Date of Report	30/12/2022
PRISCA	5.1.0.17

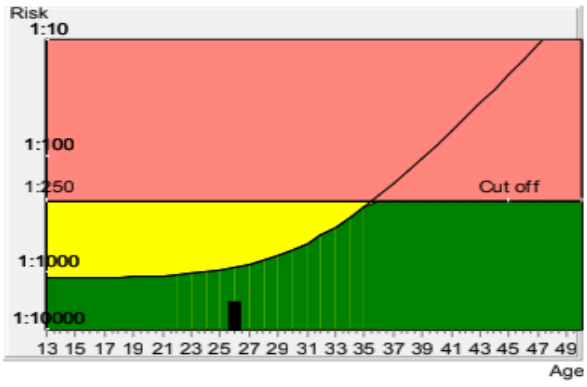
Patient Data			
Name	MRS. MANJULA RATHOR	Patient ID	012212290165
Birthday	18/01/1997	Sample ID	11551121
Age at term	26.03	Sample Date	29/12/2022
Gestational age	13+0		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	64	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom		12+6
PAPP-A	3.28 mIU/ml	0.55	Method	CRL (<>Robinson)
fb-hCG	24.6 ng/ml	0.77	Scan date	28/12/2022
			Nuchal translucency	1.2
			Nuchal translucency MoM	0.72
			Nasal bone	Present

Risks at sampling date	
Age Risk	1:933
Biochemical T21 risk	1:2452
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000

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 21with NT test it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 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 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