

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

\*Free Home Sample Collection 9999 778 778



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 15/9/2021 5.2.0.13 PRISCA Patient Data MRS. ANJU Patient ID 012109130226 Name 18/03/1996 Sample ID 10922663 Birthday Age at delivery 26 Sample Date 14/9/2021 Gestational age 13+4 Correction factors 1 IVF Fetuses unknown Previous trisomy 21 unknown NO Pregnancies Weight in kg 45 Diabetes unknown NO Origin Smoker Asian **Biochemical Data** Ultrasound Data 13+3 Value Corr Mom Gestational age **Parameter** PAPP-A 9 mIU/mlCRL(<>Robinson 1.1 Method 0.88 13/9/2021 fb-hCG 36.8 ng/ml Scan date Risks at sampling date Nuchal translucency in mm 2.3 Age Risk 1:1320 1.22 Nuchal translucency MOM Biochemical T21 risk <1:10000 Nasal bone Present <1:10000 Combined Trisomy 21 Risk Trisomy 13/18 + NT <1:10000 Qualification in measuring NT CR Risk Down's Syndrome Risk (Trisomy 21 Screening) Risk 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 (with NT) it is expected that among more than 10000 women with the 1: 00 same data, there is one woman with a trisomy 21 Cut off 1:250 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