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Date of Report 04-02-2020 **PRISCA** 5.0.2.37 Patient Data MRS KANCHAN GARG Patient ID Name 012002030092 28-04-1990 Sample ID 10414100 Birthday 03/02/2020 Age at delivery 29.8 Sample Date 13+3 Gestational age Correction factors 1 IVF Fetuses unknown Previous trisomy 21 unknown Weight in kg 72 Diabetes unknown Pregnancies unknown Smoker Unknown Origin Asian **Biochemical Data** Ultrasound Data Parameter Value Corr Mom Gestational age 13+2CRL(<>Robinson PAPP-A 2.41 mIU/ml0.49 Method fb-hCG 20.6 ng/ml 0.52 Scan date 02-02-2020 Risks at sampling date Crown rump length in mm 71 1:685 1.03 Age Risk Nuchal translucency MOM Biochemical T21 risk 1:2965 Nasal bone Present Combined Trisomy 21 Risk <1:10000 Sonographer DR.SAMRITI SHARMA Trisomy 13/18 + NT <1:10000 Qualification in measuring NT 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 (with NT) it is expected that among more than 10000 women with the 1: 00 same data, there is one woman with a trisomy 21 1:250 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 approaches and

Risk Above Cut Off

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

<1:10000, which indicates a low risk

Trisomy 13/18 + NT

13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49

Risk above Age Risk

values

have no diagnostic value!

Ri

The laboratory cannot be hold responsible for their impact

on the risk assessment! Calculated risks have no diagnostic

Risk below Age risk