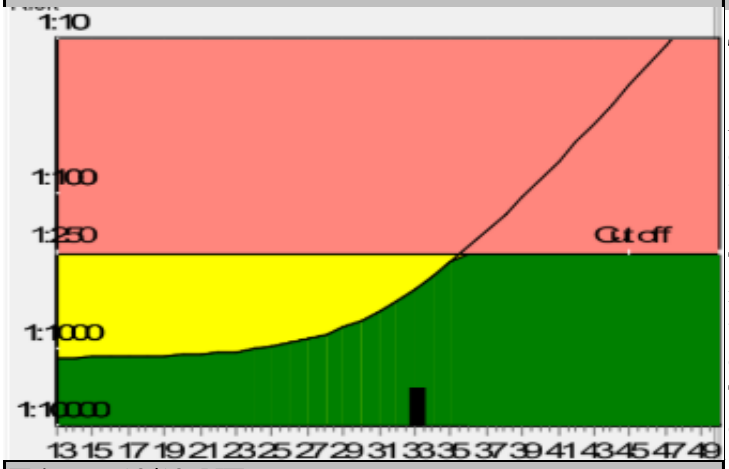
Date of Report 22-01-2025
PRISCA 5.2.0.13

Patient Data			
Name	MRS. TANVI SHRIVASTAVA	Patient ID	12501200195
Birthday	26-12-1991	Sample ID	11959679
Age at Sample date	33.1	Sample Date	20-01-2025
Gestational age	13+2		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	57.2	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+4
PAPP-A	6.7 mIU/ml	0.88	Method	CRL (<>Robinson)
fb-hCG	23.5 ng/ml	0.77	Scan date	16-01-2025
			Crown rump length in mm	61.2
Risks at sampling date			Nuchal translucency MoM	0.81
Age Risk		1:413	Nasal bone	PRESENT
Biochemical T21 risk		1:3469	Sonographer	DR.
Combined trisomy 21 risk		<1:10000	Qualifications in measuring NT	MBBS
Trisomy 13/18 + NT		<1:10000		

Risk



Trisomy 13/18+NT

The calculated risk for Trisomy 13/18 (with NT) is <1:10000 , which indicates a low risk

Down's Syndrome Risk (Trisomy 21 Screening)

The calculated risk for Trisomy 21(with NT) is below the cut off, which represents 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 the risk calculations are statistical aapproaches and have no diagnostic value!

The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998).

The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

Risk Above Cut Off Risk above Age Risk Risk below Age risk