

Prisca 5.1.0.17
Date of report: 26-01-2026

Patient data			
Name	Mrs. T ASHRITHA		Patient ID
Birthday	17-04-1991		Sample ID
Age at sample date	34.8		Sample Date
Gestational age	11 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	59.6	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.05 mIU/mL	0.73	11 + 5
fb-hCG	50.89 ng/mL	1.02	Method
			CRL Robinson
			Scan date
			24-01-2026
Risks at sampling date			Trisomy 21
Age risk			1:276
Biochemical T21 risk			1:815
Combined trisomy 21 risk			1:3848
Trisomy 13/18 + NT			<1:10000
			Qualifications in measuring NT
			Sonographer
			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 3848 women with the same data, there is one woman with a trisomy 21 pregnancy and 3847 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 held 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

below cut off
 Below Cut Off, but above Age Risk
 above cut off