

Prisca 5.1.0.17
Date of report: 12-01-2026

Patient data				
Name	Mrs. PRATIKSHA GURLE		Patient ID	0372601110018
Birthday	14-09-2002		Sample ID	B4262145
Age at sample date	23.3		Sample Date	11-01-2026
Gestational age	13 + 5			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies
Weight	64	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 6
PAPP-A	5.32 mIU/mL	0.94	Method	CRL Robinson
fb-hCG	30.2 ng/mL	1.01	Scan date	05-01-2026
Risks at sampling date			Crown rump length in mm	
Age risk	1:1061		68.1	
Biochemical T21 risk	1:5549		Nuchal translucency MoM	
Combined trisomy 21 risk	<1:10000		0.64	
Trisomy 13/18 + NT	<1:10000		Nasal bone	
			present	
			Sonographer	
			NA	
			Qualifications in measuring NT	
			Sonographer	
Risk			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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.</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 < 1:10000, which represents a low risk.</p>				

Sign of Physician

below cut off
 Below Cut Off, but above Age Risk
 above cut off