

\*Free Home Sample Collection 9999 778 778



Book a Test Online www.molq.in

 Date of Report
 13-03-2024

 PRISCA
 5.2.0.13

					TRISCIT	5.2.0.10
Patient Data						
Name	MRS. POONAM			Patient ID		12403120111
Birthday	19-10-1991			Sample ID		11792242
Age at Sample date	32.4			Sample Date		12-03-2024
Gestational age	estational age 13+1					
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	52	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	e	12+1
PAPP-A	5.9	mIU/ml	0.73	Method		CRL (<>Robinson)
fb-hCG	25.6 ng/ml		0.78	Scan date		05-03-2024
Risks at sampling date				Crown rump length in mm 555		
Age Risk			1:463	Nuchal translu	icency MoM	0.96
Biochemical T21 risk			1:2469	Nasal bone		PRESENT
Combined trisomy 21 risk			<1:10000	Sonographer		DR. PALLAVI GUPTA
Trisomy 13/18 + NT			<1:10000	Qualifications	in measuring NT	MBBS
Risk 1:10				Down's Syndrome Risk (Trisomy 21 Screening)		
1:100  1:1000  1:10000  1:10000  13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49  Trisomy 13/18+NT  The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk				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 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).  The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		
which indicates a low risk				<u> </u>		