

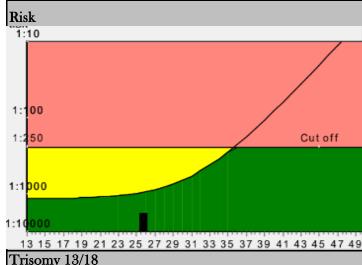
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Date of Report 26-04-2024 **PRISCA** 5.2.0.13 Patient Data MRS. ASHITA Patient ID Name 12404250130 11835043 Birthday 23-07-1998 Sample ID Age at Sample date 25.8 Sample Date 25-04-2024 Gestational age 13+6Correction factors 1 IVF Fetuses unknown Previous trisomy 21 unknown Weight in kg 91 Diabetes NO Pregnancies unknown Smoker NO Origin Asian **Biochemical Data** Ultrasound Data 13+6 **Parameter** Value Corr Mom Gestational age 6.8 mIU/ml1.32 CRL (<>Robinson) Method 2.25 25-04-2024 49.5 ng/ml Scan date

PAPP-A fb-hCG 79.1 Risks at sampling date Crown rump length in mm 1:968 0.73 Age Risk Nuchal translucency MoM Biochemical T21 risk 1:1477 Nasal bone present Overall population risk 1:7672 Sonographer DR. DHRUV TANEJA <1:10000 Trisomy 13/18 + NT Qualifications in measuring NT **MBBS** Down's Syndrome Risk (Trisomy 21 Screening)



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 7672 women with the same data, there is one woman with a trisomy 21 pregnancy and 7671 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).

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