

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

Book a Test Online www.molq.in

Date of Report 8/3/2022 5.1.0.17 PRISCA Patient Data MRS. NISHA SHARMA Patient ID 052203070016 Name 10.02.1989 Sample ID 11248085 Birthday Age at term 33.7 Sample Date 7/3/2022 Gestational age 12+1 Correction factors 1 IVF Fetuses unknown Previous trisomy 21 unknown NO Pregnancies Weight in kg 59.3 Diabetes unknown NO Origin Smoker Asian **Biochemical Data** Ultrasound Data 12+1 Value Corr Mom Gestational age Parameter PAPP-A 2.38 mIU/ml 0.6 CRL (<>Robinson) Method 0.63 7/3/2022 fb-hCG 30.6 ng/ml Scan date Risks at sampling date Nuchal translucency 1.9 Age Risk 1:396 Nuchal translucency MoM 1.3 Biochemical T21 risk 1:2059 Nasal Bone Presesnt Combined trisomy 21 risk 1:3570 Trisomy 13/18+NT <1:10000 Risk Down's Syndrome Risk (Trisomy 21 Screening) Risk 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 it is expected that among 3570 women with the 1-100 same data, there is one woman with a trisomy 21 50 Cut off pregnancy and 3569 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