

Risk Above Cut Off

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Risk below Age risk

 Date of Report
 21-01-2021

 PRISCA
 5.0.2.37

|  |                  |  | PRISCA              | 5.0.2.37     |
|--|------------------|--|---------------------|--------------|
| Patient Data   |                  |  |                     |              |
| Name M   | IRS NITIN KUMARI | Patient ID   |                     | 012101200026 |
| Birthday   | 14-01-1992       | Sample ID  |                     | 10867475     |
| Age at delivery  | 29               | Sample Date  |                     | 20/01/2021   |
| Gestational age  |                  |  |                     |              |
| Correction factors   |                  |  |                     |              |
| Fetuses 1 $\Gamma$   | VF               | unknown  | Previous trisomy 21 | unknown      |
| Weight in kg 60 $\Gamma$   | Diabetes         | unknown  | Pregnancies         | unknown      |
| Smoker Unknown C   | Origin           | Asian  |                     |              |
| Biochemical Data   | Ultrasound Data  |  |                     |              |
| Parameter Value  | Corr Mom         | Gestational ago  | е                   | 11+5         |
| PAPP-A 1.84 n  | nIU/ml 0.56      | Method   |                     | LMP          |
| fb-hCG 44.4 n  | g/ml 0.87        |  |                     |              |
| Risks at sampling date   |                  |  |                     |              |
| Age Risk   | 1:700            |  |                     |              |
| Biochemical T21 risk   | 1:600            |  |                     |              |
| Combined Trisomy 21 Risk   | 1:1535           |  |                     |              |
| Trisomy 13/18 + NT   | 1:7003           |  |                     |              |
|  |                  | Down's Syndrome Risk (Trisomy 21 Screening)  |                     |              |
| 1:10   |                  | The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.  |                     |              |
| 1:1000<br>1:1000<br>1:10000  |                  | After the result of the Trisomy 21 test (with NT) it is expected that among more than 1535 women with the same data, there is one woman with a trisomy 21 pregnancy and 1534 women with not affected pregnancies. 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! |                     |              |
| Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 (with NT) is 1:7003, 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  |                     |              |

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