Prisca 5.2.0.13

Date of report: 14/05/2025

Patient data				
Name	MRS. GAYATRI			
Birthday	25/06/2000	Sample ID		2505024807/NOD
Age at sample date	24.9	Sample Date)	13/05/2025
Gestational age	12 + 1	·		
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	n
Weight 56.6	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound da	ıta	
Parameter Value	Corr. MoM	Gestational age 12 + (
PAPP-A 5.3 mIU/m	l 1.47	Method CRL Robinso		
fb-hCG 56.5 ng/ml	1.18	Scan date 12/05/20		
Risks at sampling date		Crown rump length in mm		56.
Age risk	1:953	Nuchal translucency MoM		1.4
Biochemical T21 risk	1:8689	Nasal bone pres		
Combined trisomy 21 risk	1:6529	Sonographer		DR. ANU KAUSHII
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT		M.I
1:100 1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 38 Trisomy 13/18 + NT The calculated risk for trisomy 13 translucency) is < 1:10000, which 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 6529 women with the same data, there is one woman with a trisomy 21 pregnancy and 6528 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 Diagr 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