

Biochemical Data

*Free Home Sample Collection 9999 778 778



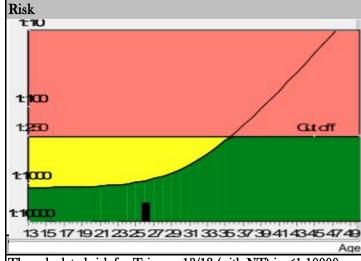
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Date of Report 22/02/2025 PRISCA 5.2.0.13

					TRISCA	J.Z.U.10
Patient Data						
Name			SWATI	Patient ID		12502200217
Birthday			18/1/1999	Sample ID		11925394
Age at Sample date			26.1'	Sample Date		20/02/2025
Gestational age			13+3			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	74.1	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		

Ultrasound Data

Parameter Value		Corr Mom	Gestational age	11+2	
PAPP-A	$7.2~\mathrm{mIU/ml}$	1.24	Method	CRL (<>Robinson)	
fb-hCG	23.5 ng/ml	0.87	Scan date	5/2/2025	
Risks at sampling date			Crown rump length in mm	45	
Age Risk		1:938	Nuchal translucency MoM	0.64	
Biochemical T21 risk		<1:10000	Nasal bone	PRESENT	
Combined trisomy 21 risk		<1:10000	Sonographer	DR. JAG MOHAN	
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MD	



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 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 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