

Prisca 5.1.0.17
Date of report: 21-01-2026

SWATHI

Patient data			
Name	Mrs. POOJA.	Patient ID	0012601190352.
Birthday	20-06-1998	Sample ID	B4283346.
Age at sample date	27.6	Sample Date	19-01-2026
Gestational age	10 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	57	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	1.9 mIU/mL	0.99	
fb-hCG	54 ng/mL	0.90	
Risks at sampling date			
Age risk	1:770	Gestational age	10 + 6
Biochemical T21 risk	1:5899	Method	CRL Robinson
Combined trisomy 21 risk	1:1942	Scan date	19-01-2026
Trisomy 13/18 + NT	<1:10000	Crown rump length in mm	41.9
		Nuchal translucency MoM	0.85
		Nasal bone	absent
		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 1942 women with the same data, there is one woman with a trisomy 21 pregnancy and 1941 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