

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

which indicates a low risk

*Free Home Sample Collection 9999 778 778



risk assessment! Calculated risks have no diagnostic values

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Date of Report 20-05-2024 PRISCA 5.2.0.13

			PRISCA	5.2.0.13
Patient Data				
Name	MRS. NI	ΓU SHARMA	Patient ID	12405190186
Birthday		22-08-1997	Sample ID	12001233
Age at Sample date		26.7	Sample Date	19-05-2024
Gestational age		13+6		
Correction factors				
Fetuses	1 IVF		unknown Previous trison	ny 21 unknown
Weight in kg	65 Diabetes		NO Pregnancies	unknown
Smoker	NO Origin		Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+6
PAPP-A	6.8 mIU/ml	0.87	Method	CRL (<>Robinson)
fb-hCG	52.8 ng/ml	2.19	Scan date	19-05-2024
Risks at sampling date			Crown rump length in mm	80
Age Risk		1:914	Nuchal translucency MoM	0.73
Biochemical T21 risk		1:632	Nasal bone	PRESENT
Combined trisomy 21 risk	ζ.	1:3640	Sonographer	DR.DEEPIKA
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring N	Т мр
Risk			Down's Syndrome Risk (Trisc	omy 21 Screening)
1:100 1:250 Cut off 1:1000			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 3640 women with the same data, there is one woman with a trisomy 21 pregnancy and 3639 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	
13 15 17 19 21 23 25 27	7 29 31 33 35 37 39	41 43 45 47 49	done according to accepted guidelines (Prenat Diagn 18:511-523; 1998).	
Trisomy 13/18+NT The calculated risk for Trise	omer 19/10 (with NT)	:- <1.10000	The laboratory cannot be hold re-	sponsible for their impact on the