Prisca 5.2.0.13

Date of report: 22/06/2024

Patient data					
Name M	MRS. GUNJAN GARG				
Birthday	8/06/1996			2406039224/NOD	
Age at sample date	28.0		е	21/06/2024	
Gestational age	13 +	3			
Correction factors					
Fetuses 1	IVF	no	Previous trisomy 21	no	
Weight 59	diabetes	no	pregnancies		
Smoker no	Origin	Asian			
Biochemical data	Ultrasound data				
Parameter Value	Corr. Mo	M Gestational	Gestational age 13 +		
PAPP-A 5.1 mIU/m	I 0.8	7 Method	Method CRL Robinson		
fb-hCG 47.1 ng/ml	1.	6 Scan date	Scan date 21/06/2024		
Risks at sampling date		Crown rum	o length in mm	75.6	
Age risk	1:8		Nuchal translucency MoM 0.70 Nasal bone present		
Biochemical T21 risk				present	
Combined trisomy 21 risk	00 Sonograph				
Trisomy 13/18 + NT	<1:100		Qualifications in measuring NT Trisomy 21		
1:1000 1:250 Cut off 1:10000 1:10000 1:10000 1:10000 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!		