

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

Book a Test Online www.molq.in

				Date of Report PRISCA	20/5/2022 5.1.0.17
Patient Data					
Name	MRS. SURE	BHI SHARMA	Patient ID		012205210090
Birthday		29/4/1994	Sample ID		11470719
Age at term		28.07	Sample Date		19/5/2022
Gestational age		12+0			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	70.3 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational ag	2	12+0
PAPP-A	3.1 mIU/ml	1.02	Method		CRL (<>Robinson)
fb-hCG	70.4 ng/ml	1.49	Scan date		21/5/2022
Risks at sampling date			Nuchal Translucency 0.9		
Age Risk	1:776		Nuchal Translucency MoM 0.6		
Biochemical T21 risk	1:1985		Nasal Bone	Nasal Bone Presen	
Combined T21 risk		<1:10000			
Trisomy 13/18+NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:250 Cut off 1:250 Cut off 1:1000 1:			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 9999 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 values		
Risk	Above Cut Off		Risk above Ag	e Risk	Risk below Age risk