

NA

Patient data			
Name	Mrs. PARIJAT DEVI	Patient ID	0692507080225
Birthday	15-08-1998	Sample ID	B3197341
Age at sample date	26.9	Sample Date	08-07-2025
Gestational age	12 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	61.1	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.33 mIU/mL	0.89	12 + 2
fb-hCG	40.92 ng/mL	0.96	Method
			CRL Robinson
			Scan date
			07-07-2025
Risks at sampling date			Crown rump length in mm
Age risk		1:863	60.4
Biochemical T21 risk		1:4569	Nuchal translucency MoM
Combined trisomy 21 risk		<1:10000	0.96
Trisomy 13/18 + NT		<1:10000	Nasal bone
			present
			Sonographer
			NA
			Qualifications in measuring NT
			NA
Risk			Trisomy 21
1:10	<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>		
1:100			
1:250			
1:1000			
1:10000			
1:10000			
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49	Age		
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