

\*Free Home Sample Collection 9999 778 778



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Date of Report 05-02-2025
PRISCA 5 2 0 13

					PRISCA	5.2.0.13	
Patient Data							
Name	MRS. MADHURI			Patient ID		12502040151	
Birthday	nday 17-09-2000					11932351	
Age at Sample date 24.4				Sample Date		04-02-2025	
Gestational age			12+2	2			
Correction factors		1					
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	56	Diabetes		NO	Pregnancies	unknown	
Smoker	NO	Origin		Asian			
Biochemical Data				Ultrasound Data			
Parameter	Value		Corr Mom	Gestational ag	e	12+2	
PAPP-A	4.5	mIU/ml	0.84	Method		CRL (<>Robinson)	
fb-hCG	41.8	ng/ml	1.03	Scan date		04-02-2025	
Risks at sampling date				Crown rump length in mm 57.2			
Age Risk	1:978			Nuchal translucency MoM 0.61			
Biochemical T21 risk			1:3885	Nasal bone		PRESENT	
Combined trisomy 21 risk			<1:10000	Sonographer		DR.VARUN	
Trisomy 13/18 + NT			<1:10000	Qualifications	in measuring NT		
Risk 1:10				Down's Syndrome Risk (Trisomy 21 Screening)			
1:100 1:20 Gt off 1:100				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 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,				-	risk assessment! Calculated risks have no diagnostic values		

which indicates a low risk