Date of Report 10-05-2024
PRISCA 5.2.0.13

Patient Data

Name	ANAMIKA KUMARI	Patient ID	012405080277
Birthday	02-02-1996	Sample ID	11844526
Age at Sample date	28.3	Sample Date	08-05-2024
Gestational age	12+5		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	74	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	5.6 mIU/ml	1.23
fb-hCG	54.1 ng/ml	1.62

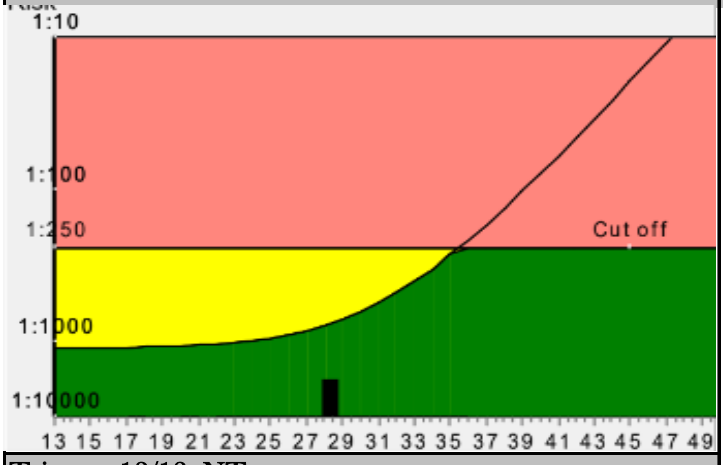
Risks at sampling date

Age Risk	1:782
Bicochemical T21 risk	1:2402
Commbined Trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	12+4
Method	CRL (<>Robinson)
Scan date	08-05-2024
Crown curm length in mm	62.2
Nuchal translucency MoM	0.69
Nasal bone	present
Sonographer	DR.VIKASH GOYAL
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 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