

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 8/9/2021 PRISCA 5.2.0.13 Patient Data 012109080079 MRS. SHUCHI GUPTA Patient ID Name 15/08/1993 Sample ID 10969212 Birthday Age at delivery 28.58 Sample Date 8/9/2021 Gestational age 13+2 Correction factors 1 IVF Fetuses unknown Previous trisomy 21 unknown NO Pregnancies Weight in kg 70 Diabetes unknown NO Origin Smoker Asian **Biochemical Data** Ultrasound Data 12+6 Value Corr Mom Gestational age **Parameter** PAPP-A 4.29 mIU/ml 0.92 CRL(<>Robinson Method 2.98 5/9/2021 fb-hCG 109 ng/ml Scan date Risks at sampling date Nuchal translucency in mm 0.9 Age Risk 1:1111 0.53 Nuchal translucency MOM Biochemical T21 risk 1:393 Nasal bone Present 1:2323 Combined Trisomy 21 Risk Trisomy 13/18 + NT <1:10000 Qualification in measuring NT CR Risk 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. After the result of the Trisomy 21 test (with NT) it is expected that among 2323 women with the 1:100 same data, there is one woman with a trisomy 21 Cut off pregnancy and 2322 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

values

Risk above Age Risk