

*Free Home Sample Collection 9999 778 778 Download "MOLQ" App on Book a Test Online www.molq.in

				Date of Report PRISCA	28/6/2023 5.2.0.13
Patient Data					
Name	MR	S. POONAM	Patient ID		012306270191
Birthday		15/10/1991	Sample ID		11664072
Age at term		32.3	Sample Date		27/6/2023
Gestational age		12+6			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	56 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			12+0
PAPP-A	6.24 mIU/ml	0.93	Method		CRL (<>Robinson)
fb-hCG	122.4 ng/ml	3.51	Scan date		23/6/2023
Risks at sampling date			Crown rump length in mm 54.4		
Age Risk		1:515	Nuchal translucency MoM		0.83
Biochemical T21 risk		1:121	Nasal bone		Present
Combined trisomy 21 risk		1:696			
Trisomy 13/18 + NT		<1:10000			
Risk 1:10			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 Cut off 1:1000 1:10000 1:10000 1:10000 1:10000 The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk			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 696 women with the same data, there is one woman with a trisomy 21 pregnancy and 695 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 🛛 🚺 R	lisk below Age risk