

Prisca 5.1.0.17
Date of report: 25-09-2025

MANISHA SAO

Patient data			
Name	Mrs. SABNAM KUMARI	Patient ID	0622509220043
Birthday	12-12-2007	Sample ID	b2801354
Age at sample date	17.8	Sample Date	22-09-2025
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	42	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.32 mIU/mL	0.34	12 + 4
fb-hCG	39.05 ng/mL	0.89	Method
			CRL Robinson
			Scan date
			20-09-2025
Risks at sampling date			Crown rump length in mm
Age risk		1:1123	64.5
Biochemical T21 risk		1:580	Nuchal translucency MoM
Combined trisomy 21 risk		1:3775	0.57
Trisomy 13/18 + NT		<1:10000	Nasal bone
			present
			Sonographer
			N A
			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 3775 women with the same data, there is one woman with a trisomy 21 pregnancy and 3774 women with not affected pregnancies. The PAPP-A level is low. 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