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Date of Report 11/3/2022 5.1.0.17 PRISCA Patient Data MRS. MANNA DEVI (F1) Patient ID 012203100027 Name 10/7/1992 Sample ID 11257843 Birthday Age at term 30.01 Sample Date 10/3/2022 Gestational age 13+2 Correction factors 2 IVF Fetuses unknown Previous trisomy 21 unknown 60 Diabetes NO Pregnancies Weight in kg unknown NO Origin Smoker Asian **Biochemical Data** Ultrasound Data 13+4 Value Corr Mom Gestational age **Parameter** PAPP-A 5.28 mIU/ml 0.42 CRL (<>Robinson) Method 0.31 9/3/2022 fb-hCG 27.2 ng/ml Scan date Risks at sampling date Nuchal translucency 1:700 Age Risk 1.09 Nuchal translucency MoM Biochemical T21 risk 1:4646 Nasal Bone Presesnt Combined trisomy 21 risk <1:10000 Trisomy 13/18+NT 1:2875 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 more than 10000 women with the 1:100 same data, there is one woman with a trisomy 21 1:250 Cut off pregnancy 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 aapproaches and

Trisomy 13/18+NT

The calculated risk for Trisomy 13/18 with NT is 1:2875, which indicates a low risk

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 on the risk assessment! Calculated risks have no diagnostic values

Risk Above Cut Off

Risk above Age Risk

have no diagnostic value!

Risk below Age risk