translu   | cency MOM                | 0.95                   |
| Biochemical T21 risk   |                  | 1:4100     | Nasal bone   |                          | Present                |
| Combined Trisomy 21  | Risk             | <1:10000   | Sonographer  |                          | DR INDRAJEET KUNDU     |
| Trisomy 13/18 + NT   |                  | <1:10000   | Qualification in   | n measuring NT           | MD                     |
| Risk<br>1:10   |                  | 1          | Down's Syndrome Risk (Trisomy 21 Screening)  |                          |                        |
|  |                  |            | The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.  |                          |                        |
| 1:1000<br>1:10000<br>13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49<br>Age<br>Trisomy 13/18 + NT<br>The calculated risk for Trisomy 13/18 (with NT) is |                  |            | After the result of the Trisomy 21 test (with NT) it is<br>expected that among more than 10000 women with the<br>same data, there is one woman with a trisomy 21<br>pregnancy and 9999 women with not affected pregnancies.<br>The calculated risk by <b>PRISCA</b> depends on the accuracy of<br>the information provided by the referring physician. Please<br>note that the risk calculations are statistical aapproaches and<br>have no diagnostic value!<br>The laboratory cannot be hold responsible for their impact<br>on the risk assessment! Calculated risks have no diagnostic<br>values |                          |                        |
|  | sk Above Cut Off |            | Risk above Ag  | e Risk                   | Risk below Age risk    |