Date of Report	09-08-2024
PRISCA	5.2.0.13

Patient Data

Name	MRS. MANISHA	Patient ID	12408080136
Birthday	26-12-1995	Sample ID	11883803
Age at Sample date	28.6	Sample Date	08-08-2024
Gestational age	12+2		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	55	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	5.1 mIU/ml	0.72
fb-hCG	71.7 ng/ml	1.61

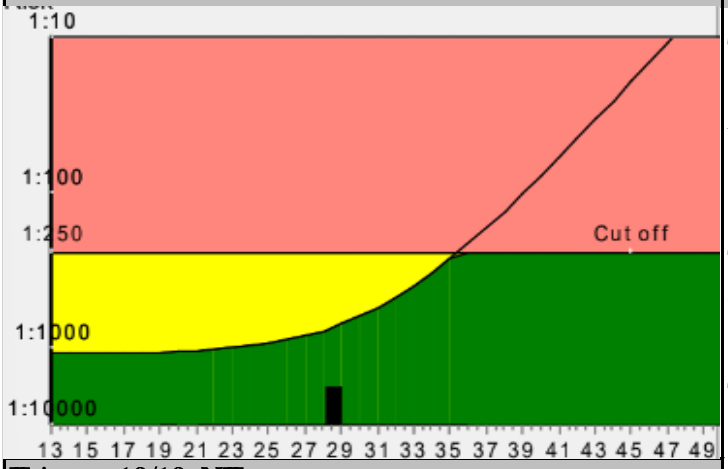
Risks at sampling date

Age Risk	1:745
Biochemical T21 risk	1:728
Combined trisomy 21 risk	1:4189
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	12+2
Method	CRL (<>Robinson)
Scan date	08-08-2024
Crown rump length in mm	56.9
Nuchal translucency MoM	0.80
Nasal bone	PRESENT
Sonographer	DR. RAKHI
Qualifications in measuring NT	MBBS

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)

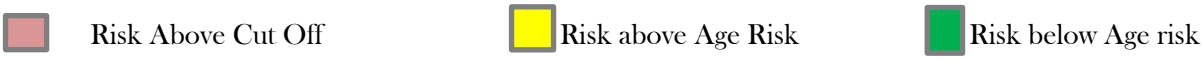
The calculated risk for Trisomy 21(with NT) is below the cut off, which represents a low risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 4189 women with the same data, there is one woman with a trisomy 21 pregnancy and 4188 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