Date of Report	16/05/2024
PRISCA	5.2.0.13

Patient Data			
Name	**SARITA**	Patient ID	012405150157
Birthday	25/07/1991	Sample ID	11485117
Age at sample	32.8	Sample Date	15/05/2024
Gestational age	12+2		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	79	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	4.60 mIU/ml	1.30
fb-hCG	31.5 ng/ml	0.86

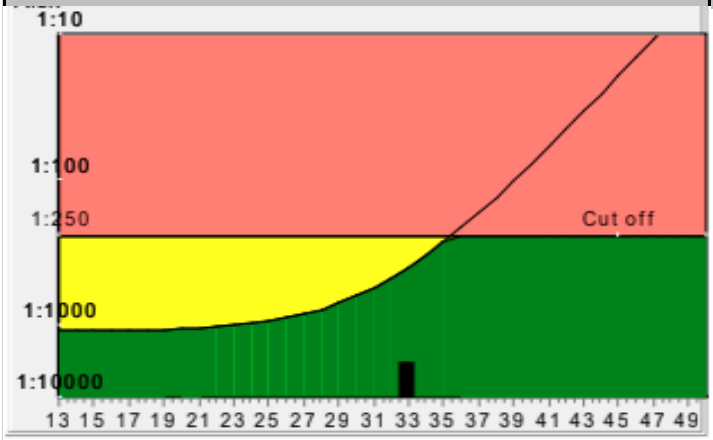
Risks at sampling date

Age Risk	1:418
Biochemical T21 risk	1:6230
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	12+2
Method	CRL (<>Robinson)
Scan date	15/05/2024
Crown rump length in mm	57.3
Nuchal translucency MoM	0.86
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 < 1:10000, 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