

Prisca 5.1.0.17
Date of report: 29-10-2025

N A

Patient data			
Name	Mrs. RUTUJA INGLE		Patient ID 0662510280037
Birthday	16-03-2001		Sample ID B3861055
Age at sample date	24.6		Sample Date 28-10-2025
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	56	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 12 + 3
PAPP-A	1.81 mIU/mL	0.41	Method CRL Robinson
fb-hCG	42.02 ng/mL	0.99	Scan date 27-10-2025
Risks at sampling date			Crown rump length in mm 60.9
Age risk	1:978		Nuchal translucency MoM 1.08
Biochemical T21 risk	1:692		Nasal bone present
Combined trisomy 21 risk	1:2574		Sonographer N A
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT MD
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 2574 women with the same data, there is one woman with a trisomy 21 pregnancy and 2573 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