

Prisca 5.1.0.17
Date of report: 16-01-2026

Patient data			
Name	Ms. SEEMA KUMARI		Patient ID
Birthday	14-10-2002		Sample ID
Age at sample date	23.3		Sample Date
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	41	diabetes	no
Smoker	no	Origin	Asian
			Previous trisomy 21 pregnancies
			unknown
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.71 mIU/mL	0.23	12 + 6
fb-hCG	38 ng/mL	0.89	Method
			CRL Robinson
			Scan date
			14-01-2026
			Crown rump length in mm
			68.4
			Nuchal translucency MoM
			1.04
			Nasal bone
			unknown
			Sonographer
			NA
			Qualifications in measuring NT
			Sonographer
Risks at sampling date			
Age risk		1:1040	
Biochemical T21 risk		1:162	
Combined trisomy 21 risk		1:736	
Trisomy 13/18 + NT		1:2418	
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 736 women with the same data, there is one woman with a trisomy 21 pregnancy and 735 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 1:2418, which represents a low risk.</p>			

Sign of Physician

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