

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

Book a Test Online www.molq.in

Date of Report 20-06-2024 PRISCA 5.2.0.13

				PRISCA	5.2.0.13
Patient Data					
Name MRS. CHANCHAL W/O VICKY			Patient ID		12406190172
Birthday 31-03-2006			Sample ID		11893027
Age at Sample date 18.2			2 Sample Date		19-06-2024
Gestational age		11+3	3		
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	44.1 Diabetes	S	NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	e	11+3
PAPP-A	3.1 mIU/ml	0.63	Method		CRL (<>Robinson)
fb-hCG	67.6 ng/ml	1.24	Scan date		19-06-2024
Risks at sampling date			Crown rump length in mm 46.6		
Age Risk	age Risk 1:1063		Nuchal translucency MoM 1.4		
Biochemical T21 risk		1:1405	Nasal bone		PRESENT
Combined trisomy 21 risk 1:1471			Sonographer DR.VIKAS		DR.VIKASH
Trisomy 13/18 + NT <1:10000			Qualifications in measuring NT MD		
Risk			Down's Syndr	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 1471 women with the same data, there is one woman with a trisomy 21 pregnancy and 1470 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		
1:1000 1:1000 13 15 17 19 21 23 25 Trisomy 13/18+NT The calculated risk for T	27 29 31 33 35 37 39		the risk calculated diagnostic value. The patient condone according 1998).	ions are statistical aapproa	ches and have no NT measurement was renat Diagn 18:511-523;