		Date of Report	29/3/2023
		PRISCA	5.2.0.13

Patient Data			
Name	**MRS. SARITA**	Patient ID	012303270305
Birthday	28/04/1988	Sample ID	11551776
Age at term	35.3	Sample Date	27/3/2023
Gestational age	12+6		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	52	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom		12+5
PAPP-A	6.5 mIU/ml	0.89	Method	CRL (<>Robinson)
fb-hCG	72.1 ng/ml	2.01	Scan date	27/3/2023
Risks at sampling date			Nuchal translucency	1.2
Age Risk		1:281	Nuchal translucency MoM	0.75
Biochemical T21 risk		1:258	Nasal bone	Present
Combined trisomy 21 risk		1:1456		
Trisomy 13/18 + NT		<1:10000		

Risk	Down's Syndrome Risk (Trisomy 21 Screening)

Down's Syndrome Risk (Trisomy 21 Screening)
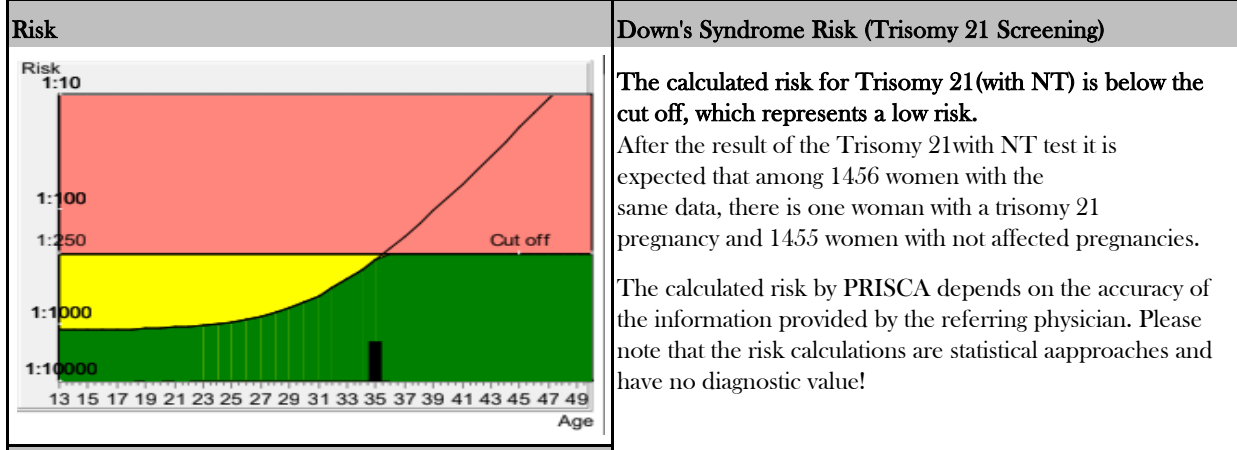
The calculated risk for Trisomy 21(with NT) is below the cut off, which represents a low risk.

After the result of the Trisomy 21with NT test it is expected that among 1456 women with the same data, there is one woman with a trisomy 21 pregnancy and 1455 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 the risk calculations are statistical aapproaches and have no diagnostic value!

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

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