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

Prisca 5.0.2.37

Date of report: 16-06-2019

Patient data				
Name	MRS. RITU			1906220662/AMB
Birthday	02-03-1992		Sample ID 19062	
Age at sample date	27.3)	15-06-2019
Gestational age	11 + 5			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 75	diabetes	no	pregancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound data		
Parameter Value	Value Corr. MoM G		Gestational age 11 + 3	
PAPP-A 3.67 mIU/m	l 1.94	Method CRL Robinson		
fb-hCG 66.3 ng/ml	1.55	Scan date 13-06-2019		
		Crown rump length in mm 49.0 Nuchal translucency MoM 1.2		
Age risk			Nuchal translucency MoM	
Biochemical T21 risk	1:6294	Nasal bone		present
•		Sonographer		
Trisomy 13/18 + NT			Qualifications in measuring NT MD Trisomy 21	
1:100 1:1000 1:10000 1:10000 1:10000 1:10000 Trisomy 13/18 + NT The calculated risk for trisomy 13 translucency) is < 1:10000, which risk.	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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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