

Prisca 5.1.0.17
Date of report: 19-01-2026

Patient data			
Name	Mrs. SHILPA VERMA		Patient ID 0442601170007
Birthday	16-03-1994		Sample ID B4115546
Age at sample date	31.8		Sample Date 17-01-2026
Gestational age	11 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	73.2	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 11 + 5
PAPP-A	2.04 mIU/mL	0.87	Method CRL Robinson
fb-hCG	45.83 ng/mL	1.00	Scan date 16-01-2026
Risks at sampling date			Crown rump length in mm 52.4
Age risk 1:484			Nuchal translucency MoM 1.07
Biochemical T21 risk 1:2209			Nasal bone present
Combined trisomy 21 risk 1:7417			Sonographer NA
Trisomy 13/18 + NT <1:10000			Qualifications in measuring NT Sonographer
Risk			Trisomy 21
Age			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 7417 women with the same data, there is one woman with a trisomy 21 pregnancy and 7416 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