

Prisca 5.1.0.17
Date of report: 28-01-2026

Patient data			
Name	Mrs. SPANDANA	Patient ID	0042601270009
Birthday	16-01-1991	Sample ID	B3701909
Age at sample date	35.0	Sample Date	27-01-2026
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	76	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.72 mIU/mL	1.10	
fb-hCG	38.1 ng/mL	1.06	
Risks at sampling date			
Age risk	1:273	Crown rump length in mm	68.7
Biochemical T21 risk	1:1807	Nuchal translucency MoM	0.75
Combined trisomy 21 risk	1:8871	Nasal bone	present
Trisomy 13/18 + NT	<1:10000	Sonographer	NA
		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 8871 women with the same data, there is one woman with a trisomy 21 pregnancy and 8870 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 held 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