

Trisomy 13/18 + NT

The calculated risk for Trisomy 13/18 (with NT) is

Risk Above Cut Off

<1:10000, which indicates a low risk

The laboratory cannot be hold responsible for their impact

on the risk assessment! Calculated risks have no diagnostic

Risk below Age risk

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Date of Report 25/12/2021 PRISCA 5.2.0.13 Patient Data MRS. YASODA Patient ID 012112230147 Name 21/03/1986 Sample ID 11283299 Birthday Age at delivery 36.28 Sample Date 23/12/2021 Gestational age 13+0 Correction factors 1 IVF Fetuses unknown Previous trisomy 21 unknown NO Pregnancies Weight in kg 50 Diabetes unknown NO Origin Smoker Asian **Biochemical Data** Ultrasound Data 12+5 Value Corr Mom Gestational age **Parameter** PAPP-A 17.8 mIU/ml2.81 CRL(<>Robinson Method 22/12/2021 fb-hCG 64.7 ng/ml 1.51 Scan date Risks at sampling date Nuchal translucency 1.3 Age Risk 1:323 0.77 Nuchal translucency MoM Biochemical T21 risk 1:2776 Nasal bone Present Combined T21 risk <1:10000 Trisomy 13/18+NT <1:10000 Risk Down's Syndrome Risk (Trisomy 21 Screening) The calculated risk for Trisomy 21(with NT) is below the 1:10 cut off, which represents a low risk. After the result of the Trisomy 21 test it is expected that among more than 10000 women with the 1:100 same data, there is one woman with a trisomy 21 1:250 Cut off pregnancy and 9999 women with not affected pregnancies. The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please 1:1000 note that the risk calculations are statistical aapproaches and have no diagnostic value! 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age

values

Risk above Age Risk