

N A

Patient data							
Name	Mrs. KIRAN KUMARI	Patient ID	0472503240107				
Birthday	18-08-1988	Sample ID	B2490696				
Age at sample date	36.6	Sample Date	24-03-2025				
Gestational age	13 + 1						
Correction factors							
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown		
Weight	58	diabetes	no				
Smoker	no	Origin	Asian				
Biochemical data							
Parameter	Value	Corr. MoM					
PAPP-A	3.1 mIU/mL	0.59					
fb-hCG	39.2 ng/mL	1.08					
Risks at sampling date							
Age risk	1:193						
Biochemical T21 risk	1:295						
Combined trisomy 21 risk	1:1702						
Trisomy 13/18 + NT	<1:10000						
Ultrasound data							
			Gestational age	13 + 1			
			Method	CRL Robinson			
			Scan date	24-03-2025			
			Crown rump length in mm	72			
			Nuchal translucency MoM	0.78			
			Nasal bone	unknown			
			Sonographer	N A			
			Qualifications in measuring NT	MD			
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 1702 women with the same data, there is one woman with a trisomy 21 pregnancy and 1701 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