Date of Report 27-01-2025
PRISCA 5.2.0.13

Patient Data			
Name	MRS. SEJAL SHARMA	Patient ID	12501230198
Birthday	14-08-1998	Sample ID	11484389
Age at Sample date	26.5	Sample Date	23-01-2025
Gestational age	13+0		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	52	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	7.6 mIU/ml	0.99
fb-hCG	51.4 ng/ml	1.49

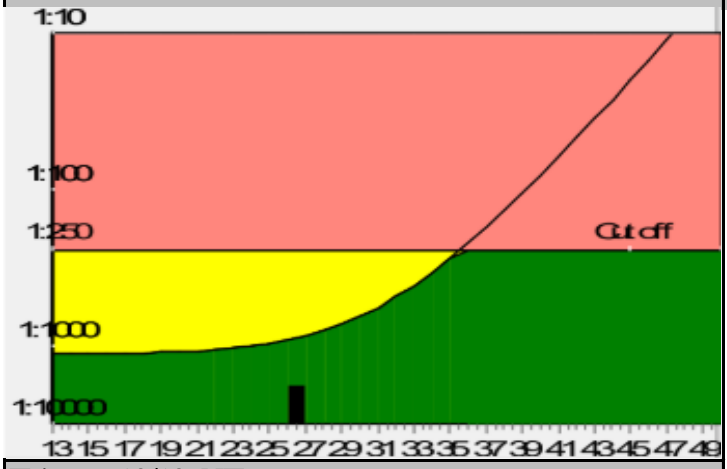
Risks at sampling date

Age Risk	1:906
Biochemical T21 risk	1:2173
Combined trisomy 21 risk	1:6401
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	12+5
Method	CRL (<>Robinson)
Scan date	23-01-2025
Crown rump length in mm	69
Nuchal translucency MoM	1.11
Nasal bone	PRESENT
Sonographer	DR.SHRUTI SANGWAN
Qualifications in measuring NT	MBBS

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 6401 women with the same data, there is one woman with a trisomy 21 pregnancy and 6400 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