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

Prisca 5.0.2.37

Date of report: 10-05-2018

Patient data					
Name M	MRS. KHUSHMINDER				
Birthday	29-01-1992	Sample ID		1805220200/AMB	
Age at sample date	26.3	Sample Date		09-05-2018	
Gestational age	12 + 1				
Correction factors					
Fetuses 1	IVF	no	Previous trisomy 21	no	
Weight 69	diabetes	no	pregancies		
Smoker no	Origin	Asian			
Biochemical data		Ultrasound data			
Parameter Value	Corr. MoM	M Gestational age		12 + 0	
PAPP-A 3.8 mIU/m	nl 1.51	Method CRL Robinson			
fb-hCG 75.1 ng/ml	2.08	Scan date 08-05-2018			
Risks at sampling date	C		length in mm	56.6	
Age risk	1:888	Nuchal translucency MoM		1.21	
Biochemical T21 risk	1:2146	Nasal bone pro		present	
Combined trisomy 21 risk	somy 21 risk 1:4321		Sonographer		
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT		MD	
			Trisomy 21 The calculated risk for Trisomy 21 (with nuchal		
1: 00 1: 50 1: 100 1: 1000 1: 10000 1: 10000		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 4321 women with the same data, there is one woman with a trisomy 21 pregnancy and 4320 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