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

5.0.2.37

Date of report: 02-05-2019

Prisca

Patient data				
Name	MRS. NEETOO		181905010019	
Birthday	06-05-1996		181905010019	
Age at sample date	23		Sample Date 01-05-2019	
Gestational age	12 + 0			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21 no	
Weight 44.4	diabetes	no	pregancies	
Smoker no	Origin	Asian		
Biochemical data	emical data		Ultrasound data	
Parameter Value	Corr. MoM	Gestational	age 12 + 0	
PAPP-A 2.94 mIU/m	nl 1.29	Method LMP		
fb-hCG 75.2 ng/ml	1.97 Scan date			
Risks at sampling date			Crown rump length in mm	
Age risk	1:999			
Biochemical T21 risk	1:2075			
Combined trisomy 21 risk				
Trisomy 13/18 + NT			Qualifications in measuring NT MD	
			Trisomy 21 The calculated risk for Trisomy 21 (with nuchal	
1:100 1:250 Cut off 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:10000 1:10000 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 3154 women with the same data, there is one woman with a trisomy 21 pregnancy and 3153 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