Date of Report 27-03-2024
PRISCA 5.2.0.13

Patient Data			
Name	MRS. PAYAL KUMARI	Patient ID	12403260099
Birthday	28-09-1996	Sample ID	11844534
Age at Sample date	27.5	Sample Date	26-03-2024
Gestational age	11+3		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	46	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	3.1 mIU/ml	0.68
fb-hCG	37.2 ng/ml	0.69

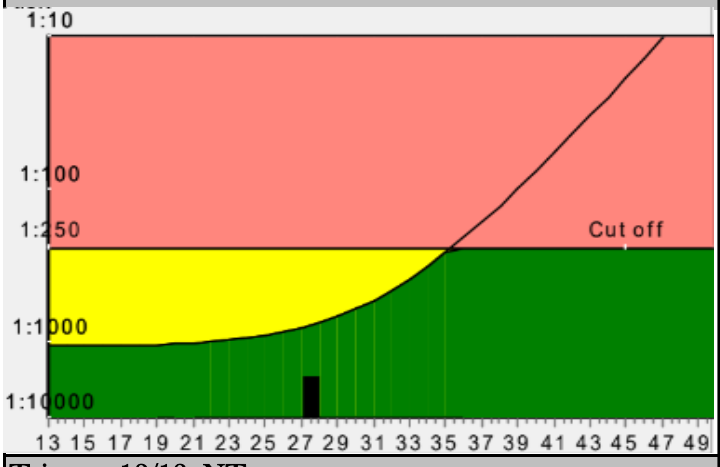
Risks at sampling date

Age Risk	1:795
Biochemical T21 risk	1:4233
Combined trisomy 21 risk	1:1730
Trisomy 13/18 + NT	1:7012

Ultrasound Data

Gestational age	11+3
Method	CRL (<>Robinson)
Scan date	26-03-2024
Crown rump length in mm	46.3
Nuchal translucency MoM	1.65
Nasal bone	PRESENT
Sonographer	DR. VIKAS GOYAL
Qualifications in measuring NT	MBBS

Risk



Trisomy 13/18+NT

The calculated risk for Trisomy 13/18 (with NT) is 1:7012, 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 1730 women with the same data, there is one woman with a trisomy 21 pregnancy and 1729 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