

NA

Patient data			
Name	Mrs. SHALINI KUMARI	Patient ID	0812504180017
Birthday	18-10-2003	Sample ID	A1781454
Age at sample date	21.5	Sample Date	18-04-2025
Gestational age	13 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	68	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	6.28 mIU/mL	1.38	Gestational age 13 + 1
fb-hCG	35.1 ng/mL	1.06	Method CRL Robinson
Risks at sampling date			Scan date 17-04-2025
Age risk		1:1092	Crown rump length in mm 71.7
Biochemical T21 risk		<1:10000	Nuchal translucency MoM 0.86
Combined trisomy 21 risk		<1:10000	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer NA
			Qualifications in measuring NT NA
Risk			Trisomy 21
<p>The graph plots risk on a logarithmic scale from 1:10000 to 1:10 against age from 13 to 49. A diagonal cut-off line separates the risk levels. The patient's risk is indicated by a vertical bar at age 21.5, which is below the cut-off line.</p>			<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