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

5.0.2.37

Date of report: 08-04-2017

Prisca

Patient data		•		
Name	MRS. POOJA	Patient ID		
Birthday	05-04-1991	Sample ID 1704220183/AMB		
Age at sample date	26.0			07-04-2017
Gestational age	12 + 4			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 48	diabetes	no	pregancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound da	ta	
Parameter Value	Corr. MoM	Gestational age 12 + 3		
PAPP-A 6.5 mIU/m	nl 1.42	Method CRL Robinson		
fb-hCG 102.1 ng/ml	2.70	Scan date 06-04-2017		
Risks at sampling date		Crown rump	length in mm	62.6
Age risk	1:916	Nuchal trans	luchal translucency MoM 0.62	
Biochemical T21 risk	1:1007	Nasal bone		present
		Sonographe	r	
Trisomy 13/18 + NT	<1:10000	Qualification	s in measuring NT	MD
Risk 1:10	1.	Trisomy 21	ted risk for Trisomy 21 (w	
1100 1250 Cut off 1:1000 1:1000 1:10000 1:10000 1:10000 Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low		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 5254 women with the same data, there is one woman with a trisomy 21 pregnancy and 5253 women with not affected pregnancies. The free beta HCG level is high. 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