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|-------------------|--|--|--|--|
| Sample Collection |  |  |  |  |
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|  |                    |            |  | Date of Report<br>PRISCA   | 08-02-2021<br>5.0.2.37 |
|--|--------------------|------------|--|--|------------------------|
| Patient Data   |                    |            |  |  |                        |
| Name   |                    | MRS NEHA   | Patient ID   |  | 012102060084           |
| Birthday   |                    | 04-08-1991 | Sample ID  |  | 10881466               |
| Age at delivery  |                    | 29.5       | Sample Date  |  | 06/02/2021             |
| Gestational age  |                    | 13+4       |  |  |                        |
| Correction factors   |                    |            |  |  |                        |
| Fetuses  | 1 IVF              |            | unknown  | Previous trisomy 21  | unknown                |
| Weight in kg   | 58.3 Diabetes      |            | unknown  | Pregnancies  | unknown                |
| Smoker   | Unknown Origin     |            | Asian  |  |                        |
| <b>Biochemical Data</b>  |                    |            | Ultrasound Da  | ata  |                        |
| Parameter  | Value              | Corr Mom   | Gestational age  | 2  | 13+4                   |
| PAPP-A   | 5.3 mIU/ml         | 0.8        | Method   |  | CRL(<>Robinson         |
| fb-hCG   | 25.8 ng/ml         | 0.62       | Scan date  |  | 06-02-2021             |
| Risks at sampling dat  | te                 |            | Crown rump le  | ength in mm  | 75                     |
| Age Risk   |                    | 1:709      | Nuchal translu   | cency MOM  | 0.6                    |
| Biochemical T21 risk   | ζ.                 | 1:7457     | Nasal bone   |  | Present                |
| Combined Trisomy 2   | 21 Risk            | <1:10000   | Sonographer  |  | )R.PRAKASH LALCHANDANI |
| Trisomy 13/18 + NT   |                    | <1:10000   | Qualification in   | n measuring NT   | MD                     |
| Risk<br>1:10   |                    | _          | Down's Syndrome Risk (Trisomy 21 Screening)  |  |                        |
| 1:100  | /                  |            | <b>cut off, which</b><br>After the result  | <b>l risk for Trisomy 21</b><br>represents a low risk<br>t of the Trisomy 21 to<br>unong more than 100 | est (with NT) it is    |
| 1:1 <mark>000<br/>1:10000<br/>13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49</mark><br>Age             |                    |            | 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! |  |                        |
| Trisomy 13/18 + NT<br>The calculated risk for Trisomy 13/18 (with NT) is<br><1:10000, which indicates a low risk |                    |            | The laboratory cannot be hold responsible for their impact<br>on the risk assessment! Calculated risks have no diagnostic<br>values  |  |                        |
| F  | Risk Above Cut Off |            | Risk above Ag  | e Risk   | Risk below Age risk    |