KOS DIAGNOSTIC LAB 6349/1 NICHOLSON ROAD, AMBALA CANTT

Patient data						
Name	MRS. MANJIT		Patient ID		2209220487/AMB	
Birthday	14-03-1995		Sample ID		2209220487/AMB	
Age at sample date	27.5		Sample Date	e	19-09-2022	
Gestational age	11 + 0					
Correction factors						
Fetuses	1	IVF	no	Previous trisomy 21	no	
Weight	56.8	diabetes	no pregnancies			
Smoker	no	Origin	Asian			
Biochemical data			Ultrasound data			
Parameter	Value	Corr. MoN	Gestational age 10 + 5			
PAPP-A	0.65 mIU/m	nl 0.33	Method CRL Robinson			
fb-hCG	30.1 ng/ml	0.55				
Risks at sampling date			Crown rump	length in mm	41.1	
Age risk		1:779	Nuchal trans	slucency MoM	0.86	
Biochemical T21 risk				· · · · · · · ·		
•			Sonographer .			
-			Qualifications in measuring NT MD			
Risk 1:10			Trisomy 21	Trisomy 21 The calculated risk for Trisomy 21 (with nuchal		
1:1000 1:10000 1:110000 1:110000 1:10000 1:10000 1:10000 Age			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 5571 women with the same data, there is one woman with a trisomy 21 pregnancy and 5570 women with not affected pregnancies. The PAPP-A level is low. 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!			
Trisomy 13/18 + NT						
The calculated risk for translucency) is 1:62						
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Sign of Physician