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

5.0.2.37

Date of report: 26-03-2019

Prisca

Patient data					
Name MRS. PRIKSHA		Patient ID			
Birthday	01-07-1996		2	211903250007	
ge at sample date 22.9		Sample Date 25-03-201		25-03-2019	
Gestational age 12 + 5					
Correction factors					
Fetuses 1	IVF	unknown	Previous trisomy 21	unknown	
Weight 64	diabetes	unknown	pregancies		
Smoker unknown	Origin	Asian			
Biochemical data	ľ		Ultrasound data		
Parameter Value	Corr. MoM Gestational age 12 + 5				
PAPP-A 6.5 mIU/m	l 1.23	1.23 Method LMP			
fb-hCG 72.1 ng/ml	1.73 Scan date				
Risks at sampling date		Crown rump length in mm			
Age risk			Nuchal translucency MoM		
Biochemical T21 risk			Nasal bone unknow		
Combined trisomy 21 risk <1:10000 Trisomy 13/18 + NT <1:10000		0 1			
Trisomy 13/18 + NT	Qualifications in measuring NT MD				
Risk 1:10 1:100 1:250 Cutoff 1:000 1:1000 1:10000 1:10000 1:10000 Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.		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 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