	Date of Report	14-02-2024
	PRISCA	5.2.0.13

Patient Data			
Name	**MRS CHANDA**	Patient ID	012402130169
Birthday	01-04-1990	Sample ID	11851226
Age at Sample date	33.9	Sample Date	13-02-2024
Gestational age	13+6		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	59	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+1
PAPP-A	6.8 mIU/ml	0.77	Method	CRL (<>Robinson)
fb-hCG	19.6 ng/ml	0.79	Scan date	08-02-2024

Risks at sampling date		Crown rump length in mm	70
Age Risk	1:361	Nuchal translucency MoM	0.63
Biochemical T21 risk	1:2116	Nasal bone	PRESENT
Combined trisomy 21 risk	<1:10000	Sonographer	DR GARIMA GUPTA
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT	MBBS

Risk

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

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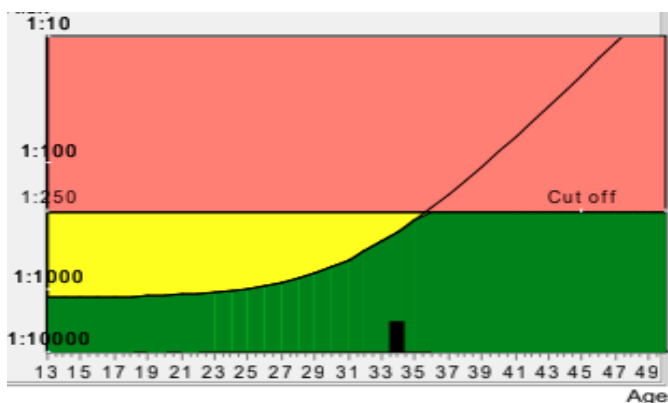
The calculated risk for Trisomy 21(with NT) is below the cut off, which represents a low risk.
After the result of the Trisomy 21 test (with NT) it is expected that among 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 women with not affected pregnancies.
The calculated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

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