Date of Report	18/12/2024
PRISCA	5.2.0.13

Patient Data

Name	Mrs Maheshwari	Patient ID	012412160059
Birthday	16/01/1998	Sample ID	11996158
Age at delivery	26.9	Sample Date	16/01/1998
Gestational age	13+1		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	45	Diabetes	No	Pregnancies	unknown
Smoker	No	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	5.6 mIU/ml	0.59
fb-hCG	45.1 ng/ml	1.30

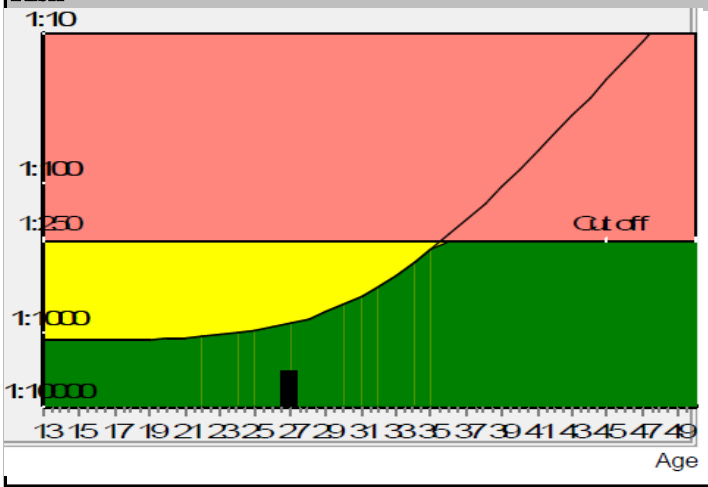
Ultrasound Data

Gestational age	12+5
Method	CRL(<>Robinson
Scan date	13/12/2024
Crown rump length in mm	63
Nuchal translucency MOM	1.23
Nasal bone	Present
Sonographer	Dr.
Qualification in measuring NT	MBBS

Risks at sampling date

Age Risk	1:883
Biochemical T21 risk	1:881
Combined Trisomy 21 risk	1:1839
Trisomy 13/18 + NT risk	<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 21 test it is expected that among 1839 women with the same data, there is one woman with a trisomy 21 pregnancy and 1838 women with no 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 the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-23, 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