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

5.0.2.37

Date of report: 22-12-2018

Prisca

Patient data				
Name	Ms.DAVINDER			211812210001
Birthday	19-11-1990	Sample ID 2		211812210001
Age at sample date	28.0	Sample Date		21-12-2018
Gestational age	12 + 5			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 69.2	2 diabetes	no	pregancies	
Smoker no	Origin	Asian		
chemical data		Ultrasound data		
Parameter Value	Corr. MoM	Gestational age 12 + 5		
PAPP-A 4.99 mIU/m	l 1.25	Method CRL Robinson		
fb-hCG 60.6 ng/ml	1.40	Scan date 20-12-2018		
Risks at sampling date		Crown rump	length in mm	
Age risk	1:987	Nuchal translucency MoM		
Biochemical T21 risk	1:4430	Nasal bone present		
Combined trisomy 21 risk	1:4827	Sonographer		
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT MD		
Risk 1:10		Trisomy 21	ated risk for Trisomy 21 (
1: 00 1250 1: 100 1: 100 1: 1000 1: 10000 1: 100000 1: 10000 1: 10000 1: 10000 1:		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 4827 women with the same data, there is one woman with a trisomy 21 pregnancy and 4826 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