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

5.1.0.17

Date of report: 24-08-2019

Prisca

Patient data					
Name	MRS. JASWANT		19	008221396/AMB	
Birthday	22-01-1992	Sample ID	1908221396/AMB		
Age at delivery	ery 28.1		Sample Date 23-08-2		
Gestational age 12 + 5					
Correction factors					
Fetuses 1	IVF	no	Previous trisomy 21	no	
Weight 57	diabetes	no pregnancies			
Smoker no	Origin	Asian			
Biochemical data		Ultrasound data			
Parameter Value	Corr. MoM Gestational age 12		12 + 2		
PAPP-A 7.5 mIU/m	nl 1.90	Method		CRL Robinson	
fb-hCG 22.5 ng/ml	0.65	Scan date		20-08-2019	
isks at term		Crown rump	length in mm	60	
Age risk	1:1155	Nuchal trans	chal translucency MoM 0		
Biochemical T21 risk	<1:10000	Nasal bone		present	
Combined trisomy 21 risk <1:10000		Sonographer			
Trisomy 13/18 + NT	<1:10000	Qualification	ns in measuring NT	MD	
			Trisomy 21 The calculated risk for Trisomy 21 (with nuchal		
1:100 1:250 Cut off 1:000 Cut off 1:1000 Cut off 1:10000 Cut off <		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