

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 01-09-2024 PRISCA 5.2.0.13

				PRISCA	5.2.0.13
Patient Data					
Name	MRS. JYOT	TI AWASTHI	Patient ID		12407310276
Birthday		12-08-1989	Sample ID		11905716
Age at Sample date		35	Sample Date		31-07-2024
Gestational age		13+1			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	73 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Da	ata.	
Parameter	Value	Corr Mom	Gestational age	2	13+1
PAPP-A	6.8 mIU/ml	1.26	Method		CRL (<>Robinson)
fb-hCG	27.5 ng/ml	0.93	Scan date		31-07-2024
Risks at sampling date			Crown rump length in mm 69.3		
Age Risk		1:280	Nuchal translu	cency MoM	0.61
Biochemical T21 risk		1:3310	Nasal bone		present
Combined trisomy 21 risk	X.	<1:10000	Sonographer		DR. INDRAJEET
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	MD
Risk 1:10			Down's Syndro	ome Risk (Trisomy 21	Screening)
1:100 1:250 Cut off			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!		
1:10000 13 15 17 19 21 23 25 2 Trisomy 13/18+NT The calculated risk for Trise			done according 1998).	abined risk presumes that it to accepted guidelines (Precannot be hold responsible	enat Diagn 18:511-523;