Date of report:	11/06/2024
Prisca	5.2.0.13

Patient data Name MRS. PRIYA Patient ID 5/05/1991 Sample ID Birthday 2406018437/NOD Age at sample date 33.1 Sample Date 10/06/2024 Gestational age 12 + 6 **Correction factors** Fetuses IVF Previous trisomy 21 1 no no pregnancies 54 diabetes Weight no Smoker Asian Origin no **Biochemical data** Ultrasound data Parameter Value Corr. MoM Gestational age 12 + 6 PAPP-A 1.18 mIU/ml 0.23 Method **CRL** Robinson 10/06/2024 fb-hCG 0.92 41.1 ng/ml Scan date Risks at sampling date Crown rump length in mm 67 Age risk 1:404 Nuchal translucency MoM 0.76 **Biochemical T21 risk** 1:54 Nasal bone present Combined trisomy 21 risk 1:398 DR. NEERJA CHOPRA Sonographer Trisomy 13/18 + NT 1:1628 Qualifications in measuring NT M.D Trisomy 21 Risk 1:10 The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 398 women with the same data, there is one woman with a trisomy 21 pregnancy and 397 1:00 women with not affected pregnancies. The PAPP-A level is low. 50 Cut off 1:: The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that risk calculations are statistical approaches and have no diagnostic value! 1:1000 The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)). 1:10 The laboratory can not be hold responsible for their impact 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 on the risk assessment ! Calculated risks have no diagnostic value! Ade Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:1628, which represents a low risk.