Date of Report	28/1/2023
PRISCA	5.1.0.17

Patient Data			
Name	MRS. MEENAKSHI	Patient ID	012301260046
Birthday	13/07/1993	Sample ID	11479693
Age at term	30	Sample Date	26/1/2023
Gestational age	12+6		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	62.9	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	3.72 mIU/ml	0.55
fb-hCG	38.2 ng/ml	0.98

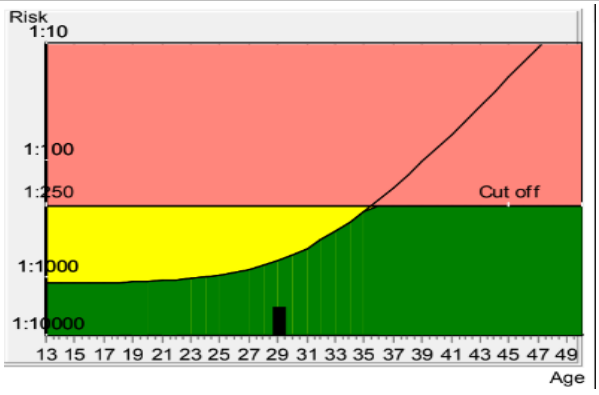
Ultrasound Data

	12+4
Method	CRL (<>Robinson)
Scan date	25/1/2023
Nuchal translucency	1.3
Nuchal translucency MoM	0.81
Nasal bone	Present

Risks at sampling date

Age Risk	1:727
Biochemical T21 risk	1:1156
Combined trisomy 21 risk	1:6655
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 6655 women with the same data, there is one woman with a trisomy 21 pregnancy and 6654 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