

N A

Patient data			
Name	Mrs. SRINIJA	Patient ID	0012505010094
Birthday	17-02-1997	Sample ID	B2528407
Age at sample date	28.2	Sample Date	01-05-2025
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	70	diabetes	no
Smoker	no	Origin	Asian
Biochemical data			
Parameter	Value	Corr. MoM	Ultrasound data
PAPP-A	2.44 mIU/mL	0.62	Gestational age 12 + 6
fb-hCG	35.74 ng/mL	1.01	Method CRL Robinson
Risks at sampling date			
Age risk	1:795		Scan date 30-04-2025
Biochemical T21 risk	1:1569		Crown rump length in mm 68.5
Combined trisomy 21 risk	1:7786		Nuchal translucency MoM 0.93
Trisomy 13/18 + NT	<1:10000		Nasal bone present
			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. <p>After the result of the Trisomy 21 test (with NT) it is expected that among 7786 women with the same data, there is one woman with a trisomy 21 pregnancy and 7785 women with not affected pregnancies.</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 held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>			
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