Date of Report	5/5/2024
PRISCA	5.2.0.13

Patient Data			
Name	**MANITA**	Patient ID	012405030230
Birthday	13/04/1995	Sample ID	12001852
Age at sample	29.1	Sample Date	3/5/2024
Gestational age	13+0		

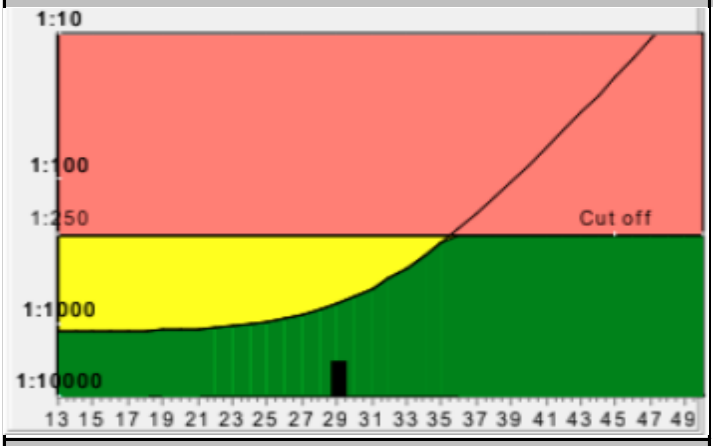
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	55	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data		
Parameter	**Value**	**Corr Mom**
PAPP-A	5.60 mIU/ml	0.78
fb-hCG	28.8 ng/ml	28.8

Risks at sampling date	
Age Risk	1:731
Biochemical T21 risk	1:3692
Combined trisomy 21 risk	1:5813
Trisomy 13/18 + NT	<1:10000

Ultrasound Data	
Gestational age	12+6
Method	CRL (<>Robinson)
Scan date	2/5/2024
Crown rump length in mm	66
Nuchal translucency MoM	1.07
Nasal bone	Absent
Sonographer	DR. HARENDRA BHASKAR
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 5813 women with the same data, there is one woman with a trisomy 21 pregnancy and 5812 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