

Trisomy 13/18 + NT

indicates a low risk

The calculated risk for Trisomy 13/18 is <1:10000, which

Risk Above Cut Off

*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 21/9/2021 5.2.0.13 PRISCA Patient Data MRS.RITU SHARMA Patient ID 012109200165 Name 21/04/1996 Sample ID 10975429 Birthday Age at delivery 28.4 Sample Date 20/9/2021 Gestational age 13+2 Correction factors 1 IVF Fetuses unknown Previous trisomy 21 unknown NO Pregnancies Weight in kg 56 Diabetes unknown NO Origin Smoker Asian **Biochemical Data** Ultrasound Data 13+1 Value Corr Mom Gestational age **Parameter** PAPP-A 2.85 mIU/ml0.45 CRL(<>Robinson Method 0.51 19/9/2021 fb-hCG 22.5 ng/mlScan date Risks at sampling date Crown rump length in mm 68.4 Age Risk 1:787 0.52 Nuchal translucency MoM Biochemical T21 risk 1:2724 Nasal bone Present Combined trisomy 21 risk <1:10000 Sonographer DR. PAWAN JOON Trisomy 13/18 <1:10000 Qualification in measuring NT Risk Down's Syndrome Risk (Trisomy 21 Screening) Risk 1:10 The calculated risk for Trisomy 21 is below the cut off, which represents a low risk. After the result of the Trisomy 21 test (with NT) it is 1:100 expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 Cut off pregnancy and 9999 women with not affected pregnancies. The calculated risk by PRISCA depends on the accuracy of 1:1000 the information provided by the referring physician. Please 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