

*Free Home Sample Collection 9999 778 778

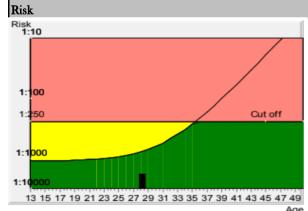


Book a Test Online www.molq.in

Date of Report 24/3/2022 PRISCA 51017

					PRISCA	5.1.0.17
Patient Data						
Name		MI	RS. DEEPA	Patient ID		012203230176
Birthday			1/1/1994	Sample ID		11433525
Age at term			28.08	Sample Date		23/3/2022
Gestational age			12+0			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	81	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data			Ultrasound Da	ata		

biochemicai Data			Ultrasound Data		
Parameter Value		Corr Mom	Gestational age	12+0	
PAPP-A	3.52 mIU/ml	0.87	Method	CRL (<>Robinson)	
fb-hCG	53.4 ng/ml	0.94	Scan date	23/3/2022	
Risks at sampling date			Nuchal translucency	1	
Age Risk		1:765	Nuchal translucency MoM	0.78	
Biochemical T21 risk		1:4050	Nasal Bone	Presesnt	
Combined trisomy 21 risk		<1:10000			
Trisomy 13/18+NT		<1:10000			



Down's Syndrome Risk (Trisomy 21 Screening)

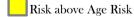
The calculated risk for Trisomy 21 with NT is below the cut off, which represents a low risk.

After the result of the Trisomy 21 test it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy 9999 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!

Trisomy 13/18+NT The calculated risk for Trisomy 13/18 with NT is <1:10000, which indicates a low risk

The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values







Risk below Age risk