Date of Report	12-12-2024
PRISCA	5.2.0.13

Patient Data

Name	S. BHARTI W/O MANISH KUMAR	Patient ID	12412110117
Birthday	21-09-1999	Sample ID	11947024
Age at Sample date	25.2	Sample Date	11-12-2024
Gestational age	13+0		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	36.9	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	5.1 mIU/ml	0.45
fb-hCG	21.22 ng/ml	0.54

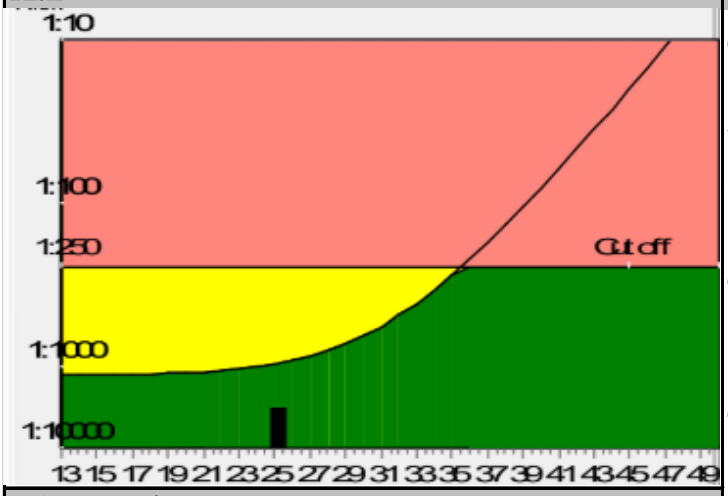
Risks at sampling date

Age Risk	1:967
Biochemical T21 risk	1:3144
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	12+6
Method	CRL (<>Robinson)
Scan date	10-12-2024
Crown rump length in mm	65.5
Nuchal translucency MoM	0.90
Nasal bone	present
Sonographer	DR. AMENDA
Qualifications in measuring NT	MD

Risk



Trisomy 13/18+NT

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

Down's Syndrome Risk (Trisomy 21 Screening)

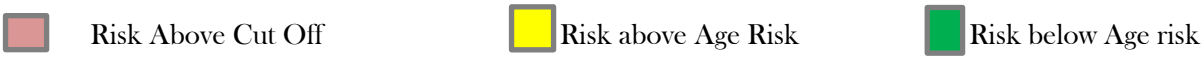
The calculated risk for Trisomy 21(with NT) is below the cut off, which represents a low risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 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


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