

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

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				Date of Report	12-08-2024
Patient Data				PRISCA	5.2.0.13
Name					12408110140
Birthday	-		Sample ID		11883707
Age at Sample date			Sample Date		11-08-2024
Gestational age 12+3			-		
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	48.5 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian	-	
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational ag	e	12+3
PAPP-A	5.7 mIU/ml	0.85	Method		CRL (<>Robinson)
fb-hCG	49.8 ng/ml	1.2	Scan date		11-08-2024
Risks at sampling date			Crown rump length in mm 58.8		
Age Risk 1:824			Nuchal translucency MoM 1.24		
Biochemical T21 risk	1:2344		Nasal bone PR		PRESENT
Combined trisomy 21 risk 1:4654		Sonographer DR. VIKASH			
Frisomy 13/18 + NT <1:10000		Qualifications in measuring NT M			
Risk				ome Risk (Trisomy 21	Screening)
1:100 1:250 Cut off 1:100 1:1000 1:1000 1:10000 1:10000 1:15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Trisomy 13/18+NT			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 4654 women with the same data, there is one woman with a trisomy 21 pregnancy and 4653 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	,	115K assessinent!	Calculated LISKS Have HO	ulaghosut values	



Risk Above Cut Off

