Prisca 5.2.0.13

Date of report: 24/09/2024

Patient data				
me MRS. RADHIKA GOYAE		Patient ID		
Birthday	19/10/1997		2409050738	/NOD
Age at sample date	te 26.9		Sample Date 24/09	
Gestational age	12 + 3			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 60	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound data		
Parameter Value	Value Corr. MoM		Gestational age 12 + 2	
PAPP-A 5.1 mIU/m	1.33	Method CRL Robinson		
fb-hCG 35.1 ng/ml	0.77			
Risks at sampling date				58.9
Age risk			Nuchal translucency MoM 0.7	
Biochemical T21 risk			Nasal bone preser	
Combined trisomy 21 risk <1:10000		Sonographer		
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT Trisomy 21		
1:10 1:250 Cut off 1:10000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.		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