

The calculated risk for Trisomy 13/18 (with NT) is <1:10000,

which indicates a low risk

*Free Home Sample Collection 9999 778 778



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Date of Report 07-06-2024 PRISCA 5.2.0.13

The laboratory cannot be hold responsible for their impact on the

risk assessment! Calculated risks have no diagnostic values

| | | | | | Date of Report | 0, 00 2021 |
|--|-------|----------|---------------------|---|---------------------|------------------|
| | | | | | PRISCA | 5.2.0.13 |
| Patient Data | | | | | | |
| Name | | MRS. PO | OJA DAYMA | Patient ID | | 12406060093 |
| Birthday | | | 20-11-2001 | Sample ID | | 11896347 |
| Age at Sample date | | | 22.5 | Sample Date | | 06-06-2024 |
| Gestational age | | | 11+6 | | | |
| Correction factors | | | | | | |
| Fetuses | 1 | IVF | | unknown | Previous trisomy 21 | unknown |
| Weight in kg | 62 | Diabetes | | NO | Pregnancies | unknown |
| Smoker | NO | Origin | | Asian | | |
| Biochemical Data | | | | Ultrasound Data | | |
| Parameter | Value | | Corr Mom | Gestational age | e | 11+6 |
| PAPP-A | 3.9 | mIU/ml | 0.98 | Method | | CRL (<>Robinson) |
| fb-hCG | 86.7 | ng/ml | 1.98 | Scan date | | 06-06-2024 |
| Risks at sampling date | | | | Crown rump l | ength in mm | 51.1 |
| Age Risk | | | 1:1017 | Nuchal translu | icency MoM | 0.60 |
| Biochemical T21 risk | | | 1:1179 | Nasal bone | | PRESENT |
| Combined trisomy 21 risk | | | - 1:6495 | Sonographer | | DR. |
| Trisomy 13/18 + NT | | | < 1:10000 | | in measuring NT | MBBS |
| Risk | | | | Down's Syndrome Risk (Trisomy 21 Screening) | | |
| 1:100 1:100 1:100 1:1000 1:10000 1:10000 1:10000 | | | | 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 6495 women with the same data, there is one woman with a trisomy 21 pregnancy and 6494 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! The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998). | | |
| Trisomy 13/18+NT | | | | 1998). | | |