

\*Free Home Sample Collection 9999 778 778

ć

Download "MOLQ" App on 

Book a Test Online www.molq.in

				Date of Report PRISCA	03-01-19 5.0.2.37
Patient Data				TRIBERT	5.0.2.57
Name		Mrs Sonam	Patient ID		011901020083
Birthday			Sample ID		10352349
Age at delivery			Sample Date		02/01/2019
Gestational age		12+1			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	67 Diabetes		no	Pregnancies	
Smoker	no Origin		Asian	-	
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+1
PAPP-A	2.63 mIU/ml	0.90	Method		CRL (<>Hadlock)
fb-hCG	73.77 ng/ml	1.63	Scan Date		02-01-19
Risks at sampling date			CRL 55.3		
Age Risk		1:981	Nuchal translucency MoM		1.09
Biochemical T21 Risk		1:1556	Nasal Bone		Present
Combined Trisomy 21 Risk		1:4869	Sonographer		DR.VIKAS GOYAL
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	DMRD
Risk Risk 1:10 1:250 1:250 1:250 1:1000 1:250 Cut off 1:10000 1:100000 1:10000 1:100000 1:10000 1:10000 1:100000 1:100000			Down's Syndrome Risk (Trisomy 21 Screening) The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 4869 women with the same data, there is one woman with a trisomy 21 pregnancy and 4868 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 risk calculations are statistical approach and have no diagnostic values! The patient combined risk presumes the NT measurement was done according to accepted guidelines The laboratory can not be held responsible for their impact on the risk assessment! Calculated value has no diagnostic value!		

