

Prisca 5.1.0.17
Date of report: 21-01-2026

Patient data			
Name	Mrs. BHAGYAREKA	Patient ID	0842601200058
Birthday	05-10-1991	Sample ID	B4141938
Age at sample date	34.3	Sample Date	20-01-2026
Gestational age	11 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	46	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	1.63 mIU/mL	0.43	
fb-hCG	46.47 ng/mL	0.84	
Risks at sampling date		Gestational age 11 + 5	
Age risk	1:306	Method CRL Robinson	
Biochemical T21 risk	1:343	Scan date 20-01-2026	
Combined trisomy 21 risk	1:2079	Crown rump length in mm 53	
Trisomy 13/18 + NT	<1:10000	Nuchal translucency MoM 0.64	
		Nasal bone present	
		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 2079 women with the same data, there is one woman with a trisomy 21 pregnancy and 2078 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