		Date of Report	19-01-2024
		PRISCA	5.2.0.13

Patient Data			
Name	**MRS PREETI**	Patient ID	012401180139
Birthday	25-06-1998	Sample ID	11813714
Age at Sample date	25.6	Sample Date	18-01-2024
Gestational age	13+1		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	60	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data				Ultrasound Data	
Parameter	Value	Corr Mom		Gestational age	13+0
PAPP-A	5.6 mIU/ml	0.82		Method	CRL (<>Robinson)
fb-hCG	31.5 ng/ml	1		Scan date	17-01-2024
Risks at sampling date				Crown rump length in mm	67.6
Age Risk		1:956		Nuchal translucency MoM	0.82
Biochemical T21 risk		1:3807		Nasal bone	PRESENT
Combined trisomy 21 risk		<1:10000		Sonographer	DR VIKRAM PRATAP
Trisomy 13/18 + NT		<1:10000		Qualifications in measuring NT	MD

Risk	Down's Syndrome Risk (Trisomy 21 Screening)

Down's Syndrome Risk (Trisomy 21 Screening)
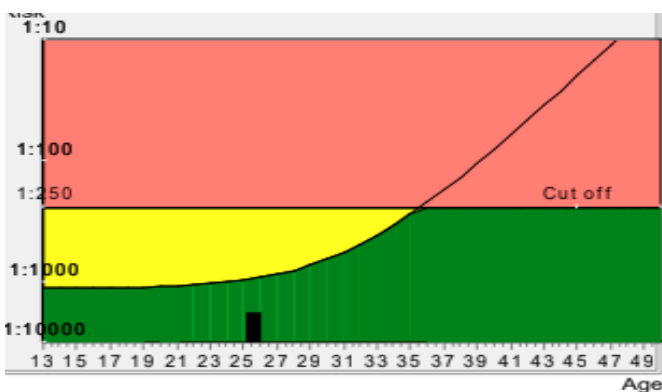
The calculated risk for Trisomy 21(with NT) is below the cut off, which represents a low risk.
After the result of the Trisomy 21 test (with NT) it is expected that among 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 women with not affected pregnancies.
The calculated 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 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

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