

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

Book a Test Online www.molq.in

				Date of Report PRISCA	1/2/2023 5.1.0.17
Patient Data					
Name		MRS. NISHA	Patient ID		012301310099
Birthday		1/9/1994	Sample ID		11408272
Age at term		28.1	Sample Date		31/1/2023
Gestational age		12+1			
Correction factors				_	
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	53 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			12+0
PAPP-A	2.89 mIU/ml	0.53	Method		CRL (<>Robinson)
fb-hCG	34.5 ng/ml	0.8	Scan date		30/1/2023
Risks at sampling date			Nuchal translucency 1.2		
Age Risk	1:756		Nuchal translucency MoM 0.8		
Biochemical T21 risk		1:1712	Nasal bone		Present
Combined trisomy 21 risk	i.	1:9718			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndr	ome Risk (Trisomy 21	Screening)
Risk 1:10 1:00 1:250 Cut off 1:100 1:100 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 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 with NT test it is expected that among 9718 women with the same data, there is one woman with a trisomy 21 pregnancy and 9717 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 laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic		
<1:10000 , which indicate	es a low risk		values		
<b>Risk</b> A	Above Cut Off		Risk above Ag	e Kisk	Risk below Age risk