

which indicates a low risk

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Date of Report 10-12-2024 PRISCA 5.2.0.13

| | | | | PRISCA | 5.2.0.13 |
|--------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|---------------------|------------------|
| Patient Data | | | | | |
| Name MRS. GUDIYA PARVEEN | | Patient ID | | 12412090060 | |
| Birthday 11-05-1997 | | Sample ID 11996 | | 11996178 | |
| Age at Sample date | | 27.6 | Sample Date | | 09-12-2024 |
| Gestational age | | 13+3 | | | |
| Correction factors | 1 | | | _ | |
| Fetuses | IVF | | unknown | Previous trisomy 21 | unknown |
| Weight in kg 55 | Diabetes | | NO | Pregnancies | unknown |
| Smoker NC | Origin | | Asian | | |
| Biochemical Data | | | Ultrasound Data | | |
| Parameter Value | ; | Corr Mom | Gestational ag | e | 13+3 |
| PAPP-A 6.9 | mIU/ml | 0.79 | Method | | CRL (<>Robinson) |
| fb-hCG 21. | i ng/ml | 0.72 | Scan date | | 09-12-2024 |
| Risks at sampling date | | | Crown rump length in mm 73.8 | | |
| Age Risk | | 1:849 | Nuchal translu | icency MoM | 0.88 |
| Biochemical T21 risk | | 1:6461 | Nasal bone | | present |
| Combined trisomy 21 risk | | <1:10000 | Sonographer | | DR. DEEPIKA |
| Trisomy 13/18 + NT | | <1:10000 | Qualifications | in measuring NT | MD |
| Risk | | | Down's Syndrome Risk (Trisomy 21 Screening) | | |
| 1:10 1:100 1:250 G.t.off | | The calculated risk for Trisomy 21(with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. | | | |
| 1:1000 1:1000 1:1000 1:1000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Trisomy 13/18+NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000, | | | 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! The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998). The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values | | |