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

5.0.2.37

Date of report: 03-05-2017

Prisca

Patient data		r		
Name	MRS. MEENA			1704220781/AMB
Birthday	25-04-1976			1704220781/AMB
Age at sample date	41.0			29-04-2017
Gestational age	12 + 3			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 78.2	diabetes	no	pregancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound da	ata	
Parameter Value	Corr. MoM	Gestational age 12 + 2		
PAPP-A 5.5 mIU/m	l 2.28	Method CRL Robinson		
fb-hCG 110.1 ng/ml	3.32	Scan date 28-04-2017		
Risks at sampling date	date		length in mm	60.2
Age risk	1:60		Nuchal translucency MoM 1.9	
Biochemical T21 risk	1:66			present
Combined trisomy 21 risk			r	
Trisomy 13/18 + NT	<1:10000	Qualification	s in measuring NT	MD
Risk 1:10	1	Trisomy 21	ated risk for Trisomy 21	
1:100 1:250 1:1000 1:1000 1:10000 1:10000 1:15 1719 212325 2729 3 Trisomy 13/18 + NT The calculated risk for trisomy 13 translucency) is < 1:10000, which	translucency) is above the cut off, which indicates an increased risk. After the result of the Trisomy 21 Test (with nuchal translucency), it is expected that among less than 50 pregnancies with the same data, there is one trisomy 21 pregnancy. The free beta HCG level is high. 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

above cut off