

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

Book a Test Online www.molq.in

				Date of Report PRISCA	21/3/2023 5.2.0.13
Patient Data					
Name	MRS. PA	LLAVI GARG	Patient ID		012303200142
Birthday		8/12/1990	Sample ID		11523937
Age at term		32.9	Sample Date		20/3/2023
Gestational age		12+1			
Correction factors				Γ	
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	38.5 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			12+1
PAPP-A	4.28 mIU/ml	0.55	Method		CRL (<>Robinson)
fb-hCG	52.9 ng/ml	1.08	Scan date		20/3/2023
Risks at sampling date			Nuchal translucency 1.1		
Age Risk 1:456		1:456	Nuchal translucency MoM 0.7		
Biochemical T21 risk	1:578		Nasal bone Preser		
Combined trisomy 21 risk		1:3397			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndr	ome Risk (Trisomy 21	Screening)
Risk 1:10 1:250 0.ut off 1:1000 1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age			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 3397 women with the same data, there is one woman with a trisomy 21 pregnancy and 3396 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		
The calculated risk for Trisomy 13/18 (with NT) is <1:10000 , which indicates a low risk			on the risk assessment! Calculated risks have no diagnostic values		
Risk A	above Cut Off		Risk above Ag	e Risk	Risk below Age risk