

Prisca 5.1.0.17
 Date of report: 03-10-2025

NA

Patient data			
Name	Mrs. TANU SINGH	Patient ID	0772510010118
Birthday	16-07-1998	Sample ID	B3057714
Age at sample date	27.2	Sample Date	01-10-2025
Gestational age	12 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	58	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	4.29 mIU/mL	1.30	
fb-hCG	43.52 ng/mL	0.91	
Risks at sampling date			
Age risk		1:830	
Biochemical T21 risk		<1:10000	
Combined trisomy 21 risk		<1:10000	
Trisomy 13/18 + NT		<1:10000	
			Gestational age 12 + 0 Method CRL Robinson Scan date 01-10-2025 Crown rump length in mm 56.6 Nuchal translucency MoM 0.87 Nasal bone present Sonographer NA Qualifications in measuring NT NA
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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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