Date of Report 14-01-2024
PRISCA 5.2.0.13

Patient Data			
Name	MRS YOJNA KUMARI	Patient ID	012401130098
Birthday	15-02-1996	Sample ID	11830799
Age at Sample date	27.9	Sample Date	13-01-2024
Gestational age	12+5		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	61.1	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	3.76 mIU/ml	0.66
fb-hCG	25.6 ng/ml	0.72

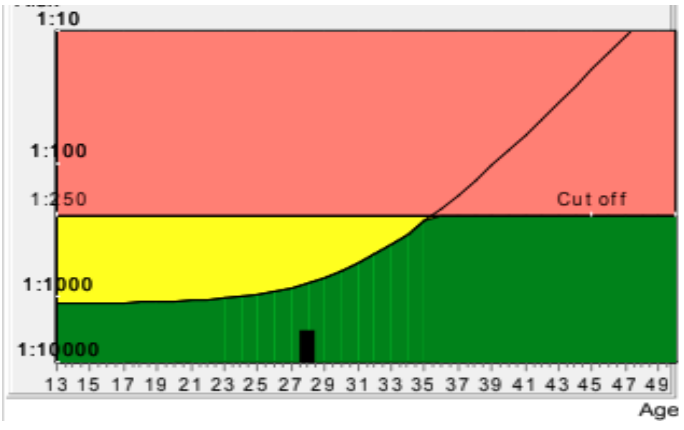
Risks at sampling date

Age Risk	1:807
Biochemical T21 risk	1:3833
Combined trisomy 21 risk	1:8160
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	12+4
Method	CRL (<>Robinson)
Scan date	12-01-2024
Crown rump length in mm	61.8
Nuchal translucency MoM	0.63
Nasal bone	ABSENT
Sonographer	DR JAG MOHAN
Qualifications in measuring NT	MD

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 8160 women with the same data, there is one woman with a trisomy 21 pregnancy and 8159 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