

Prisca 5.1.0.17
Date of report: 09-03-2025

NA

Patient data			
Name	Mrs. RUPALI RANGARI		Patient ID
Birthday	20-08-1990	Sample ID	0372503080259
Age at sample date	34.5	Sample Date	a1505085
Gestational age	11 + 1		08-03-2025
Correction factors			
Fetuses	1	IVF	no
Weight	55	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.71 mIU/mL	0.73	11 + 0
fb-hCG	53.39 ng/mL	0.93	Method
			CRL Robinson
			Scan date
			07-03-2025
			Crown rump length in mm
			44
			Nuchal translucency MoM
			0.57
			Nasal bone
			present
			Sonographer
			NA
			Qualifications in measuring NT
			MD
Risks at sampling date			
Age risk	1:283		
Biochemical T21 risk	1:1033		
Combined trisomy 21 risk	1:5510		
Trisomy 13/18 + NT	<1:10000		
Risk		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 5510 women with the same data, there is one woman with a trisomy 21 pregnancy and 5509 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 hold 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