

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

Book a Test Online www.molq.in

				Date of Report PRISCA	26/6/2023 5.2.0.13
Patient Data					
Name	Ν	ARS. MEENU	Patient ID		012306250179
Birthday		24/12/2002	Sample ID		11656221
Age at term		21.1	Sample Date		25/6/2023
Gestational age		13+2			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	44 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			13+2
PAPP-A	4.47 mIU/ml	0.44	Method		CRL (<>Robinson)
fb-hCG	10.8 ng/ml	0.32	Scan date		22/6/2023
Risks at sampling date			Crown rump length in mm 72		
Age Risk		1:1109	Nuchal translucency MoM		1.11
Biochemical T21 risk		1:8095	Nasal bone		Present
Combined trisomy 21 risk		<1:10000			
Trisomy 13/18 + NT		1:5080			
Risk 1:10			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 1:1000 1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			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 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! The laboratory cannot be hold responsible for their impact		
The calculated risk for Trisomy 13/18 (with NT) is 1:5080, which indicates a low risk			on the risk assessment! Calculated risks have no diagnostic values		
Risk	Above Cut Off		Risk above Ag	e Risk 📃 F	Risk below Age risk