

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

Book a Test Online www.molq.in

				Date of Report PRISCA	18-06-19 5.0.2.37
Patient Data					
Name		Mrs. PREETI	Patient ID		011906160087
Birthday		28-07-92	Sample ID		10468801
Age at delivery		26.9	Sample Date		16/6/2019
Gestational age		12+6			
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	12+6
PAPP-A	3.04 mIU/ml	0.62	Method		CRL (<>Robinson)
fb-hCG	36.3 ng/ml	0.82	Scan Date		16-06-19
Risks at sampling date			Crown rump length in mm 58.2		
Age Risk		1:876	Nuchal translucency MoM		0.92
Biochemical T21 risk		1:2830	Nasal Bone		Present
Combined Trisomy 21 F	Risk	<1:10000	sonographer		DR.SANJEEV KUMAR
Trisomy 13/18 + NT		<1:10000	Qualification in	n measuring NT	MBBS,PGDUS,DMRD
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		
1:100 1:250 Out off 1:1000 1:1000 1:1000 1:1000 1:1000 Age			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		Risk above Ag	e Risk	Risk below Age risk	