## KOS DIAGNOSTIC LAB 63491/1, NICHOLSON ROAD, AMBALA CANTT

5.0.2.37

Date of report: 29-04-2017

Prisca

Patient data			·	
Name	MRS. RAJNI			1704220745/AMB
Birthday	24-08-1986		1704220745/AM	
Age at sample date	30.7		Sample Date 28-04-20	
Gestational age	12 + 2			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 43.4	diabetes	no	pregancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound da	ta	
Parameter Value	Corr. MoM	Gestational age 11 + 6		
PAPP-A 4.2 mIU/m	ıl 0.92	Method CRL Robinson		
fb-hCG 64.6 ng/ml	1.56	Scan date 25-04-2017		
Risks at sampling date	<s at="" date<="" sampling="" td=""><td colspan="2">Crown rump length in mm 55</td></s>		Crown rump length in mm 55	
Age risk	1:586		Nuchal translucency MoM 0.96	
Biochemical T21 risk	1:1074			present
Combined trisomy 21 risk	-			
Trisomy 13/18 + NT	<1:10000	Qualification	s in measuring NT	MD
Risk 1:10	1.	Trisomy 21	ted risk for Trisomy 21 (v	
1 100 1 250 1: 1000 1: 1000 1: 1000 1: 10000 1: 10000 Trisomy 13/18 + NT The calculated risk for trisomy 13 translucency) is < 1:10000, which risk.	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 4717 women with the same data, there is one woman with a trisomy 21 pregnancy and 4716 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

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