

Trisomy 13/18+NT

which indicates a low risk

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

\*Free Home Sample Collection 9999 778 778



Book a Test Online www.molq.in

Date of Report 18-09-2024 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.	ΓANIMA	SRIVATAVA	Patient ID		12409170182
Birthday			10-11-1991	Sample ID		11858744
Age at Sample date			32.9	Sample Date		17-09-2024
Gestational age			12+0			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 2	21 unknown
Weight in kg	86	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data			Ultrasound Data			
Parameter	Value		Corr Mom	Gestational age	e	12+0
PAPP-A	4.3	mIU/ml	1.52	Method		CRL (<>Robinson)
fb-hCG	49.8	ng/ml	1.29	Scan date		17-09-2024
Risks at sampling date				Crown rump l	ength in mm	52
Age Risk			1:409	Nuchal translu	icency MoM	0.86
Biochemical T21 risk			1:3216	Nasal bone		PRESENT
Combined trisomy 21 ris	sk		<b>&lt;</b> 1:10000	Sonographer		DR. PRAKASH LALCHANDAN
Trisomy 13/18 + NT			<b>-</b> <1:10000	Qualifications	in measuring NT	MD
				Down's Syndrome 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		
1:1(000				diagnostic value!  The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998).		