

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	unknown			
Weight	51	diabetes	no					
Smoker	no	Origin	Asian					
Biochemical data								
Parameter	Value	Corr. MoM	Ultrasound data					
PAPP-A	1.31 mIU/mL	0.32	Gestational age 11 + 6					
fb-hCG	43.52 ng/mL	0.90	Method CRL Robinson					
Risks at sampling date								
Age risk	1:1020		Scan date 08-01-2026					
Biochemical T21 risk	1:426		Crown rump length in mm 53.5					
Combined trisomy 21 risk	1:1425		Nuchal translucency MoM 1.12					
Trisomy 13/18 + NT	1:8064		Nasal bone present					
Sonographer NA								
Qualifications in measuring NT Sonographer								
Trisomy 21								
The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk. <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.</p> <p>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 held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>								
Trisomy 13/18 + NT								
The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:8064, which represents a low risk.								

Sign of Physician