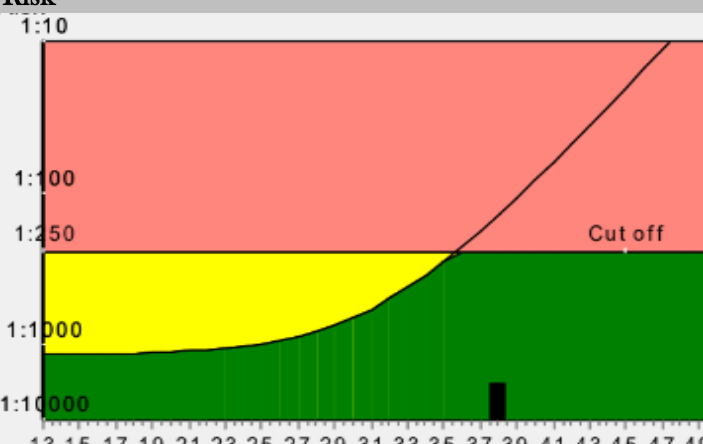


Date of Report 14-05-2024  
PRISCA 5.2.0.13

| Patient Data  |                   |          |   |                     |         |
|---|-------------------|----------|---|---------------------|---------|
| Name  | MRS. MEENU TIWARI |          | Patient ID  | 12405090201         |         |
| Birthday  | 21-05-1986        |          | Sample ID   | 11888373            |         |
| Age at Sample date  | 38.0              |          | Sample Date   | 09-05-2024          |         |
| Gestational age   | 13+6              |          |   |                     |         |
| Correction factors  |                   |          |   |                     |         |
| Fetuses   | 1                 | IVF      | unknown   | Previous trisomy 21 | unknown |
| Weight in kg  | 61                | Diabetes | NO  | Pregnancies         | unknown |
| Smoker  | NO                | Origin   | Asian   |                     |         |
| Biochemical Data  |                   |          | Ultrasound Data   |                     |         |
| Parameter   | Value             | Corr Mom | Gestational age   | 13+6                |         |
| PAPP-A  | 6.8 mIU/ml        | 0.80     | Method  | CRL (<>Robinson)    |         |
| fb-hCG  | 24.1 ng/ml        | 0.98     | Scan date   | 09-05-2024          |         |
| Risks at sampling date  |                   |          | Crown curm length in mm   | 78.7                |         |
| Age Risk  | 1:141             |          | Nuchal translucency MoM   | 0.57                |         |
| Bicochemical T21 risk   | 1:563             |          | Nasal bone  | present             |         |
| Commbined Trisomy 21 risk   | 1:2960            |          | Sonographer   | DR.PAVEEN           |         |
| Trisomy 13/18 + NT  | <1:10000          |          | Qualifications in measuring NT  | C/R                 |         |
| Risk  |                   |          | Down's Syndrome Risk (Trisomy 21 Screening)   |                     |         |
|       |                   |          | <p>The calculated risk for Trisomy 21 is below the cut off, which represents a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 2960 women with the same data, there is one woman with a trisomy 21 pregnancy and 2959 women with not affected pregnancies.</p> <p>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 approaches and have no diagnostic value!</p> <p>The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998).</p> |                     |         |
| Trisomy 13/18+NT  |                   |          | The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values   |                     |         |
| The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk |                   |          |   |                     |         |

Risk Above Cut Off
  Risk above Age Risk
  Risk below Age risk