

\*Free Home Sample Collection **9999 778 778**  Download "MOLQ" App on

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				Date of Report	29-10-2024
				PRISCA	5.2.0.13
Patient Data Name	MI	RS. SHWETA	Patient ID		12410270113
Birthday			Sample ID		11996491
Age at Sample date			Sample Date		27-10-2024
Gestational age		12+2	-		
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	60 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	e	12+1
PAPP-A	5.3 mIU/ml	1.07	Method		CRL (<>Robinson)
fb-hCG	25.6 ng/ml	0.64	Scan date		26-10-2024
Risks at sampling date			Crown rump length in mm 58.4		
Age Risk		1:114	Nuchal translu	icency MoM	0.72
Biochemical T21 risk	iochemical T21 risk 1:2132		Nasal bone PRESENT		
Combined trisomy 21 risk		1:9801	Sonographer		DR. DEEPIKA
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	MD
Risk 1:10			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 Cut off 1:100 1:1000			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 9801 women with the same data, there is one woman with a trisomy 21 pregnancy and 9800 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		
which indicates a low risk			risk assessment!	Calculated risks have no o	diagnostic values



Risk Above Cut Off

