

Prisca 5.1.0.17
Date of report: 27-01-2026

Patient data			
Name	Mrs. PRIYANKA VAHULE	Patient ID	0672601260091
Birthday	02-01-2006	Sample ID	B4415522
Age at sample date	20.1	Sample Date	26-01-2026
Gestational age	13 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	53	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	5.78 mIU/mL	0.94	Gestational age 12 + 6
fb-hCG	34.12 ng/mL	0.95	Method CRL Robinson
			Scan date 23-01-2026
Risks at sampling date			Crown rump length in mm 66.74
Age risk		1:1115	Nuchal translucency MoM 1.34
Biochemical T21 risk		1:6891	Nasal bone present
Combined trisomy 21 risk		1:9308	Sonographer NA
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT Sonographer
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 9308 women with the same data, there is one woman with a trisomy 21 pregnancy and 9307 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

below cut off
 Below Cut Off, but above Age Risk
 above cut off