

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

Book a Test Online www.molq.in

				Date of Report PRISCA	20/6/2022 5.1.0.17
Patient Data					
Name		MRS. SEEMA	Patient ID		012206190092
Birthday		20/05/1990	Sample ID		11461464
Age at term		31.07	Sample Date		19/6/2022
Gestational age		12+5			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	63 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+4
PAPP-A	5.24 mIU/ml	0.95	Method		CRL(<>Robinson)
fb-hCG	38.7 ng/ml	1.11	Scan date		18/6/2022
Risks at sampling date			Nuchal Translucency 1.3		
Age Risk	1:481		Nuchal Translucency MoM 0.8		
Biochemical T21 risk		1:2132	Nasal Bone		Present
Combined T21 risk		<1:10000			
Trisomy 13/18+NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:100 1:250 1:1000 1:1000 1:10000 1:10000 1:3 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 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!		
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	x Above Cut Off		Risk above Ag	e Risk	lisk below Age risk