

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 25-06-2024 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.LAXMI	Patient ID		12406240163
Birthday			13-01-1999	Sample ID		11860942
Age at Sample date			25.4	Sample Date		24-06-2024
Gestational age 13+5						
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	62	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational ag	e	13+5
PAPP-A	7.1	mIU/ml	0.89	Method		CRL (<>Robinson)
fb-hCG	15.6	ng/ml	0.61	Scan date		24-06-2024
Risks at sampling date				Crown rump l	ength in mm	77.5
Age Risk			1:979	Nuchal translu	icency MoM	0.90
Biochemical T21 risk			<1:10000	Nasal bone		PRESENT
Combined trisomy 21 ris	k		<1:10000	Sonographer		DR. Deepika
Trisomy 13/18 + NT			<1:10000	Qualifications	in measuring NT	MD
Risk				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		
1:1000 1:10000 13 15 17 19 21 23 25 2 Trisomy 13/18+NT	27 29 31 33	3 35 37 39	41 43 45 47 49	the risk calculate diagnostic value The patient con done according	vided by the referring phy ions are statistical aapproa! hbined risk presumes that to accepted guidelines (Pr	ches and have no NT measurement was