Date of Report	16/1/2024
PRISCA	5.2.0.13

Patient Data			
Name	MANISHA KUMARI	Patient ID	12401150103
Birthday	29/02/2000	Sample ID	11819512
Age at sample	23.90	Sample Date	15/1/2024
Gestational age	12+1		

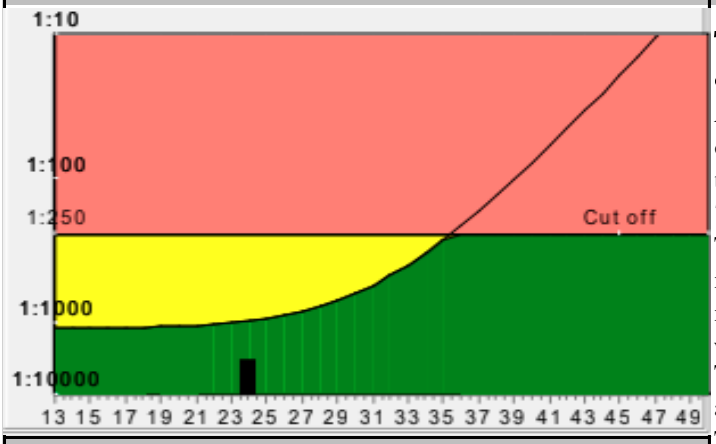
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	54	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data		
Parameter	**Value**	**Corr Mom**
PAPP-A	3.90 mIU/ml	0.74
fb-hCG	45.4 ng/ml	1.06

Risks at sampling date	
Age Risk	1:990
Biochemical T21 risk	1:2707
Combined trisomy 21 risk	1:7050
Trisomy 13/18 + NT	<1:10000

Ultrasound Data	
Gestational age	11+6
Method	CRL (<>Robinson)
Scan date	13/1/2024
Crown rump length in mm	51.2
Nuchal translucency MoM	1.16
Nasal bone	Present
Sonographer	DR.MANJU
Qualifications in measuring NT	MD

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 7050 women with the same data, there is one woman with a trisomy 21 pregnancy and 7049 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