

Prisca 5.1.0.17
Date of report: 19-01-2026

Patient data			
Name	Mrs. RAJUL JAIN	Patient ID	0382601150083
Birthday	12-10-1989	Sample ID	D0094241
Age at sample date	36.3	Sample Date	15-01-2026
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	72	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.57 mIU/mL	0.67	
fb-hCG	35.74 ng/mL	1.02	
Risks at sampling date			
Age risk	1:208	Gestational age	13 + 0
Biochemical T21 risk	1:499	Method	CRL Robinson
Combined trisomy 21 risk	1:2762	Scan date	15-01-2026
Trisomy 13/18 + NT	<1:10000	Crown rump length in mm	70
		Nuchal translucency MoM	0.74
		Nasal bone	present
		Sonographer	NA
		Qualifications in measuring NT	Sonographer
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 2762 women with the same data, there is one woman with a trisomy 21 pregnancy and 2761 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
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