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

Date of report: 22-09-2016

Patient data		I		
Name	MRS. ANUSHA			1609202011/AMB
Birthday	30-09-1991			1609202011/AMB
Age at sample date	25.0	Sample Date 21-09-201		21-09-2016
Gestational age	12 + 5			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 41	diabetes	no	pregancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound data		
Parameter Value	Corr. MoM	Gestational	age	12 + 4
PAPP-A 2.8 mIU/n	nl 0.49	Method		CRL Robinson
fb-hCG 55.5 ng/ml	1.41	Scan date		20-09-2016
Risks at sampling date		Crown rump length in mm		63.7
Age risk	1:969	Nuchal translucency MoM 1.10		
Biochemical T21 risk	1:481	Nasal bone present		
Combined trisomy 21 risk	1:1620	Sonographer .		
Trisomy 13/18 + NT	<1:10000	Qualification	s in measuring NT	MD
Risk 1¦10		Trisomy 21	ited risk for Trisomy 21	
1:100 1:250 1:1000 1:1000 1:10000	Age /18 (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 1620 women with the same data, there is one woman with a trisomy 21 pregnancy and 1619 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

above cut off