

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

Book a Test Online www.molq.in

				Date of Report PRISCA	17/8/2023 5.2.0.13
Patient Data					
Name	S	AMYUKTHA	Patient ID		012308160035
Birthday		14/05/1996	Sample ID		11792866
Age at term		27.9	Sample Date		16/8/2023
Gestational age		12+6			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	54.6 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			12+5
PAPP-A	3.75 mIU/ml	0.55	Method		CRL (<>Robinson)
fb-hCG	28.3 ng/ml	0.8	Scan date		15/8/2023
Risks at sampling date			Crown rump length in mm 63.5		
Age Risk		1:854	Nuchal translucency MoM		0.98
Biochemical T21 risk		1:2026	Nasal bone		Present
Combined trisomy 21 risk		1:9123			
Trisomy 13/18 + NT		<1:10000			
Risk 1:10			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 Cut off			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 expected that among 9123 women with the same data, there is one woman with a trisomy 21 pregnancy and 9122 women with not affected pregnancies.		
:1000 10000 13 15 17 19 21 23 25 27 2	29 31 33 35 37 39	41 43 45 47 49	the information	l risk by PRISCA depend n provided by the referri isk calculations are statist ostic value!	ing physician. Please
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 4	Above Cut Off		Risk above Ag	e Risk 📃 R	lisk below Age risk