

Prisca 5.1.0.17
Date of report: 29-01-2026

Patient data			
Name	Mrs. SANGITA CHAKRABORTY	Patient ID	0692601280131
Birthday	26-09-1992	Sample ID	D0080090
Age at sample date	33.3	Sample Date	28-01-2026
Gestational age	13 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	79.7	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	4.07 mIU/mL	0.98	Gestational age 12 + 6
fb-hCG	32.66 ng/mL	1.12	Method CRL Robinson
Risks at sampling date			Scan date 23-01-2026
Age risk	1:396		Crown rump length in mm 67.7
Biochemical T21 risk	1:1842		Nuchal translucency MoM 0.70
Combined trisomy 21 risk	1:9361		Nasal bone present
Trisomy 13/18 + NT	<1:10000		Sonographer NA
			Qualifications in measuring NT Sonographer
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 9361 women with the same data, there is one woman with a trisomy 21 pregnancy and 9360 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

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