

Prisca 5.1.0.17
Date of report: 11-01-2026

Patient data				
Name	Mrs. B.MAHESHWARI		Patient ID	0902601100034
Birthday	06-03-2003		Sample ID	A1976740
Age at sample date	22.9		Sample Date	10-01-2026
Gestational age	12 + 1			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies
Weight	51	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	11 + 6
PAPP-A	1.31 mIU/mL	0.32	Method	CRL Robinson
fb-hCG	43.52 ng/mL	0.90	Scan date	08-01-2026
Risks at sampling date			Crown rump length in mm	
Age risk	1:1020		53.5	
Biochemical T21 risk	1:426		Nuchal translucency MoM	
Combined trisomy 21 risk	1:1425		1.12	
Trisomy 13/18 + NT	1:8064		Nasal bone	
			present	
			Sonographer	
			NA	
			Qualifications in measuring NT	
			Sonographer	
Trisomy 21				
<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 1425 women with the same data, there is one woman with a trisomy 21 pregnancy and 1424 women with not affected pregnancies.</p> <p>The PAPP-A level is low.</p> <p>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!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>				
Trisomy 13/18 + NT				
<p>The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:8064, which represents a low risk.</p>				

Sign of Physician