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

5.0.2.37

Date of report: 19-05-2019

Prisca

Patient data		-		
Name MRS.	MRS. CHARANJEET KAUR			
Birthday	24-08-1993		Sample ID 2119	
Age at sample date	25.0	Sample Date		18-05-2019
Gestational age	12 + 6			
Correction factors		-		
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 58.5	diabetes	no	pregancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound data		
Parameter Value	Corr. MoM	Gestational age 12 + 6		
PAPP-A 4.80 mIU/m	nl 1.33	Method CRL Robinson		
fb-hCG 30.7 ng/ml	0.97	Scan date 18-05-2019		
Risks at sampling date	at sampling date		Crown rump length in mm 64.3	
Age risk 1:1060		Nuchal translucency MoM 0.60		
Biochemical T21 risk	<1:10000	Nasal bone		present
Combined trisomy 21 risk <1:10000		Sonographer		
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT		MD
Risk 1:10 1:00 1:250 Cut off 1:1000 1:1000 1:1000 1:1000 1:1 000 1:1 000 1:1 000 1:1 000 Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.		Trisomy 21 The calculated risk for Trisomy 21 (with nuchal 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