

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

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				Date of Report	14-12-2024
				PRISCA	5.2.0.13
Patient Data Name		MRS. NEHA	Patient ID		12412130090
Birthday			Sample ID		11860038
Age at Sample date			Sample Date		13-12-2024
Gestational age		12+2	-		
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	78.9 Diabete	es	NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+2
PAPP-A	5.7 mIU/m	1 1.60	Method		CRL (<>Robinson)
fb-hCG	23.1 ng/ml	0.63	Scan date		13-12-2024
Risks at sampling date			Crown rump length in mm 59		
Age Risk		1:305	Nuchal translu	cency MoM	1.10
Biochemical T21 risk	ochemical T21 risk <1:10000		Nasal bone present		
Combined trisomy 21 risk		<1:10000	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 Cutoff 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 and 1472 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 calculated risk presence of the responsible for their impact on the risk calculations are statistical ways and the responsible for the responsed of the respon		
which indicates a low risk					



Risk Above Cut Off

