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

5.1.0.17

Date of report: 14-08-2019

Prisca

Patient data				
Name M	MRS. KULJEET KAUR		1908220723/AMB	
Birthday	17-01-1992		1908220723/AMB	
Age at delivery	28.1		Sample Date 13-08-2019	
Gestational age	13 + 3			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21 no	
Weight 52	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound data		
Parameter Value	Value Corr. MoM C		Gestational age 12 + 4	
PAPP-A 10.1 mIU/m	ıl 1.78	8 Method CRL Robinson		
fb-hCG 49.3 ng/ml	1.56	1.56 Scan date 07-08-2019		
Risks at term		Crown rump length in mm 63		
Age risk	1:1158	Nuchal translucency MoM 0.		
Biochemical T21 risk	1:7711	Nasal bone present		
Combined trisomy 21 risk <1:10000		Sonographer .		
Trisomy 13/18 + NT			Qualifications in measuring NT MD	
Risk 1:10		Trisomy 21	ated risk for Trisomy 21 (with nuchal	
1:100 1:250 Cutoff 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:10000 1:10000 With nuchal translucency) is < 1:10000, which represents a low 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 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