

*Free Home Sample Collection 9999 778 778



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Date of Report 22-10-2024 PRISCA 5.2.0.13

					PRISCA	5.2.0.13
Patient Data						
Name	MRS. NEELAM			Patient ID		12410210101
Birthday	15-03-1984			Sample ID		11996713
Age at Sample date 40.6			Sample Date		21-10-2024	
Gestational age			11+6			
Correction factors		ı				
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	58.3 Diabetes			NO	Pregnancies	unknown
Smoker	NO Origin			Asian	ı	
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational ag	e	11+6
PAPP-A	5.7	mIU/ml	1.33	Method		CRL (<>Robinson)
fb-hCG	31.2	ng/ml	0.7	Scan date		21-10-2024
Risks at sampling date				Crown rump length in mm 51.1		
Age Risk			1:65	Nuchal translu	icency MoM	0.87
Biochemical T21 risk			1:1585	Nasal bone		present
Combined trisomy 21 risk			1:6924	Sonographer		DR. SUMIT
Trisomy 13/18 + NT			<1:10000	Qualifications	in measuring NT	MD
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
1:10 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 Trisomy 13/18+NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000,				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 6924 women with the same data, there is one woman with a trisomy 21 pregnancy and 6923 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). The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		
which indicates a low risk						-