

Trisomy 13/18

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



on the risk assessment! Calculated risks have no diagnostic

Risk below Age risk

Book a Test Online www.molq.in

Date of Report 31/3/2022 5.1.0.17 PRISCA Patient Data MRS. SUPRIYA Patient ID 012203300133 Name 22-9-1998 Sample ID 11474991 Birthday Age at term 23.11 Sample Date 30/3/2022 Gestational age 12+0Correction factors 1 IVF Fetuses unknown Previous trisomy 21 unknown NO Pregnancies Weight in kg 71 Diabetes unknown NO Origin Smoker Asian **Biochemical Data** Ultrasound Data 12+0 Value Corr Mom Gestational age **Parameter** PAPP-A 3.52 mIU/ml 1.17 CRL (<>Robinson) Method 30/03/2022 fb-hCG 66.7 ng/ml 1.42 Scan date Risks at sampling date Nuchal Translucency 1.3 1:996 0.91 Age Risk Nuchal Translucency MoM Biochemical T21 risk 1:3837 Nasal Bone Present Combined T21 risk <1:10000 Trisomy 13/18+NT <1:10000 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 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 9999 women with not affected pregnancies. The calculated risk by PRISCA depends on the accuracy of 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 The laboratory cannot be hold responsible for their impact

values

Risk above Age Risk