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|   |                         |           |   | Date of Report<br>PRISCA | 6/10/2022<br>5.1.0.17 |
|---|-------------------------|-----------|---|--------------------------|-----------------------|
| Patient Data  |                         |           |   |                          |                       |
| Name  | MRS. ANURADHA           | A AGARWAL | Patient ID  |                          | 012210040197          |
| Birthday  |                         | 5/9/1987  | Sample ID   |                          | 11574437              |
| Age at term   |                         | 35.07     | Sample Date   |                          | 4/10/2022             |
| Gestational age   |                         | 11+3      |   |                          |                       |
| Correction factors  |                         |           |   |                          |                       |
| Fetuses   | 1 IVF                   |           | unknown   | Previous trisomy 21      | unknown               |
| Weight in kg  | 67 Diabetes             |           | NO  | Pregnancies              | unknown               |
| Smoker  | NO Origin               |           | Asian   |                          |                       |
| Biochemical Data  |                         |           | Ultrasound Data   |                          |                       |
| Parameter   | Value                   | Corr Mom  | Gestational age   | e                        | 11+2                  |
| PAPP-A  | 3.1 mIU/ml              | 1.03      | Method  |                          | CRL (<>Robinson)      |
| fb-hCG  | 43.2 ng/ml              | 0.92      | Scan date   |                          | 4/10/2022             |
| Risks at sampling date  |                         |           | Nuchal translucency 1.3   |                          |                       |
| Age Risk 1:255  |                         | 1:255     | Nuchal translucency MoM 1.02  |                          |                       |
| Biochemical T21 risk  | nemical T21 risk 1:2049 |           | Nasal bone  | Nasal bone Presen        |                       |
| Combined trisomy 21   | risk                    | 1:7656    |   |                          |                       |
| Trisomy 13/18 + NT  |                         | <1:10000  |   |                          |                       |
| Risk  |                         |           | Down's Syndrome Risk (Trisomy 21 Screening)   |                          |                       |
| Risk<br>1:10<br>1:250 Cut off<br>1:1000   |                         |           | The calculated risk for Trisomy 21 (with NT) is below the<br>cut off, which represents a low risk.<br>After the result of the Trisomy 21 with NT test it is<br>expected that among 7656 women with the<br>same data, there is one woman with a trisomy 21<br>pregnancy and 7655 women with not affected pregnancies.<br>The calculated risk by PRISCA 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! |                          |                       |
| 13 15 17 19 21 23 24<br>Trisomy 13/18+NT<br>The calculated risk for<br><1:10000 , which indic |                         | Age       | The laboratory cannot be hold responsible for their impact<br>on the risk assessment! Calculated risks have no diagnostic<br>values<br>Risk above Age Risk  |                          |                       |