

Prisca 5.1.0.17
Date of report: 21-06-2025

N A

Patient data			
Name	Mrs. ROOPAL YADAV	Patient ID	0852506190072
Birthday	20-12-1991	Sample ID	B3012465
Age at sample date	33.5	Sample Date	19-06-2025
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	68	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	1.94 mIU/mL	0.56	
fb-hCG	42.41 ng/mL	1.07	
Risks at sampling date		Gestational age	
Age risk		12 + 3	
Biochemical T21 risk		Method	
Combined trisomy 21 risk		CRL Robinson	
Trisomy 13/18 + NT		Scan date	
		18-06-2025	
		Crown rump length in mm	
		62	
		Nuchal translucency MoM	
		1.12	
		Nasal bone	
		present	
		Sonographer	
		N A	
		Qualifications in measuring NT	
		MD	
Risk		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 1546 women with the same data, there is one woman with a trisomy 21 pregnancy and 1545 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!	
Trisomy 13/18 + NT			
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			

Sign of Physician