

Date of Report 19/08/2024
PRISCA 5.0.2.13

Patient Data			
Name	Mrs. Kamini Tyagi	Patient ID	012408160109
Birthday	01/02/1995	Sample ID	11863762
Age at delivery	29.5	Sample Date	16/08/2024
Gestational age	12+5		

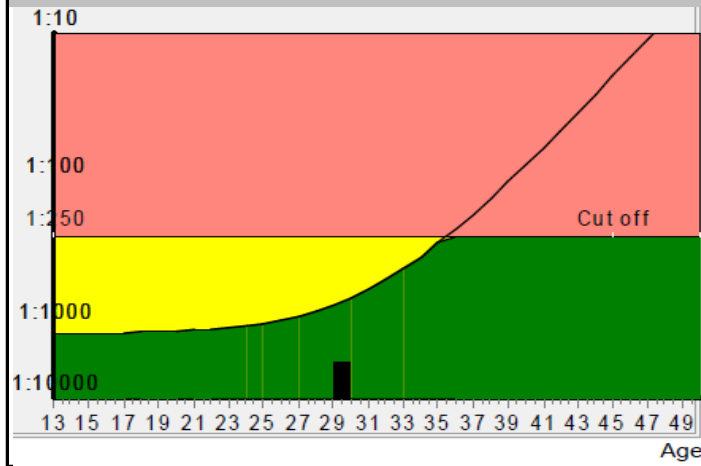
Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	56	Diabetes	No
Smoker	Unknown	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	

Biochemical Data			Ultrasound Data	
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Parameter	Value	Corr Mom		
PAPP-A	5.6 mIU/ml	0.88	Gestational age	12+4
fb-hCG	23.5 ng/ml	0.65	Method	CRL(<>Robinson)
			Scan date	15/08/2024

Risks at sampling date			Ultrasound Data	
Age Risk	1:687		Crown rump length in mm	62.1
Biochemical T21 risk	1:8351		Nuchal translucency MOM	0.72
Combined Trisomy 21 Risk	<1:10000		Nasal bone	Present
Trisomy 13/18 + NT	<1:10000		Sonographer	Dr Inderjeet
			Qualification in measuring NT	MBBS, MD Radio-diagnosis

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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.

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 the 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

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

Risk Above Cut Off
 Risk below Cut Off but above Age Risk
 Risk below Cut Off