

*Free Home				
Sample Collection 9999 778 778				

Download "MOLQ" App on

Book a Test Online www.molq.in

				Date of Report PRISCA	17/1/2022 5.2.0.13
Patient Data					
Name	MRS. VI	NITA BHATI	Patient ID		012201160006
Birthday		17/01/1992	Sample ID		11307524
Age at delivery		30	Sample Date		16/1/2022
Gestational age		13+0			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	73 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+3
PAPP-A	1.93 mIU/ml	0.46	Method		CRL (<>Robinson)
fb-hCG	8.29 ng/ml	0.2	Scan date		12/1/2022
Risks at sampling date			Nuchal translucency 2.02		
Age Risk		1:657	Nuchal translucency MoM		1.32
Overall population risk		1:5521	Nasal Bone		Present
Biochemical T21 risk		1:9311			
Trisomy 13/18		1:430			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:100 1:250 Cut off 1:1000			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 it is expected that among 9311 women with the same data, there is one woman with a trisomy 21 pregnancy and 9310 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		
1:10000 13 15 17 19 21 23 25 2 Trisomy 13/18 (with NT The calculated risk for T which indicates a low ris Risk	) Frisomy 13/18 with	Age	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 Age Risk Risk below Age risk		