

Trisomy 13/18 + NT
The calculated risk for Trisomy 13/18 (with NT) is 1:10000, which indicates a low risk

Risk Above Cut Off



The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

Risk below Age risk

Risk above Age Risk

2/5/2019 Date of Report PRISCA Patient Data Name Mrs Revati Patient ID 011905010116 27/05/1986 Sample ID Birthday 10431463 Sample Date Age at sample date 01/05/2019 32.9 Gestational age by CRL 12+6 Correction factors 1 IVF unknown Previous trisomy 21 Fetuses unknow Weight in kg 59 Diabetes unknown Pregnancies unknow Smoker Biochemical Data Ultrasound Data Parameter Value Corr Mom Gestational age by CRL 12+6 PAPP-A 2.62 mIU/mlMethod CRL (<>Robinson) 0.59 fb-hCG Scan Date 1/5/2019 Risks at sampling date Crown rump length (mm) 65 1:417 1.14 Age Risk Nuchal translucency MoM Biochemical T21 Risk 1:523 Nasal Bone Present Combined Trisomy 21 Risk 1:1493 Sonographer Trisomy 13/18 + NT Qualifications in measuring NT <1:10000 Down's Syndrome Risk (Trisomy 21 Screening) 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 523 women with the ame data, there is one woman with a trisomy 21 pregnancy. 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