

The calculated risk for Trisomy 13/18 (with NT) is

Risk Above Cut Off

<1:10000, which indicates a low risk

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on the risk assessment! Calculated risks have no diagnostic

Risk below Age risk

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Date of Report 30/9/2021 PRISCA 5.2.0.13 Patient Data MRS. SAPNA Patient ID 012109280346 Name 18/05/1996 Sample ID 11188976 Birthday Age at delivery 25.91 Sample Date 28/9/2021 Gestational age 11+6 Correction factors 1 IVF Fetuses unknown Previous trisomy 21 unknown NO Pregnancies Weight in kg 65.7 Diabetes unknown NO Origin Smoker Asian **Biochemical Data** Ultrasound Data 11+5 Value Corr Mom Gestational age **Parameter** PAPP-A 10.5 mIU/ml3.38 CRL(<>Robinson Method Scan date 28/9/2021 fb-hCG 82.4 ng/ml1.78 Risks at sampling date Nuchal translucency in mm 1.1 Age Risk 1:1326 0.79 Nuchal translucency MOM Biochemical T21 risk 1:7660 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) 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 more than 10000 women with the 1:100 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 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 Trisomy 13/18 + NT The laboratory cannot be hold responsible for their impact

values

Risk above Age Risk