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

 Prisca
 5.1.0.17

 Date of report:
 31-07-2021

Patient data		1			
Name	MRS. JAGMOHAN			2107220966/AMB	
Birthday	25-04-1987			2107220966/AMB	
Age at sample date	34.3	Sample Date	9	30-07-2021	
Gestational age	13 + 4				
Correction factors					
Fetuses 1	IVF	no	Previous trisomy 21	no	
Weight 78	diabetes	no	pregnancies		
Smoker no	Origin	Asian			
Biochemical data		Ultrasound da	ata		
Parameter Value	Corr. MoM	Gestational age 13 + 3			
PAPP-A 3.75 mIU/m	I 0.85	Method CRL Robinson			
fb-hCG 46.6 ng/ml	1.27	Scan date 29-07-2021			
Risks at sampling date			Crown rump length in mm 75.		
Age risk	1:330		Nuchal translucency MoM 0.6		
Biochemical T21 risk	1:840	Nasal bone		present	
Combined trisomy 21 risk	somy 21 risk 1:4506		Sonographer		
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT		MD	
1:10 1:100 1:250 Cut off 1:1000 1:10000			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 4506 women with the same data, there is one woman with a trisomy 21 pregnancy and 4505 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