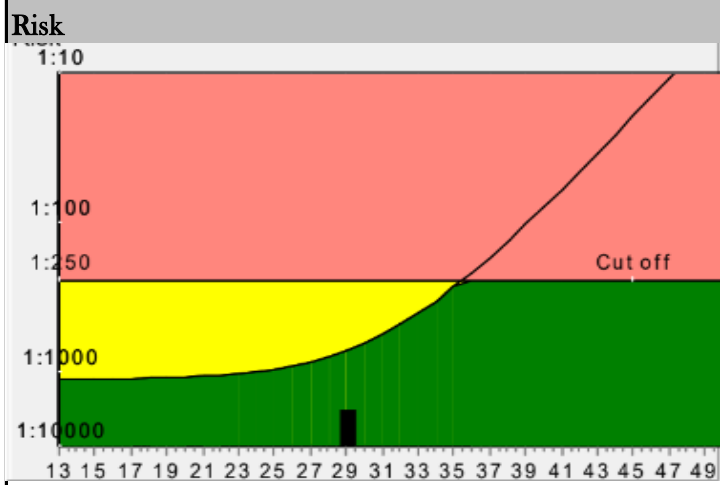


Date of Report 08-10-2024
PRISCA 5.2.0.13

Patient Data			
Name	MRS. CHANCHAL W/O BHARAT	Patient ID	12410070161
Birthday	12-09-1995	Sample ID	11966779
Age at Sample date	29.1	Sample Date	07-10-2024
Gestational age	12+5		

Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	76	Diabetes	NO
Smoker	NO	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+5
PAPP-A	5.1 mIU/ml	1.16	Method	CRL (<>Robinson)
fb-hCG	38.3 ng/ml	1.16	Scan date	07-10-2024
Risks at sampling date			Crown rump length in mm	63.1
Age Risk		1:723	Nuchal translucency MoM	1.05
Biochemical T21 risk		1:4405	Nasal bone	PRESENT
Combined trisomy 21 risk		<1:10000	Sonographer	DR.AMENDA
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MD



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 approaches 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).

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

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