

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

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					Date of Report PRISCA	11-07-2024 5.2.0.13
Patient Data						
Name	MRS. NEETU			Patient ID		12407100124
Birthday	23-04-1988			Sample ID		11858929
Age at Sample date	36.2			Sample Date		10-07-2024
Gestational age			12+3			
Correction factors					Γ	
Fetuses	1	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	72 Diabetes		NO	Pregnancies	unknown	
Smoker	NO Origin		Asian			
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	2	12+3
PAPP-A	4.5	mIU/ml	1.07	Method		CRL (<>Robinson)
fb-hCG	22.3 ng/ml 0.61		0.61	Scan date		10-07-2024
Risks at sampling date				Crown rump length in mm 59		
Age Risk			1:206	Nuchal translu	cency MoM	0.84
Biochemical T21 risk			1:4209	Nasal bone		PRESENT
Combined trisomy 21 risk			<1:10000	Sonographer		DR. Prakash Lalchandan
Trisomy 13/18 + NT			<1:10000	Qualifications	in measuring NT	MD
Risk 1:10				Down's Syndro	ome Risk (Trisomy 21	Screening)
1:100 1:250 1:1000 1:1000 1:1000 13 15 17 19 21 23 25 27 Trisomy 13/18+NT The calculated risk for Triso which indicates a low risk			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 amog 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 risk assessment! Calculated risks have no diagnostic values			



Risk Above Cut Off

