

*Free Home				
Sample Collection 9999 778 778				
3333 110 110				

Download "MOLQ" App on

Book a Test Online www.molq.in

				Date of Report PRISCA	20/2/2022 5.2.0.13	
Patient Data						
Name	М	RS. ANTIMA	Patient ID		012202190019	
Birthday		9/11/1996	Sample ID		11307300	
Age at delivery		25.3	Sample Date		19/02.2022	
Gestational age		11+5				
Correction factors	1			-		
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown	
Weight in kg	43 Diabetes	43 Diabetes		Pregnancies	unknown	
Smoker	NO Origin	NO Origin				
Biochemical Data			Ultrasound Data			
Parameter	Value	Corr Mom	Gestational age	2	11+4	
PAPP-A	3.1 mIU/ml	0.65	Method		CRL (<>Robinson)	
fb-hCG	26.7 ng/ml	0.46	Scan date		18/02/2022	
Risks at sampling date			Nuchal translucency 1.5			
Age Risk		1:921	Nuchal translu	cency MoM	1.17	
Biochemical T21 risk		<1:10000	Nasal Bone		Present	
Combined T21 risk		<1:10000				
Trisomy 13/18+NT		<1:10000				
Risk			Down's Syndrome Risk (Trisomy 21 Screening)			
Risk 1:100 1:250 1:1000 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:1000 1:1000 1:1000 1:1000 1:250 1:1000 1:250 1:1000 1:250 1:1000 1:250 1:1000 1:250 1:10000 1:100000 1:10000 1:10000 1:10000 1:10000 1:10000 1:1000	Frisomy 13/18 with	1 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 test 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 Age Risk Risk below Age risk						