Date of Report	14-01-2025
PRISCA	5.2.0.13

Patient Data			
Name	MRS. RIYA W/O SUMIT	Patient ID	12501130107
Birthday	15-06-1996	Sample ID	11914828
Age at Sample date	28.6	Sample Date	13-01-2025
Gestational age	12+4		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	68.3	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data				Ultrasound Data	
Parameter	Value	Corr Mom		Gestational age	12+4
PAPP-A	6.3 mIU/ml	1.33		Method	CRL (<>Robinson)
fb-hCG	27.8 ng/ml	0.78		Scan date	13-01-2025
Risks at sampling date				Crown rump length in mm	61.6
Age Risk	1:756			Nuchal translucency MoM	0.75
Biochemical T21 risk	<1:10000			Nasal bone	PRESENT
Combined trisomy 21 risk	<1:10000			Sonographer	DR. AMENDA
Trisomy 13/18 + NT	<1:10000			Qualifications in measuring NT	MD

Risk

Down's Syndrome Risk (Trisomy 21 Screening)
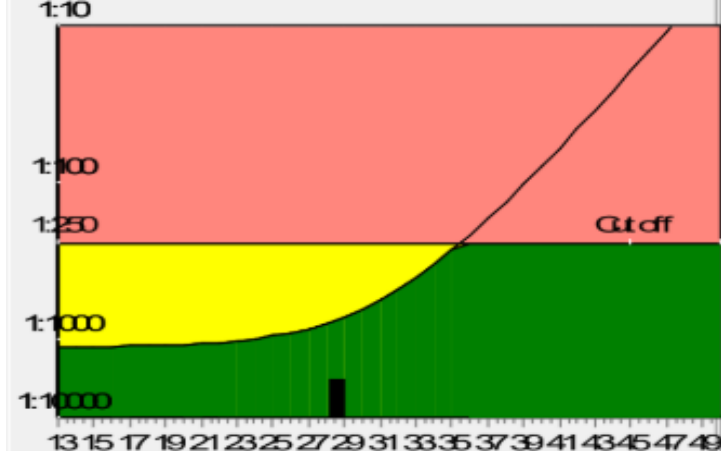
The calculated risk for Trisomy 21(with NT) is below the cut off, which represents a low risk.
After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.

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

Trisomy 13/18+NT

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

🟥 Risk Above Cut Off	🟨 Risk above Age Risk	🟩 Risk below Age risk