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

5.0.2.37

Date of report: 27-12-2018

Prisca

Patient data					
Name	MRS. ANU	Patient ID		101812260011	
Birthday	13/08/18	Sample ID		101812260011	
Age at sample date	35	Sample Date	e	26/12/18	
Gestational age	12 + 2				
Correction factors					
Fetuses 1	IVF	no	Previous trisomy 21	no	
Weight 76	diabetes	no	pregancies		
Smoker no	Origin	Asian			
Biochemical data		Ultrasound data			
Parameter Value	Corr. MoM	Gestational a	age	12 + 2	
PAPP-A 3.27 mIU/m	l 0.87	Method		CRL Robinson	
fb-hCG 33.1 ng/ml	1.05	Scan date 26/12/18			
Risks at sampling date		Crown rump length in mm			
Age risk	1:317		Nuchal translucency MoM		
Biochemical T21 risk	1:1317	Nasal bone		present	
Combined trisomy 21 risk	21 risk 1:600		Sonographer		
Trisomy 13/18 + NT	1:8290	Qualifications in measuring NT		MD	
1:10 1:00 1:50 Cutoff 1:100 1:1000			Trisomy 21 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 301 women with the same data, there is one woman with a trisomy 21 pregnancy and 300 women with not affected pregnancies. 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! The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)). The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!		

Sign of Physician