Date of Report 05-12-2023

PRISCA 5.2.0.13

Patient Data			
Name	MRS SANTOSH	Patient ID	012312040061
Birthday	01-02-1994	Sample ID	11848503
Age at Sample date	29.8	Sample Date	04-12-2023
Gestational age	13+0		

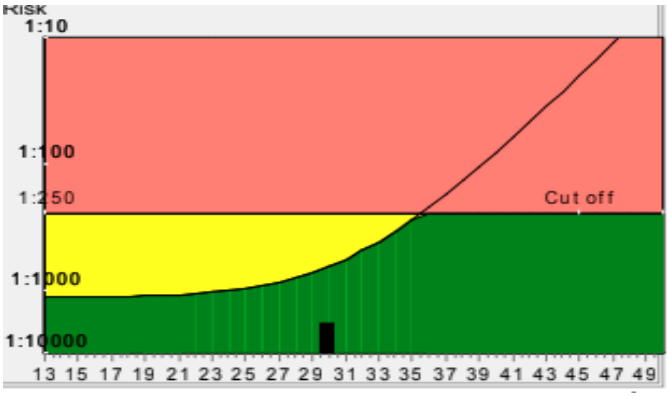
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	73.8	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data		
Parameter	**Value**	**Corr Mom**
PAPP-A	5.82 mIU/ml	1.15
fb-hCG	48.9 ng/ml	1.59

Risks at sampling date	
Age Risk	1:670
Biochemical T21 risk	1:1884
Combined trisomy 21 risk	1:9647
Trisomy 13/18 + NT	<1:10000

Ultrasound Data	
Gestational age	12+5
Method	CRL (<>Robinson)
Scan date	02-12-2023
Crown rump length in mm	63
Nuchal translucency MoM	0.80
Nasal bone	PRESENT
Sonographer	DR ABHISHEK
Qualifications in measuring NT	MBBS

Risk



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 9647 women with the same data, there is one woman with a trisomy 21 pregnancy and 9646 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