

*Free Home Sample Collection				
9999 778 778				

Download "MOLQ" App on

Book a Test Online www.molq.in

				Date of Report PRISCA	8/9/2021 5.2.0.13
Patient Data					
Name	MRS. SE	EMA YADAV	Patient ID		012109080146
Birthday		15/07/1989	Sample ID		11030577
Age at delivery		32.7	Sample Date		8/9/2021
Gestational age		11+4			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	62 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	11+4
PAPP-A	1.04 mIU/ml	0.35	Method		CRL(<>Robinson
fb-hCG	47.6 ng/ml	0.97	Scan date		8/9/2021
Risks at sampling date			Nuchal translucency in mm 1		
Age Risk		1:661	Nuchal translu	cency MOM	0.74
Biochemical T21 risk		1:304	Nasal bone		Present
Combined Trisomy 21 Risl	ζ.	1:1990			
Trisomy 13/18 + NT		<1:10000	Qualification i	n measuring NT	CR
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10			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 1990 women with the		
1:100 1:250 Cut off 1:1000 1:1000 1:10000 1:3 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age			same data, there is one woman with a trisomy 21 pregnancy and 1989 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!		
Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk			The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		
Risk A	bove Cut Off		Risk above Ag	e Risk 📃 Ri	isk below Age risk