KOS DIAGNOSTIC LAB 6349/1, NICHOLSON ROAD, AMBALA CANTT

 Prisca
 5.0.2.37

 Date of report:
 18-01-2018

Patient data Name MRS. POOJA Patient ID 1801220298/AMB Birthday 08-03-1992 Sample ID 1801220298/AMB Sample Date 17-01-2018 Age at sample date 25.9 Gestational age 11 + 3**Correction factors** IVF Fetuses 1 no Previous trisomy 21 no pregancies Weight 39 diabetes no Smoker Origin Asian no Biochemical data Ultrasound data Value Corr. MoM 11 + 2Parameter Gestational age PAPP-A 1.1 mIU/ml **CRL** Robinson 0.31 Method fb-hCG 70.1 ng/ml 1.39 Scan date 16-01-2018 Risks at sampling date Crown rump length in mm 47.5 Age risk 1:885 Nuchal translucency MoM 2.01 1:125 **Biochemical T21 risk** Nasal bone present Combined trisomy 21 risk >1:50 Sonographer Trisomy 13/18 + NT 1:117 Qualifications in measuring NT MD Trisomy 21 Risk 1:10 The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk. After the result of the Trisomy 21 Test (with nuchal translucency), it is expected that among less than 50 pregnancies with the same data, there is one trisomy 21 1:100 pregnancy. The PAPP-A level is low. 1:250 Cutoff 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)). 000 1:10 The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 diagnostic value! Aae Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:117, which represents a low risk.

Sign of Physician