

Prisca 5.1.0.17
Date of report: 16-11-2025

NA

Patient data				
Name	Mrs. PURNIMA VISHWAKARMA		Patient ID	0622511140008
Birthday	15-11-1993		Sample ID	A1445238
Age at sample date	32.0		Sample Date	14-11-2025
Gestational age	13 + 3			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies
Weight	68	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 6
PAPP-A	1.58 mIU/mL	0.33	Method	CRL Robinson
fb-hCG	32.8 ng/mL	1.03	Scan date	10-11-2025
Risks at sampling date			Crown rump length in mm	67.6
Age risk	1:500		Nuchal translucency MoM	0.76
Biochemical T21 risk	1:169		Nasal bone	present
Combined trisomy 21 risk	1:1134		Sonographer	NA
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT	NA
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 1134 women with the same data, there is one woman with a trisomy 21 pregnancy and 1133 women with not affected pregnancies. The PAPP-A level is low. 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