Date of Report	4/6/2024
PRISCA	5.2.0.13

Patient Data

Name	**POOJA MAMAIN**	Patient ID	012406030068
Birthday	16/8/1996	Sample ID	11876932
Age at sample	27.8	Sample Date	3/6/2024
Gestational age	13+0		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	65	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	6.10 mIU/ml	1.03
fb-hCG	67.4 ng/ml	2.11

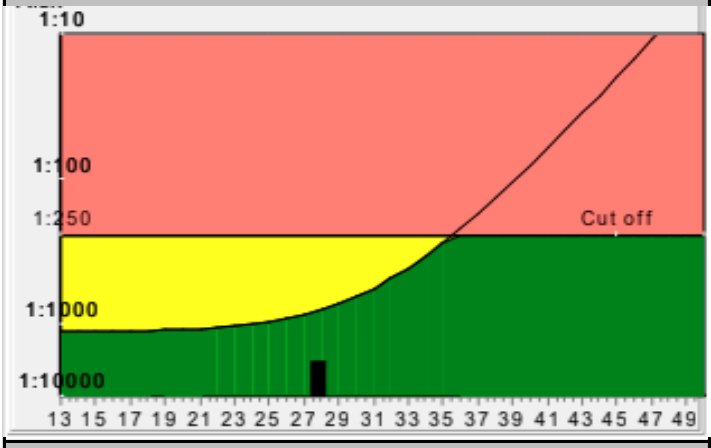
Risks at sampling date

Age Risk	1:822
Biochemical T21 risk	1:915
Combined trisomy 21 risk	1:5011
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	12+6
Method	CRL (<>Robinson)
Scan date	2/6/2024
Crown rump length in mm	62.2
Nuchal translucency MoM	0.72
Nasal bone	Present
Sonographer	DR.INDRAJEET
Qualifications in measuring NT	MD

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 (with NT) it is expected that among 5011 women with the same data, there is one woman with a trisomy 21 pregnancy and 5010 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