

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 02-12-2024 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. VISHAKI	IA SHARMA	Patient ID		12412010077
Birthday		11-08-1995	Sample ID		11996245
Age at Sample date		29.3	Sample Date		01-12-2024
Gestational age		12+3			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	43.5 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Da	ata	
Parameter	Value	Corr Mom	Gestational ago	e	12+3
PAPP-A	4.9 mIU/ml	0.65	Method		CRL (<>Robinson)
fb-hCG	65.4 ng/ml	1.52	Scan date		01-12-2024
Risks at sampling date			Crown rump l	ength in mm	60.7
Age Risk		1:698	Nuchal translu	icency MoM	0.63
Biochemical T21 risk		1:604	Nasal bone		PRESENT
Combined trisomy 21 risk		1:3571	Sonographer		DR. DEEPIKA
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 1:1000 1:10000 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 3571 women with the same data, there is one woman with a trisomy 21 pregnancy and 3570 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 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).	. 1 1 11 21	