

Date of Report 24-01-2025
PRISCA 5.2.0.13

Patient Data			
Name	MRS. PRITI KUMARI	Patient ID	12501230086
Birthday	06-05-1995	Sample ID	11971618
Age at Sample date	29.7	Sample Date	23-01-2025
Gestational age	13+2		

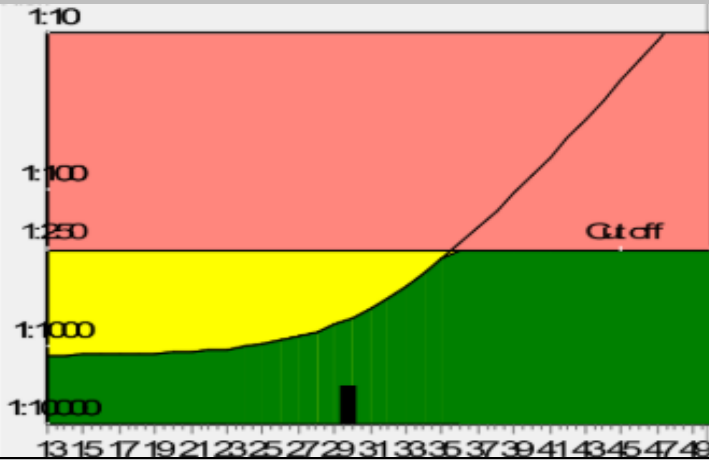
Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	40	Diabetes	NO
Smoker	NO	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data			Ultrasound Data	
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Parameter	Value	Corr Mom		
PAPP-A	6.9 mIU/ml	0.61	Gestational age	13+2
fb-hCG	53.4 ng/ml	1.53	Method	CRL (<>Robinson)
			Scan date	23-01-2025

Risks at sampling date			Ultrasound Data	
Age Risk		1:686	Crown rump length in mm	71.1
Biochemical T21 risk		1:496	Nuchal translucency MoM	0.68
Combined trisomy 21 risk		1:2988	Nasal bone	PRESENT
Trisomy 13/18 + NT		<1:10000	Sonographer	DR.
			Qualifications in measuring NT	MBBS

Risk	Down's Syndrome Risk (Trisomy 21 Screening)
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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 2988 women with the same data, there is one woman with a trisomy 21 pregnancy and 2987 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 approaches 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).

Trisomy 13/18+NT
The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk

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