

Prisca 5.1.0.17
Date of report: 22-12-2025

Patient data			
Name	Mrs. J.MADHURIMA	Patient ID	0902512220022
Birthday	15-01-1991	Sample ID	A1976782
Age at sample date	34.9	Sample Date	22-12-2025
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	70.05	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	4.34 mIU/mL	1.04	Gestational age 12 + 4
fb-hCG	35.68 ng/mL	1.05	Method CRL Robinson
			Scan date 18-12-2025
Risks at sampling date			Trisomy 21
Age risk	1:282		Crown rump length in mm 64.1
Biochemical T21 risk	1:1726		Nuchal translucency MoM 0.49
Combined trisomy 21 risk	1:8566		Nasal bone present
Trisomy 13/18 + NT	<1:10000		Sonographer NA
			Qualifications in measuring NT Sonographer
Age			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 8566 women with the same data, there is one woman with a trisomy 21 pregnancy and 8565 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