

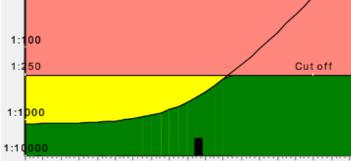
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Date of Report 23-09-2024

				PRISCA	5.2.0.13
Patient Data				TMSC/1	0.2.0.10
Name	MRS. SHIVANI SINGLA		Patient ID		12409220203
Birthday	11-07-1992		Sample ID		11863749
Age at Sample date	32.2		Sample Date		22-09-2024
Gestational age	13+2				
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	70 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational ag	e	13+1
PAPP-A	7.6 mIU/ml	1.28	Method		CRL (<>Robinson)
fb-hCG	66.29 ng/ml	2.32	Scan date		21-09-2024
Risks at sampling date			Crown rump length in mm 69.7		
Age Risk 1:481		Nuchal translucency MoM 0.81			
Biochemical T21 risk 1:642		Nasal bone PRESEN		PRESENT	
Combined trisomy 21 risk 1:3345		Sonographer Dr. INDRAJEF		DR. INDRAJEET	
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	MD
Risk 1:100 1:450 Cut off			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 3345 women with the same data, there is one woman with a trisomy 21 pregnancy and 3344 women with not affected pregnancies. The calculated risk by PRISCA depends on the accuracy of the		



13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 1998). Trisomy 13/18+NT

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

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;

The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values