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

Prisca 5.0.2.37

Date of report: 25-02-2019

Patient data				
Name	MRS. NEHA		Patient ID	
Birthday	19-03-1997		131902230005	
Age at sample date	22	Sample Date	23-02-2019	
Gestational age	12 + 5			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21 no	
Weight 60	diabetes	no	pregancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound data		
Parameter Value	Corr. MoM	Gestational a	age	
PAPP-A 4.74 mIU/n	nl 1.33	Method	LMP	
fb-hCG 30.3 ng/ml	0.97	Scan date		
Risks at sampling date		Crown rump length in mm		
Age risk	1:1083	Nuchal translucency MoM		
Biochemical T21 risk	<1:10000	Nasal bone present		
Combined trisomy 21 risk	<1:10000	Sonographer		
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT MD		
Risk 1:10 Trisomy 21 The calculated risk for Trisomy 21 (with nuchal				
1:: 000 1::1000 1::1000 1::1000 1::1000 1::1000 1::1000 1::1000 Age Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.		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

below cut off Below Cut Off, but above Age Risk

above cut off