

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

which indicates a low risk

\*Free Home Sample Collection 9999 778 778



Book a Test Online www.molq.in

Date of Report 16-08-2024 PRISCA 5.2.0.13

The laboratory cannot be hold responsible for their impact on the

risk assessment! Calculated risks have no diagnostic values

					PRISCA	5.2.0.13
Patient Data						
Name			MRS. NEHA	Patient ID		12408140077
Birthday			04-08-1999	Sample ID		11883590
Age at Sample date			25	Sample Date		14-08-2024
Gestational age			13+3			
Correction factors					T	
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	54.5	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational ag	e	12+6
PAPP-A	6.8	mIU/ml	0.81	Method		CRL (<>Robinson)
fb-hCG	23.6	ng/ml	0.8	Scan date		10-08-2024
Risks at sampling date				Crown rump l	ength in mm	65.8
Age Risk			1:989	Nuchal translu	icency MoM	0.66
Biochemical T21 risk			1:6306	Nasal bone		PRESENT
Combined trisomy 21 risk			<1:10000	Sonographer		DR. RAHUL
Trisomy 13/18 + NT			<1:10000	Qualifications	in measuring NT	MBBS
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 Cut off 1:1000 1:1000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49				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).		
Trisomy 13/18+NT				1330).		