

which indicates a low risk

\*Free Home Sample Collection 9999 778 778



Book a Test Online www.molq.in

Date of Report 27-08-2024 PRISCA 5.2.0.13

				PRISCA	5.2.0.13
Patient Data					
Name	MRS. DEEPALI		Patient ID		12408260129
Birthday	10-03-1998		Sample ID		11863984
Age at Sample date		26.5	Sample Date		26-08-2024
Gestational age		12+4	•		
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	90 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian	1	
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational ag	ge	12+4
PAPP-A	5.3 mIU/ml	1.58	Method		CRL (<>Robinson)
fb-hCG	22.4 ng/ml	0.68	Scan date		26-08-2024
Risks at sampling date			Crown rump length in mm 59.7		
Age Risk	ge Risk 1:892		Nuchal translucency MoM 0.		0.77
Biochemical T21 risk <1:10000		Nasal bone		PRESENT	
Combined trisomy 21 risk		<1:10000	Sonographer		DR. SANJEEV KUMAR
Trisomy 13/18 + NT		<1:10000	Qualifications	s 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 information provided by the referring physician. Please note that the risk calculations are statistical aapproaches and have no		
1:10000 13 15 17 19 21 23 25 27 Trisomy 13/18+NT	41 43 45 47 49	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			
The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk			risk assessment! Calculated risks have no diagnostic values		