

Prisca 5.1.0.17
Date of report: 24-03-2025

N A

Patient data			
Name	Mrs. SARANYA	Patient ID	0352503210008
Birthday	11-10-1997	Sample ID	B2410705
Age at sample date	27.4	Sample Date	21-03-2025
Gestational age	12 + 0		
Correction factors			
Fetuses	1	IVF	unknown
Weight	62	diabetes	unknown
Smoker	unknown	Origin	Caucasian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.58 mIU/mL	0.99	11 + 6
fb-hCG	45.27 ng/mL	1.01	Method
			CRL Robinson
			Scan date
			20-03-2025
Risks at sampling date		Trisomy 21	
Age risk	1:816	Crown rump length in mm	54
Biochemical T21 risk	1:4846	Nuchal translucency MoM	1.12
Combined trisomy 21 risk	<1:10000	Nasal bone	present
Trisomy 13/18 + NT	<1:10000	Sonographer	N A
		Qualifications in measuring NT	MD
		<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>	
		<p>Trisomy 13/18 + NT</p> <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